

APPROVED DRUG PRODUCTS

WITH

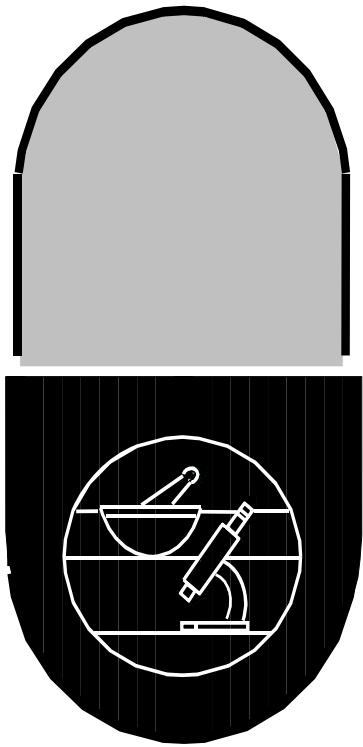
THERAPEUTIC
EQUIVALENCE
EVALUATIONS

32nd 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
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF PHARMACEUTICAL SCIENCE
OFFICE OF GENERIC DRUGS

2012



APPROVED DRUG PRODUCTS

WITH

THERAPEUTIC
EQUIVALENCE
EVALUATIONS

32nd 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
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF PHARMACEUTICAL SCIENCE
OFFICE OF GENERIC DRUGS

2012

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, 2011.

32nd EDITION



**U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF PHARMACEUTICAL SCIENCE
OFFICE OF GENERIC DRUGS**

2012

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
APPROVED DRUG PRODUCTS
with
Therapeutic Equivalence Evaluations**

CONTENTS

	<i>PAGE</i>
PREFACE TO THIRTY SECOND EDITION.....	iv
2 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
 DRUG PRODUCT LISTS	
Prescription Drug Product List	3-1
OTC Drug Product List	4-1
Drug Products with Approval under Section 505 of the 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
 APPENDICES	
A. Product Name Index	A-1
B. Product Name Index Listed by Applicant	B-1
C. Uniform Terms	C-1
 PATENT AND EXCLUSIVITY INFORMATION ADDENDUM	
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 SECOND 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 Act). Drugs on the market approved only on the basis of safety (covered by the ongoing Drug Efficacy Study Implementation [DESI] review [e.g., Donnatal® Tablets and Librax® Capsules] or pre-1938 drugs [e.g., Phenobarbital Tablets]) are not included in this publication. The main criterion for the inclusion of any product is that the product is the subject of an application with an effective approval that has not been withdrawn for safety or efficacy reasons. Inclusion of products on the List is independent of any current regulatory action through administrative or judicial means against a drug product. In addition, the List 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 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 stating 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 List 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 Act.

The therapeutic equivalence evaluations in the List reflect FDA's application of specific criteria to the multisource prescription drug products on the List approved under Section 505 of the Act. These evaluations are presented in the form of code letters that indicate the basis for the evaluation made. An explanation of the code 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, October 1980, of the final version of the List incorporated appropriate corrections and additions. Each subsequent edition has included the 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 (1984 Amendments). The 1984 Amendments require that FDA, among other things, make publicly available a list of approved drug products with monthly supplements. The *Approved Drug Products with Therapeutic Equivalence Evaluations* publication and its monthly Cumulative Supplements satisfy this requirement. The *Addendum* to this publication identifies drugs that qualify under the 1984 Amendments for periods of exclusivity (during which ANDAs or applications described in Section 505(b)(2) of the Act for those drugs may not be submitted for a specified period of time and, if allowed to be submitted, would be tentatively approved) and provides patent information concerning the listed drugs which also may delay the approval of ANDAs or Section 505(b)(2) applications. The *Addendum* also provides additional information that may be helpful to those submitting a new drug application 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 Labeling and Program Support, HFD-610, Office of Generic Drugs, Center for Drug and Evaluation and Research, 7620 Standish Place, Rockville, MD 20855. Comments received are publicly available to the extent allowable under the Freedom of Information regulations.

1. INTRODUCTION

1.1 Content and Exclusion

The List 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 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, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing.¹ This publication also includes indices of prescription and OTC drug products by trade or established name (if no trade name exists) and by applicant name (holder of the approved application). All established names for active ingredients generally conform to official compendial names or *United States Adopted Names* (USAN) as prescribed in (21 CFR 299.4(e)). The latter list includes applicants' names as abbreviated in this publication; in addition, a list of uniform terms is provided.

An *Addendum* contains drug patent and exclusivity information for the Prescription, OTC, Discontinued Drug Product Lists, and for the Drug Products with Approval under Section 505 of the 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 Act administered by the Center for Biologics Evaluation and Research because the main purpose of the publication was to provide information to states regarding FDA's recommendation as to which generic prescription drug products were acceptable candidates for drug product selection. The 1984 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 1984 Amendments, some drug products are given tentative approvals. The Agency will not include drug products with tentative approval in the List. Tentative approval lists are available at [ANDA \(Generic\) Drug Approvals](#). When the tentative approval becomes a full approval through a subsequent action letter to the application holder, the Agency will list the drug product and the final approval date in the appropriate approved drug product list.

Distributors or repackagers of products on the List are not identified. Because distributors or repackagers are not required to notify FDA when they shift their sources of supply from one approved manufacturer to another, it is not possible to maintain complete information linking product approval with the distributor or repackager handling the products.

1.2 Therapeutic Equivalence-Related Terms

Pharmaceutical Equivalents. Drug products are considered pharmaceutical equivalents if they contain the same active ingredient(s), are of the same

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

dosage form, route of administration and are identical in strength or concentration (e.g., chlordiazepoxide hydrochloride, 5mg capsules). Pharmaceutically equivalent drug products are formulated to contain the same amount of active ingredient in the same dosage form and to meet the same or compendial or other applicable standards (i.e., strength, quality, purity, and identity), but they may differ in characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration 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). Data are generally not available for FDA to make the determination of tablet to capsule bioequivalence. 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 and if they can be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.

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; (5) they are manufactured in compliance with Current Good Manufacturing Practice regulations. *The concept of therapeutic equivalence, as used to develop the List, 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., propoxyphene hydrochloride vs. pentazocine hydrochloride for the treatment of pain).* Any drug product in the List repackaged and/or distributed by other than the application holder is considered to be therapeutically equivalent to the application holder's drug product even if the application holder's drug product is single source or coded as non-equivalent (e.g., BN). Also, distributors or repackagers of an application holder's drug product are considered to have the same code as the application holder. Therapeutic equivalence determinations are not made for unapproved, off-label indications.

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 minor 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 particular brand be dispensed as a medical necessity. With this limitation, however, FDA believes that 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.

Bioavailability. This term means 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 action. For drug products that are not intended to be absorbed into the bloodstream, bioavailability may be assessed by measurements intended to reflect the rate and extent to which the active ingredient or active moiety becomes available at the site of action.

Bioequivalent Drug Products. This term describes pharmaceutical equivalent or pharmaceutical alternative products that display comparable bioavailability when studied under similar experimental conditions. Section 505 (j)(8)(B) of the 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 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 *in vivo* or *in vitro* test methods to demonstrate bioequivalence may be appropriate.

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 Statistical Criteria for Bioequivalence

Under the Drug Price Competition and Patent Term Restoration Act of 1984, manufacturers seeking approval to market a generic drug product must submit data demonstrating that the drug product is bioequivalent to the pioneer (innovator) drug product. A major premise underlying the 1984 law is that bioequivalent drug products are therapeutically equivalent, and therefore, interchangeable.

Bioavailability refers to the rate and extent to which the active ingredient or therapeutic ingredient is absorbed from a drug product and becomes available at the site of drug action (Federal Food, Drug and Cosmetic Act, section 505(j)(8)). Bioequivalence refers to equivalent release of the same drug substance from two or more drug products or formulations. This leads to an equivalent rate and extent of absorption from these formulations. Underlying the concept of bioequivalence is the thesis that, if a drug product contains a drug substance that is chemically identical and is delivered to the site of action at the same rate and extent as another drug product, then it is equivalent and can be substituted for that drug product. Methods used to define bioequivalence can be found in 21 CFR 320.24, and include (1) pharmacokinetic (PK) studies, (2) pharmacodynamic (PD) studies, (3) comparative clinical trials, and (4) *in-vitro* studies. The choice of study used is based on the site of action of the drug and the ability of the study design to compare drug delivered to that site by the two products.

The standard bioequivalence (PK) study is conducted using a two-treatment crossover study design in a limited number of volunteers, usually 24 to 36

adults. Alternately, a four-period, replicate design crossover study may also be used. Single doses of the test and reference drug products are administered and blood or plasma levels of the drug are measured over time. Pharmacokinetic parameters characterizing rate and extent of drug absorption are evaluated statistically. The PK parameters of interest are the resulting area under the plasma concentration-time curve (AUC), calculated to the last measured concentration ($AUC_{(0-t)}$) and extrapolated to infinity ($AUC_{(0-\infty)}$), for extent of absorption; and the maximum or peak drug concentrations (Cmax), for rate of absorption. Crossover studies may not be practical in drugs with a long half-life in the body, and a parallel study design may be used instead. Alternate study methods, such as in-vitro studies or equivalence studies with clinical or pharmacodynamic endpoints, are used for drug products where plasma concentrations are not useful to determine delivery of the drug substance to the site of activity (such as inhalers, nasal sprays and topical products applied to the skin).

The statistical methodology for analyzing these bioequivalence studies is called the two one-sided test procedure. Two situations are tested with this statistical methodology. The first of the two one-sided tests determines whether a generic product (test), when substituted for a brand-name product (reference) is significantly less bioavailable. The second of the two one-sided tests determines whether a brand-name product when substituted for a generic product is significantly less bioavailable. Based on the opinions of FDA medical experts, a difference of greater than 20% for each of the above tests was determined to be significant, and therefore, undesirable for all drug products. Numerically, this is expressed as a limit of test-product average/reference-product average of 80% for the first statistical test and a limit of reference-product average/test-product average of 80% for the second statistical test. By convention, all data is expressed as a ratio of the average response (AUC and Cmax) for test/reference, so the limit expressed in the second statistical test is 125% (reciprocal of 80%).

For statistical reasons, all data is log-transformed prior to conducting statistical testing. In practice, these statistical tests are carried out using an analysis of variance procedure (ANOVA) and calculating a 90% confidence interval for each pharmacokinetic parameter (Cmax and AUC). The confidence interval for both pharmacokinetic parameters, AUC and Cmax, must be entirely within the 80% to 125% boundaries cited above. Because the mean of the study data lies in the center of the 90% confidence interval, the mean of the data is usually close to 100% (a test/reference ratio of 1). Different statistical criteria are sometimes used when bioequivalence is demonstrated through comparative clinical trials pharmacodynamic studies, or comparative in-vitro methodology.

The bioequivalence methodology and criteria described above simultaneously control for both differences in the average response between test and reference, as well as the precision with which the average response in the population is estimated. This precision depends on the within-subject (normal volunteer or patient) variability in the pharmacokinetic parameters (AUC and Cmax) of the two products and on the number of subjects in the study. The width of the 90% confidence interval is a reflection in part of the within-subject variability of the test and reference products in the bioequivalence study. A test product with no differences in the average response when compared to the reference might still fail to pass the bioequivalence criteria if the variability of one or both products is high and the bioequivalence study has insufficient statistical power (i.e., insufficient number of subjects). Likewise, a test product with low variability may pass the bioequivalence criteria, when there are somewhat larger differences in the average response.

This system of assessing bioequivalence of generic products assures that these substitutable products do not deviate substantially in in-vivo performance from the reference product. The Office of Generic Drugs has conducted two surveys to quantify the differences between generic and brand name products. The first survey included 224 bioequivalence studies submitted

in approved applications during 1985 and 1986. The observed average differences between reference and generic products for AUC was 3.5% (JAMA, Sept. 4, 1987, Vol. 258, No. 9). The second survey included 127 bioequivalence studies submitted to the agency in 273 ANDAs approved in 1997.

The three measures reviewed include $AUC_{(0-t)}$, $AUC_{(0-inf)}$, and Cmax. The observed average differences between the reference and generic products were $\pm 3.47\%$ ($SD 2.84$) for $AUC_{(0-t)}$, $\pm 3.25\%$ ($SD 2.97$) for $AUC_{(0-inf)}$, and $\pm 4.29\%$ ($SD 3.72$) for Cmax (JAMA, Dec. 1, 1999, Vol. 282, No. 21).

The primary concern from the regulatory point of view is the protection of the patient against approval of products that are not bioequivalent. The current practice of carrying out two one-sided tests at the 0.05 level of significance ensures that there is no more than a 5% chance that a generic product that is not truly equivalent to the reference will be approved.

1.4 Reference Listed Drug

A reference listed drug (21 CFR 314.94(a)(3)) means the listed drug identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.

FDA has identified in the Prescription Drug Product and OTC Drug Product Lists those reference listed drugs to which the *in vivo* bioequivalence (reference standard) and, in some instances, the *in vitro* bioequivalence of the applicant's product is compared. By designating a single reference listed drug as the standard to which all generic versions must be shown to be bioequivalent, FDA hopes to avoid possible significant variations among generic drugs and their brand name counterpart. Such variations could result if generic drugs were compared to different reference listed drugs. However, in some instances when listed drugs are approved for a single drug product, a product not designated as the reference listed drug and not shown to be bioequivalent to the reference listed drug may be shielded from generic competition. A firm wishing to market a generic version of a listed drug that is not designated as the reference listed drug may petition the Agency through the Citizen Petition procedure (see 21 CFR 10.25(a) and CFR 10.30). When the Citizen Petition is approved, the second listed drug will be designated as an additional reference listed drug and the petitioner may submit an Abbreviated New Drug Application citing the designated reference listed drug. Section 1.7, *Therapeutic Equivalence Evaluations Codes products meeting necessary bioequivalence requirements* explains the (**AB**, **AB1**, **AB2**, **AB3**...) coding system for multisource drug products listed under the same heading with two reference listed drugs.

In addition, there are two situations in which two listed drugs that have been shown to be bioequivalent to each other may both be designated as reference listed drugs. The first situation occurs when the *in vivo* determination of bioequivalence is self-evident and a waiver of the *in vivo* bioequivalence may be granted. The second situation occurs when the bioequivalence of two listed products may be determined through *in vitro* methodology. The reference listed drug is identified by the symbol "+" in the Prescription and Over-the-Counter (OTC) Drug Product Lists. These identified reference listed drugs represent the best judgment of the Division of Bioequivalence at this time. The Prescription and OTC Drug Product Lists identify reference drugs for oral dosage forms, Injectables, ophthalmics, otics, and topical products. It is recommended that a firm planning to conduct an *in vivo* bioequivalence study, or planning to manufacture a batch of a drug product for which an *in vivo* waiver of bioequivalence will be requested, contact the Division of Bioequivalence, Office of Generic Drugs, to confirm the appropriate reference listed drug.

1.5 General Policies and Legal Status

The List contains public information and advice. It does not mandate the drug products which is purchased, prescribed, dispensed, or substituted for one another, nor does it, conversely, mandate the products that should be avoided. To the extent that the List 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 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 List identifies drug products approved under Section 505 of the 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 List does not necessarily mean that the drug product is either in violation of Section 505 of the Act, or 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 is 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 List. 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. There may also be better stability of one product over another under adverse storage conditions, or 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 specification of that product is appropriate. Pharmacists must also be familiar with the expiration dates/times and labeling directions for storage of the different products, particularly for reconstituted products, to assure that patients are properly advised when one product is substituted for another.

Multisource and single-source drug products. FDA has evaluated for therapeutic equivalence only multisource prescription drug products approved under Section 505 of the 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, in addition, 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 on the List, but no therapeutic equivalence code is included with such products. Any drug product

in the List repackaged and/or distributed by other than the application holder is considered to be therapeutically equivalent to the application holder's drug product even if the application holder's drug product is single source or coded as non-equivalent (e.g., BN). Also, although not identified in the List, distributors or repackagers of an application holder's drug product are considered to have the same code as the application holder. The details of these codes and the policies underlying them are discussed in Section 1.7, *Therapeutic Equivalence Evaluations Codes*.

Products on the List are identified by the names of the holders of approved applications (applicants) who may not necessarily be the manufacturer of the 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 most 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 marketer.

Although the products on the List 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 on the List are, in most cases, correct.

To relate firm name information on a product label to that on the List, the following should be noted: the applicant's name always appears on the List. This applies whether the applicant (firm name on the Form FDA 356h in the application) is the 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 on the List 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 on the List 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 on the List; 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 List, 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 on the List, the following should be noted: if the applicant is the marketer, its name appears on the List 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 drug name if no trade name exists) appears on the List. If a corporate relationship exists between an application holder 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 application holder to appear on the List. If there is no known corporate relationship between the applicant and the marketer, the established drug name appears on the List.

Every product on the List 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 Act. In such circumstances, the Agency will 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 on the List. The main criterion for

inclusion of a product is that it has an application with an effective approval that has not been withdrawn for safety or efficacy reasons. FDA believes that retention of a violative product on the List 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 data upon which the Agency's assessment of whether a product meets the criteria for therapeutic equivalence was made.

1.7 Therapeutic Equivalence Evaluations Codes

The coding system for therapeutic equivalence evaluations is constructed to allow users to determine quickly whether the Agency has evaluated a particular approved product 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 few exceptions, 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 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 actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bio-equivalence. 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 presumed and considered self-evident based on other data in the application for some dosage forms (e.g., solutions) or satisfied for

solid oral dosage forms 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 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. For example, 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. An FDA evaluation that such products are therapeutically equivalent is applicable only when each product is reconstituted, stored, and used under the conditions specified in the labeling of that product.

The Agency will 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 Special Situations*.

For example, in rare instances, there may be variations among therapeutically 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 will 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, 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 and esters of the same therapeutic moiety are regarded as pharmaceutical alternatives. For the purpose of this publication, such products are not considered to be therapeutically equivalent. There are no instances in this List where pharmaceutical alternatives are evaluated or coded with regard to therapeutic equivalence. Anhydrous and hydrated entities, as well as different polymorphs, are considered pharmaceutical equivalents and must meet the same standards and, where necessary, as in the case of ampicillin/ampicillin trihydrate, their equivalence is supported by appropriate bioavailability/bioequivalence studies.

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 may differ among some therapeutically equivalent drug products. Differences in preservatives and other inactive ingredients 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

Products coded as **AA** contain active ingredients and 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, Pharmaceutical Equivalents*) generally will be coded **AB** if a study is 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 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 drug products which are not 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 (Miles) and Procardia XL® (Pfizer), extended-release tablets, are listed under the active ingredient nifedipine. These drug products, listed under the same heading, are not bioequivalent to each other. Generic drug products deemed by FDA to be bioequivalent to Adalat® CC and Procardia XL® have been approved, Adalat® CC and Procardia XL® have been assigned ratings of **AB1** and **AB2**, respectively. The generic drug products bioequivalent to Adalat® CC would be assigned a rating of **AB1** and those bioequivalent to Procardia XL® would be 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 List, they are considered therapeutically equivalent to the application holder's drug product if the application holder's drug product is rated either with an **AB** or three-character code or is single source in the List. 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 any of several 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. 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, 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 on the List (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.

The strength of parenteral drugs products is defined as the total drug content of the container. Until recently the strength of liquid parenteral drug products in the Orange Book have not been displayed. The concentration of the liquid parenteral drug product in the Orange Book has been shown as xmg/ml. The amount of dry powder or freeze dried powder in a container has always been identified as the strength.

With the finalization of the Waxman-Hatch amendments that characterized each strength of a drug product as a listed drug, it became evident that the format of the Orange Book should be changed to reflect each strength of a parenteral solution. To this end the OGD has started to display 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 drug product had a 20 ml and 60 ml container approved the two products would be shown as 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 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 bio-equivalence 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 medicament 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 firms 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 *in vivo* bioequivalence data are submitted, 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 bio-equivalent only if *in vivo* evidence of bioequivalence is available. In those cases where *in vivo* evidence is available, the product is 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 Special Situations

Certain drugs listed in the Orange Book present special situations that merit further discussion. Following is a description 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.

Follitropin Alfa and Beta. Based on available data derived from physico-chemical tests and bioassay, follitropin alfa and follitropin beta are indistinguishable.

Gaviscon®. Gaviscon® is an OTC product which 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 which 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 which cite Gaviscon® tablets as the 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 drug products.

Levothyroxine Sodium (Mylan ANDA 76187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Unithroid (Jerome Stevens NDA 021210) tablets.

Levo-T (Alara NDA 021342), Levothyroxine Sodium (Mylan ANDA 76187), Unithroid (Jerome Stevens NDA 021210), and Levothyroxine Sodium (Merck KGAA ANDA 76752) tablets have been determined to be therapeutically equivalent to corresponding strengths of Synthroid (Abbott NDA 021402) tablets.

Levo-T (Alara NDA 021342), Unithroid (Jerome Stevens NDA 021210), Levothyroxine Sodium (Mylan ANDA 076187), and Levothyroxine Sodium (Merck KGAA ANDA 76752) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levoxyl (King Pharms NDA 021301) tablets.

Levothyroxine Sodium (Mylan ANDA 76187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levothroid (Lloyd NDA 021116) tablets.

The chart outlines TE codes for all 0.025mg products. 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 common TE code indicates therapeutic equivalence between products.

Trade Name	Applicant	Potency	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	ABBOTT	0.025MG	AB1	021402	001
LEVO-T	ALARA PHARM	0.025MG	AB1	021342	001
SYNTHROID	ABBOTT	0.025MG	AB2	021402	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB2	076187	001
LEVO-T	ALARA PHARM	0.025MG	AB2	021342	001
UNITHROID	STEVENS J	0.025MG	AB2	021210	001
LEVOTHYROXINE SODIUM	MERCK KGAA	0.025MG	AB2	076752	001
LEVOXYL	KING PHARMS	0.025MG	AB3	021301	001
LEVO-T	ALARA PHARM	0.025MG	AB3	021342	001
UNITHROID	STEVENS J	0.025MG	AB3	021210	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB3	076187	001
LEVOTHYROXINE SODIUM	MERCK KGAA	0.025MG	AB3	076752	001
LEVOTHROID	LLOYD	0.025MG	AB4	021116	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB4	076187	001

Patent Certification(s) Reference Listed Drug based upon a suitability petition. An abbreviated new drug application that refers to a Reference Listed Drug (RLD) approved pursuant to a suitability petition must demonstrate that the proposed product is bioequivalent to the RLD, and it must include appropriate patent certification(s) and an exclusivity statement with respect to the listed drug which served as the basis for the approved suitability petition. This concept also applies to an ANDA applicant that cites a RLD that was based upon an NDA that is still covered by patent (s) and/or

exclusivity, e.g. a second RLD that was selected when the *in vivo* determination of bioequivalence of the original RLD is self evident and the waiver of the *in vivo* determination of bioequivalence may be granted.

Waived exclusivity. If a new drug application (NDA) submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (Act) qualifies for exclusivity under sections 505(c)(3)(D) and 505(j)(5)(D), the exclusivity is listed in the Patent and Exclusivity Section of the Orange Book. If a drug product has received this exclusivity, the FDA will delay the approval of a 505(b)(2) application or an abbreviated new drug application (ANDA) under section 505(j) of the Act until the expiration of the exclusivity. If the listed drug is also protected by one or more patents, the approval date for the 505(b)(2) application or ANDA will be determined by the latest expiring patent or exclusivity listed in the Orange Book. However, the holder of the NDA may waive its exclusivity as to any or all 505(b)(2) and ANDA applications referencing the protected drug product. If an NDA sponsor waives its right to the exclusivity protection, qualified 505(b)(2) or ANDA applications may be approved without regard to the NDA holder's exclusivity. An NDA for which the holder has waived its exclusivity as to all 505(b)(2) and ANDA applications will be coded with a W in the Patent and Exclusivity Section of the Orange Book and be referred to this section. The applicant referencing this listed drug should indicate in the exclusivity statement that the holder of the listed drug has waived its exclusivity.

1.9 Therapeutic Equivalence Code Change for a Drug Entity

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 multi-source 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 drug products in the List (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 a 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 drugs, the Agency will announce in the *Introduction* to the Cumulative Supplement that it is considering the change and will invite comment. Comments, along with scientific data, may be sent to the Director, Division 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 submission is an *in vivo* bioavailability/bioequivalence study conducted on batches of the subject drug products. These submissions 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 such submissions are discouraged. Copies

of supporting reports published in the scientific literature or unpublished material, however, are welcome.

1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product

The aforementioned procedure 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. 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.

1.11 Discontinued Section

Those drug products in the Discontinued Section of the Orange Book in which a determination has already been made that the products were not withdrawn for safety or efficacy reasons have “**Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**” following the product strength. Those drug products are only reflective of citizen petitions determinations made since 1995. The identification of these drug products in the Discontinued Section of the Orange Book should avoid the submission of multiple citizen petitions for the same drug product. FR notices no longer applicable are removed from the Annual Edition (i.e., there is a currently marketed Reference Listed Drug and no applicable patent or exclusivity). [FR Safety or Effectiveness Determinations List](#) lists products that have current and removed notices. The list is updated quarterly. Notices issued during the year are added to the [Electronic Orange Book Query](#) in the month they become effective.

Generally, approved products are added to the Discontinued Section of the Orange Book when the applicant holder notifies the Orange Book staff of the products' not marketed status. Products may also be added if annual reports indicate the product is no longer marketed or other Agency administrative action (e.g., Withdrawal of an Application). Changes to the Orange Book are not affected by the drug registration and listing requirements of Section 510 of the Act.

1.12 Changes to the Orange Book

Every effort is made to ensure the Annual Edition is current and accurate. Applicant holders are requested to inform the FDA Orange Book Staff (OBS) of any changes or corrections. Please inform the OBS when products are no longer marketed. Notification of the Orange Book staff to include the newly approved product in the Discontinued Drug Product List rather than parts 1, 2 or 3 of the List (as discussed in Section 1.1) must occur 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

We can be contacted by email at drugproducts@fda.hhs.gov. Send Changes by FAX: 240-276-8974; mail to:

FDA/CDER Orange Book Staff
Office of Generic Drugs, HFD-610
7620 Standish Place
Rockville, MD 20855

1.13 Availability of the Edition

Commencing with the 25th edition, the Annual Edition and current monthly Cumulative Supplement are available in a Portable Document Format (PDF) at the EOB home page, [Electronic Orange Book Query](#), 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 Printing 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, application holders, 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 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.

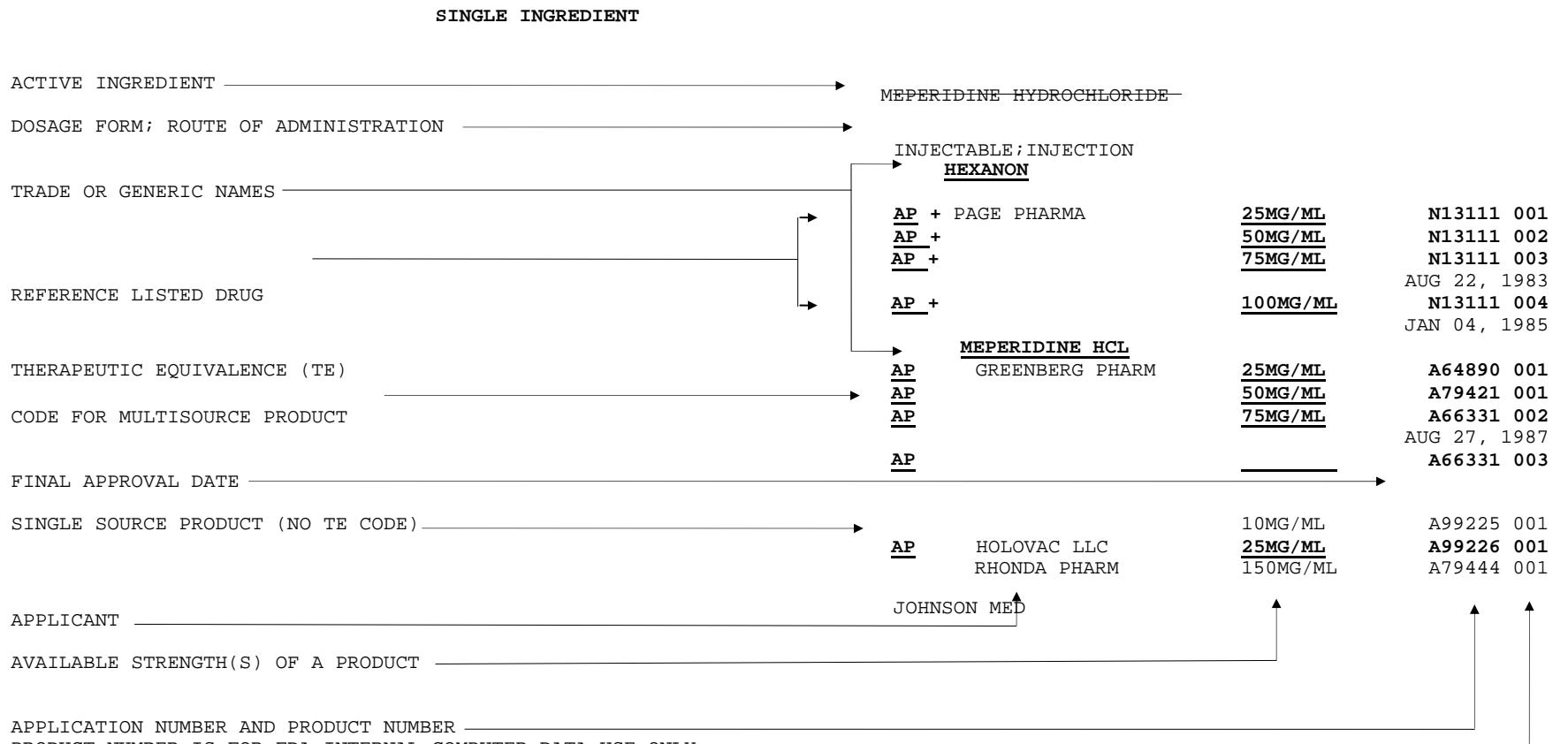
The Discontinued Drug Product List contains approved products that have never been marketed, have been discontinued from marketing, are for military use, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing. All products having a "@" in the 12th Cumulative Supplement of the 31st Edition List have been added to the Discontinued Drug Product List appearing in the 32nd 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 Section of the Orange Book.

Product Name Index (*Prescription and OTC Drug Product Lists*). This is an index of drug products by established or trade name. 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.

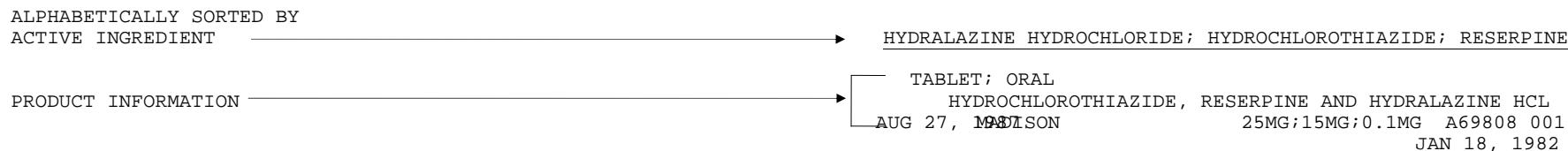
Product Name Index Listed by Applicant (*Prescription and OTC Drug Product Lists*). This is an index that cross-references applicants to drug products. The bolded and underlined entry represents the applicant name abbreviation used in this publication. Each complete applicant name that is represented 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



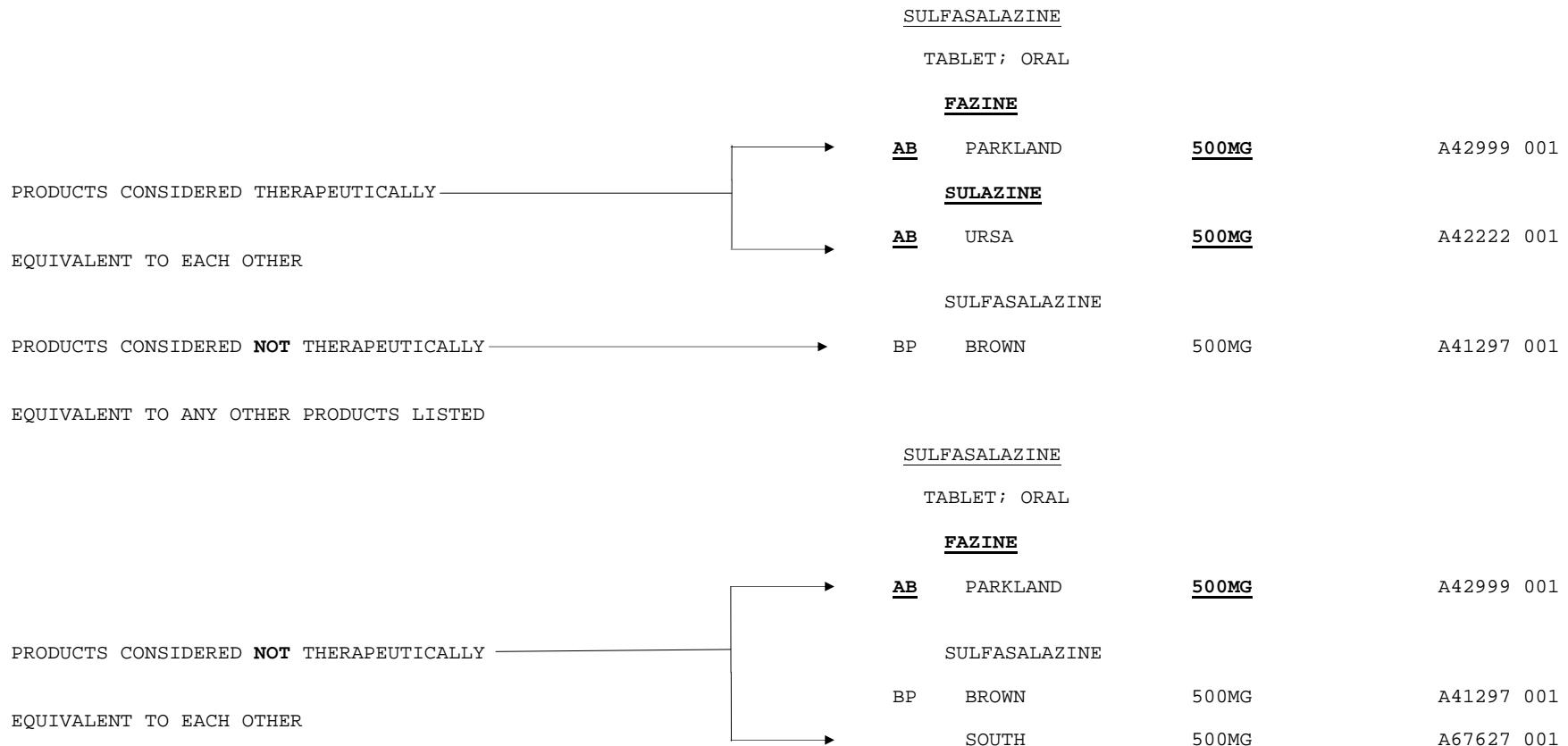
MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION



THIS EXAMPLE IS FOR PURPOSES 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.

PRESCRIPTION DRUG PRODUCT LIST

3 - 1 (of 424)

ABACAVIR SULFATE

SOLUTION; ORAL ZIAGEN			
+ VIIIV HLTHCARE	EQ 20MG BASE/ML	N020978 001	Dec 17, 1998
TABLET; ORAL ZIAGEN			
+ VIIIV HLTHCARE	EQ 300MG BASE	N020977 001	Dec 17, 1998

ABACAVIR SULFATE; LAMIVUDINE

TABLET; ORAL EPZICOM			
+ VIIIV HLTHCARE	EQ 600MG BASE;300MG	N021652 001	Aug 02, 2004

ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL TRIZIVIR			
+ VIIIV HLTHCARE	EQ 300MG BASE;150MG;300MG	N021205 001	Nov 14, 2000

ABIRATERONE ACETATE

TABLET; ORAL ZYTIGA			
+ JANSSEN BIOTECH	250MG	N202379 001	Apr 28, 2011

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL CAMPRAL			
+ FOREST LABS	333MG	N021431 001	Jul 29, 2004

ACARBOSE

TABLET; ORAL <u>ACARBOSE</u>			
<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> ROXANE	<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>AB</u> STRIDES ARCOLAB LTD	<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> 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>PRECOSE</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 <u>ACEBUTOLOL HYDROCHLORIDE</u>			
<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>AB</u> WATSON LABS	<u>EQ 200MG BASE</u>	<u>A074007 001</u>	Oct 18, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 2 (of 424)

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

<u>AB</u>	WATSON LABS	<u>EQ 400MG BASE</u>	<u>A074007 002</u>	Oct 18, 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)

+ CADENCE PHARMS	1000MG/100ML (10MG/ML)	N022450 001	Nov 02, 2010
------------------	------------------------	-------------	--------------

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

PHRENILIN FORTE

+ VALEANT	650MG;50MG
-----------	------------

A088831 001 Jun 19, 1985

TABLET; ORAL

BUTAPAP

<u>AA</u>	MIKART	<u>325MG;50MG</u>	<u>A089987 001</u>	Oct 26, 1992
	<u>PHRENILIN</u>			
<u>AA</u>	+ VALEANT	<u>325MG;50MG</u>	<u>A087811 001</u>	Jun 19, 1985
	BUTALBITAL AND ACETAMINOPHEN			
	NEXGEN PHARMA	300MG;50MG	A090956 001	Aug 23, 2011
	BUTAPAP			
	+ MIKART	650MG;50MG	A089988 001	Oct 26, 1992

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	WEST WARD	<u>500MG;50MG;40MG</u>	<u>A040261 001</u>	Oct 28, 1998
	<u>ESGIC-PLUS</u>			
<u>AA</u>	+ MIKART	<u>500MG;50MG;40MG</u>	<u>A040085 001</u>	Mar 28, 1996
	BUTALBITAL, ACETAMINOPHEN AND CAFFEINE			
	+ MIKART	325MG;50MG;40MG	A089007 001	Mar 17, 1986
	+ NEXGEN PHARMA	300MG;50MG;40MG	A040885 001	Nov 16, 2009

SOLUTION; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

+ MIKART	325MG/15ML;50MG/15ML;40MG/15ML
----------	--------------------------------

A040387 001 Jan 31, 2003

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	MALLINCKRODT	<u>325MG;50MG;40MG</u>	<u>A087804 001</u>	Jan 24, 1985
<u>AA</u>	MIKART	<u>325MG;50MG;40MG</u>	<u>A089175 001</u>	Jan 21, 1987
<u>AA</u>	MIRROR PHARMS	<u>325MG;50MG;40MG</u>	<u>A040864 001</u>	Dec 01, 2008
<u>AA</u>		<u>500MG;50MG;40MG</u>	<u>A040883 001</u>	Dec 23, 2008
<u>AA</u>	VINTAGE PHARMS	<u>325MG;50MG;40MG</u>	<u>A040511 001</u>	Aug 27, 2003
<u>AA</u>		<u>500MG;50MG;40MG</u>	<u>A040513 001</u>	Aug 25, 2003
<u>AA</u>	WATSON LABS	<u>500MG;50MG;40MG</u>	<u>A040267 001</u>	Jul 30, 1998
<u>AA</u>	WEST WARD	<u>325MG;50MG;40MG</u>	<u>A089718 001</u>	Jun 12, 1995
<u>AA</u>		<u>500MG;50MG;40MG</u>	<u>A040336 001</u>	Aug 18, 1999
	<u>ESGIC-PLUS</u>			
<u>AA</u>	+ MIKART	<u>500MG;50MG;40MG</u>	<u>A089451 001</u>	May 23, 1988
	<u>FIORICET</u>			
<u>AA</u>	+ WATSON PHARMS	<u>325MG;50MG;40MG</u>	<u>A088616 001</u>	Nov 09, 1984
	BUTALBITAL, ACETAMINOPHEN AND CAFFEINE			
	+ MIKART	750MG;50MG;40MG	A040496 001	Dec 23, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 3 (of 424)

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

<u>AB</u>	NEXGEN PHARMA INC	<u>325MG;50MG;40MG;30MG</u>	<u>A076560</u> <u>001</u>	Jun 10, 2004
<u>AB</u>	VINTAGE PHARMS	<u>325MG;50MG;40MG;30MG</u>	<u>A075929</u> <u>001</u>	Apr 22, 2002
<u>AB</u>	WEST WARD	<u>325MG;50MG;40MG;30MG</u>	<u>A075618</u> <u>001</u>	Mar 23, 2001
	<u>FIORICET W/ CODEINE</u>			
<u>AB</u>	+ WATSON LABS INC	<u>325MG;50MG;40MG;30MG</u>	<u>N020232</u> <u>001</u>	Jul 30, 1992
	<u>PHRENILIN WITH CAFFEINE AND CODEINE</u>			
<u>AB</u>	VALEANT	<u>325MG;50MG;40MG;30MG</u>	<u>A074911</u> <u>001</u>	Aug 22, 2001

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

<u>AA</u>	+ MIKART	<u>356.4MG;30MG;16MG</u>	<u>A040109</u> <u>001</u>	Aug 26, 1997
<u>AA</u>	WRASER PHARMS LLC	<u>356.4MG;30MG;16MG</u>	<u>A040688</u> <u>001</u>	Apr 03, 2007

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

<u>AA</u>	BOCA PHARMA	<u>712.8MG;60MG;32MG</u>	<u>A040701</u> <u>001</u>	Apr 03, 2007
<u>AA</u>	+ MIKART	<u>712.8MG;60MG;32MG</u>	<u>A040316</u> <u>001</u>	Apr 28, 1999

ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

<u>AA</u>	HI TECH PHARMA	<u>120MG/5ML;12MG/5ML</u>	<u>A040119</u> <u>001</u>	Apr 26, 1996
<u>AA</u>	MIKART	<u>120MG/5ML;12MG/5ML</u>	<u>A089450</u> <u>001</u>	Oct 27, 1992
<u>AA</u>	PHARM ASSOC	<u>120MG/5ML;12MG/5ML</u>	<u>A087508</u> <u>001</u>	
<u>AA</u>	VINTAGE PHARMS	<u>120MG/5ML;12MG/5ML</u>	<u>A091238</u> <u>001</u>	Nov 10, 2011
<u>AA</u>	WOCKHARDT	<u>120MG/5ML;12MG/5ML</u>	<u>A087006</u> <u>001</u>	

SUSPENSION; ORAL

CAPITAL AND CODEINE

<u>AA</u>	+ VALEANT	<u>120MG/5ML;12MG/5ML</u>	<u>A086024</u> <u>001</u>
-----------	-----------	---------------------------	---------------------------

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

<u>AA</u>	AMNEAL PHARMS NY	<u>300MG;30MG</u>	<u>A040779</u> <u>001</u>	May 29, 2008
<u>AA</u>	+ MALLINCKRODT INC	<u>300MG;15MG</u>	<u>A040419</u> <u>001</u>	May 31, 2001
<u>AA</u>		<u>300MG;30MG</u>	<u>A040419</u> <u>002</u>	May 31, 2001
<u>AA</u>		<u>300MG;60MG</u>	<u>A040419</u> <u>003</u>	May 31, 2001
<u>AA</u>	MIKART	<u>300MG;30MG</u>	<u>A089238</u> <u>001</u>	Feb 25, 1986
<u>AA</u>	RANBAXY	<u>300MG;30MG</u>	<u>A085868</u> <u>001</u>	
<u>AA</u>		<u>300MG;60MG</u>	<u>A087083</u> <u>001</u>	
<u>AA</u>	TEVA	<u>300MG;15MG</u>	<u>A088627</u> <u>001</u>	Mar 06, 1985
<u>AA</u>		<u>300MG;30MG</u>	<u>A088628</u> <u>001</u>	Mar 06, 1985
<u>AA</u>	+ VINTAGE	<u>300MG;60MG</u>	<u>A088629</u> <u>001</u>	Mar 06, 1985
<u>AA</u>		<u>300MG;15MG</u>	<u>A089990</u> <u>001</u>	Sep 30, 1988
<u>AA</u>		<u>300MG;30MG</u>	<u>A089805</u> <u>001</u>	Sep 30, 1988
<u>AA</u>	VINTAGE PHARMS	<u>300MG;60MG</u>	<u>A089828</u> <u>001</u>	Sep 30, 1988
	<u>TYLENOL W/ CODEINE NO. 3</u>			
<u>AA</u>	+ JANSSEN PHARMS	<u>300MG;30MG</u>	<u>A085055</u> <u>003</u>	
	<u>TYLENOL W/ CODEINE NO. 4</u>			
<u>AA</u>	JANSSEN PHARMS	<u>300MG;60MG</u>	<u>A085055</u> <u>004</u>	
	ACETAMINOPHEN AND CODEINE PHOSPHATE			
+ MIKART		<u>650MG;30MG</u>	<u>A089231</u> <u>001</u>	Mar 03, 1986
+ MIKART		<u>650MG;60MG</u>	<u>A089363</u> <u>001</u>	Sep 09, 1991

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	MIKART	<u>500MG;5MG</u>	<u>A081067</u> <u>001</u>	Nov 30, 1989
-----------	--------	------------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 4 (of 424)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	MIKART	<u>500MG;5MG</u>	<u>A089008</u> <u>001</u>	Feb 21, 1986
SOLUTION; ORAL				
		<u>HYDROCODONE BITARTRATE AND ACETAMINOPHEN</u>		
<u>AA</u>	BOCA PHARMA	<u>325MG/15ML; 7.5MG/15ML</u>	<u>A040894</u> <u>001</u>	Jul 19, 2011
<u>AA</u>	MALLINCKRODT	<u>500MG/15ML; 7.5MG/15ML</u>	<u>A040418</u> <u>001</u>	Jun 27, 2001
<u>AA</u>	+ MIKART	<u>325MG/15ML; 7.5MG/15ML</u>	<u>A040482</u> <u>001</u>	Sep 25, 2003
<u>AA</u>	+	<u>500MG/15ML; 7.5MG/15ML</u>	<u>A081051</u> <u>001</u>	Aug 28, 1992
<u>AA</u>	NESHER PHARMS	<u>500MG/15ML; 7.5MG/15ML</u>	<u>A040366</u> <u>001</u>	Jan 23, 2002
<u>AA</u>	PHARM ASSOC	<u>500MG/15ML; 7.5MG/15ML</u>	<u>A040182</u> <u>001</u>	Mar 13, 1998
<u>AA</u>	VINTAGE PHARMS	<u>500MG/15ML; 7.5MG/15ML</u>	<u>A040520</u> <u>001</u>	Oct 30, 2003
HYDROCODONE BITARTRATE AND ACETAMINOPHEN				
+ MIKART		<u>300MG/15ML; 10MG/15ML</u>	<u>A040881</u> <u>001</u>	Feb 25, 2010
+ PHARM ASSOC		<u>325MG/15ML; 10MG/15ML</u>	<u>A040834</u> <u>001</u>	Apr 18, 2008
TABLET; ORAL				
		<u>ANEXSIA</u>		
<u>AA</u>	MALLINCKRODT	<u>500MG;5MG</u>	<u>A089160</u> <u>001</u>	Apr 23, 1987
<u>AA</u>		<u>750MG;10MG</u>	<u>A040468</u> <u>001</u>	Oct 31, 2002
		<u>ANEXSIA 5/325</u>		
<u>AA</u>	MALLINCKRODT	<u>325MG;5MG</u>	<u>A040409</u> <u>001</u>	Oct 20, 2000
		<u>ANEXSIA 7.5/325</u>		
<u>AA</u>	MALLINCKRODT	<u>325MG;7.5MG</u>	<u>A040405</u> <u>001</u>	Sep 08, 2000
		<u>ANEXSIA 7.5/650</u>		
<u>AA</u>	MALLINCKRODT	<u>650MG;7.5MG</u>	<u>A089725</u> <u>001</u>	Sep 30, 1987
		<u>CO-GESIC</u>		
<u>AA</u>	UCB INC	<u>500MG;5MG</u>	<u>A087757</u> <u>001</u>	May 03, 1982
HYDROCODONE BITARTRATE AND ACETAMINOPHEN				
<u>AA</u>	AMNEAL PHARMS NY	<u>325MG;5MG</u>	<u>A040736</u> <u>001</u>	Aug 25, 2006
<u>AA</u>		<u>325MG;10MG</u>	<u>A040746</u> <u>001</u>	Aug 25, 2006
<u>AA</u>		<u>500MG;5MG</u>	<u>A040729</u> <u>001</u>	Aug 25, 2006
<u>AA</u>		<u>500MG;7.5MG</u>	<u>A040748</u> <u>001</u>	Aug 25, 2006
<u>AA</u>		<u>500MG;10MG</u>	<u>A040813</u> <u>001</u>	Feb 23, 2007
<u>AA</u>		<u>650MG;7.5MG</u>	<u>A040754</u> <u>001</u>	Aug 25, 2006
<u>AA</u>		<u>650MG;10MG</u>	<u>A040757</u> <u>001</u>	Aug 25, 2006
<u>AA</u>		<u>750MG;7.5MG</u>	<u>A040769</u> <u>001</u>	Aug 28, 2006
<u>AA</u>	BOCA PHARMA	<u>300MG;5MG</u>	<u>A090415</u> <u>001</u>	Jan 24, 2011
<u>AA</u>		<u>300MG;7.5MG</u>	<u>A090415</u> <u>002</u>	Jan 24, 2011
<u>AA</u>		<u>300MG;10MG</u>	<u>A090415</u> <u>003</u>	Jan 24, 2011
<u>AA</u>	MALLINCKRODT	<u>325MG;10MG</u>	<u>A040400</u> <u>001</u>	Jul 26, 2000
<u>AA</u>		<u>500MG;5MG</u>	<u>A040084</u> <u>002</u>	Jun 01, 1995
<u>AA</u>		<u>500MG;7.5MG</u>	<u>A040201</u> <u>001</u>	Feb 27, 1998
<u>AA</u>		<u>500MG;10MG</u>	<u>A040201</u> <u>002</u>	Feb 27, 1998
<u>AA</u>		<u>650MG;10MG</u>	<u>A040084</u> <u>004</u>	Oct 16, 1996
<u>AA</u>	+	<u>660MG;10MG</u>	<u>A040084</u> <u>003</u>	Jul 29, 1996
<u>AA</u>		<u>750MG;7.5MG</u>	<u>A040084</u> <u>001</u>	Jun 01, 1995
<u>AA</u>	+ MIKART	<u>300MG;5MG</u>	<u>A040658</u> <u>001</u>	Jan 19, 2006
<u>AA</u>	+	<u>300MG;7.5MG</u>	<u>A040556</u> <u>002</u>	Mar 24, 2006
<u>AA</u>	+	<u>300MG;10MG</u>	<u>A040556</u> <u>001</u>	Jun 23, 2004
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A040432</u> <u>001</u>	Jan 22, 2003
<u>AA</u>	+	<u>500MG;2.5MG</u>	<u>A089698</u> <u>001</u>	Aug 25, 1989
<u>AA</u>	+	<u>500MG;7.5MG</u>	<u>A089699</u> <u>001</u>	Aug 25, 1989
<u>AA</u>	+	<u>650MG;7.5MG</u>	<u>A089689</u> <u>001</u>	Jun 29, 1988
<u>AA</u>	+	<u>650MG;10MG</u>	<u>A081223</u> <u>001</u>	May 29, 1992
<u>AA</u>	SUN PHARM INDs INC	<u>325MG;5MG</u>	<u>A090118</u> <u>001</u>	Dec 23, 2008
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A090118</u> <u>002</u>	Dec 23, 2008
<u>AA</u>		<u>325MG;10MG</u>	<u>A090118</u> <u>003</u>	Dec 23, 2008
<u>AA</u>		<u>500MG;5MG</u>	<u>A090265</u> <u>001</u>	Dec 23, 2008
<u>AA</u>		<u>500MG;7.5MG</u>	<u>A090265</u> <u>002</u>	Dec 23, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 5 (of 424)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	SUN PHARM INDs INC	<u>500MG;10MG</u>	<u>A090265</u> <u>003</u>	Dec 23, 2008
<u>AA</u>		<u>650MG;7.5MG</u>	<u>A090380</u> <u>001</u>	Dec 23, 2008
<u>AA</u>		<u>650MG;10MG</u>	<u>A090380</u> <u>002</u>	Dec 23, 2008
<u>AA</u>		<u>660MG;10MG</u>	<u>A090380</u> <u>003</u>	Dec 23, 2008
<u>AA</u>		<u>750MG;7.5MG</u>	<u>A090380</u> <u>004</u>	Dec 23, 2008
<u>AA</u>	VINTAGE PHARMS	<u>325MG;5MG</u>	<u>A040655</u> <u>001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A040656</u> <u>001</u>	Jan 19, 2006
<u>AA</u>		<u>325MG;10MG</u>	<u>A040355</u> <u>001</u>	May 31, 2000
<u>AA</u>		<u>500MG;2.5MG</u>	<u>A040144</u> <u>002</u>	Apr 25, 1997
<u>AA</u>		<u>500MG;5MG</u>	<u>A089971</u> <u>001</u>	Dec 02, 1988
<u>AA</u>		<u>500MG;7.5MG</u>	<u>A040144</u> <u>001</u>	Feb 22, 1996
<u>AA</u>		<u>500MG;10MG</u>	<u>A040356</u> <u>001</u>	May 31, 2000
<u>AA</u>		<u>650MG;7.5MG</u>	<u>A040155</u> <u>001</u>	Apr 14, 1997
<u>AA</u>		<u>650MG;10MG</u>	<u>A040143</u> <u>001</u>	Feb 22, 1996
<u>AA</u>		<u>660MG;10MG</u>	<u>A040358</u> <u>001</u>	May 31, 2000
<u>AA</u>		<u>750MG;7.5MG</u>	<u>A040157</u> <u>001</u>	Apr 12, 1996
<u>AA</u>	WATSON LABS	<u>500MG;2.5MG</u>	<u>A081079</u> <u>001</u>	Aug 30, 1991
<u>AA</u>		<u>500MG;5MG</u>	<u>A089883</u> <u>001</u>	Dec 01, 1988
<u>AA</u>		<u>500MG;7.5MG</u>	<u>A081080</u> <u>001</u>	Aug 30, 1991
<u>AA</u>		<u>500MG;10MG</u>	<u>A040148</u> <u>002</u>	Feb 14, 1997
<u>AA</u>		<u>650MG;7.5MG</u>	<u>A040094</u> <u>001</u>	Sep 29, 1995
<u>AA</u>		<u>650MG;10MG</u>	<u>A040094</u> <u>002</u>	Sep 29, 1995
<u>AA</u>		<u>660MG;10MG</u>	<u>A040094</u> <u>003</u>	Aug 08, 2000
<u>AA</u>		<u>750MG;7.5MG</u>	<u>A081083</u> <u>001</u>	Aug 30, 1991
<u>AA</u>	+	<u>750MG;10MG</u>	<u>A040094</u> <u>004</u>	Mar 22, 1999
<u>AA</u>	WATSON LABS FLORIDA	<u>660MG;10MG</u>	<u>A040495</u> <u>001</u>	May 28, 2003
	<u>LORTAB</u>			
<u>AA</u>	UCB INC	<u>500MG;5MG</u>	<u>A087722</u> <u>001</u>	Jul 09, 1982
<u>AA</u>	+	<u>500MG;10MG</u>	<u>A040100</u> <u>001</u>	Jan 26, 1996
	<u>NORCO</u>			
<u>AA</u>	+	<u>325MG;5MG</u>	<u>A040099</u> <u>001</u>	Jun 25, 1997
<u>AA</u>	+	<u>325MG;7.5MG</u>	<u>A040148</u> <u>003</u>	Sep 12, 2000
<u>AA</u>	+	<u>325MG;10MG</u>	<u>A040148</u> <u>001</u>	Feb 14, 1997
	<u>VICODIN</u>			
<u>AA</u>	+	<u>500MG;5MG</u>	<u>A088058</u> <u>001</u>	Jan 07, 1983
	<u>VICODIN ES</u>			
<u>AA</u>	+	<u>750MG;7.5MG</u>	<u>A089736</u> <u>001</u>	Dec 09, 1988
	<u>VICODIN HP</u>			
<u>AA</u>	ABBOTT	<u>660MG;10MG</u>	<u>A040117</u> <u>001</u>	Sep 23, 1996
	HYDROCODONE BITARTRATE AND ACETAMINOPHEN			
	MIKART	<u>325MG;2.5MG</u>	<u>A040846</u> <u>001</u>	Jun 09, 2010
		<u>650MG;5MG</u>	<u>A040849</u> <u>001</u>	Jun 09, 2010
	ZYDONE			
+	ENDO PHARMS	<u>400MG;5MG</u>	<u>A040288</u> <u>001</u>	Nov 27, 1998
+		<u>400MG;7.5MG</u>	<u>A040288</u> <u>002</u>	Nov 27, 1998
+		<u>400MG;10MG</u>	<u>A040288</u> <u>003</u>	Nov 27, 1998

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	DURAMED PHARMS BARR	<u>500MG;5MG</u>	<u>A040289</u> <u>001</u>	Mar 16, 1999
<u>AA</u>	MALLINCKRODT	<u>500MG;5MG</u>	<u>A040257</u> <u>001</u>	Aug 04, 1998
<u>AA</u>	VINTAGE PHARMS	<u>500MG;5MG</u>	<u>A040106</u> <u>001</u>	Jul 30, 1996
<u>AA</u>	WATSON LABS	<u>500MG;5MG</u>	<u>A040234</u> <u>001</u>	Oct 30, 1997
	<u>ROXILOX</u>			
<u>AA</u>	ROXANE	<u>500MG;5MG</u>	<u>A040061</u> <u>001</u>	Jul 03, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 6 (of 424)

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

TYLOXAA + JANSSEN PHARMS 500MG;5MG A088790 001 Dec 12, 1984

SOLUTION; ORAL

ROXICETAA + ROXANE 325MG/5ML;5MG/5ML A089351 001 Dec 03, 1986

TABLET; ORAL

OXYCETAA MALLINCKRODT 325MG;5MG A087463 001 Dec 07, 1983OXYCODONE AND ACETAMINOPHENAA AMNEAL PHARMS NY 325MG;5MG A040777 001 Nov 27, 2007AA 325MG;10MG A040778 001 Nov 27, 2007AA 500MG;7.5MG A040789 001 Nov 27, 2007AA 650MG;10MG A040789 002 Nov 27, 2007AA COASTAL PHARMS 325MG;2.5MG A090177 001 Oct 20, 2008AA 325MG;5MG A090177 002 Oct 20, 2008AA 325MG;7.5MG A090177 003 Oct 20, 2008AA 325MG;10MG A090177 004 Oct 20, 2008AA 500MG;7.5MG A090177 005 Oct 20, 2008AA 650MG;10MG A090177 006 Oct 20, 2008AA MALLINCKRODT 325MG;7.5MG A040545 001 Jun 30, 2004AA 325MG;10MG A040545 002 Jun 30, 2004AA 500MG;7.5MG A040550 001 Jun 30, 2004AA 650MG;10MG A040550 002 Jun 30, 2004AA VINTAGE PHARMS 325MG;5MG A040105 001 Jul 30, 1996AA WATSON LABS 325MG;5MG A040171 001 Oct 30, 1997AA 325MG;7.5MG A040535 001 Sep 05, 2003AA 325MG;10MG A040535 002 Sep 05, 2003AA 500MG;7.5MG A040371 001 Dec 29, 2000AA 650MG;10MG A040371 002 Dec 29, 2000PERCOSETAA + ENDO PHARMS 325MG;2.5MG A040330 001 Jun 25, 1999AA + 325MG;5MG A040330 002 Jun 25, 1999AA + 325MG;7.5MG A040434 001 Nov 23, 2001AA + 325MG;10MG A040434 002 Nov 23, 2001AA + 500MG;7.5MG A040341 001 Jul 26, 1999AA + 650MG;10MG A040341 002 Jul 26, 1999ROXICETAA ROXANE 325MG;5MG A087003 001OXYCODONE 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+ 400MG;2.5MG A040679 001 May 16, 2006+ 400MG;5MG A040687 001 Apr 27, 2006+ 400MG;7.5MG A040698 001 Apr 27, 2006+ 400MG;10MG A040692 001 Apr 27, 2006+ 500MG;10MG A040676 001 Apr 19, 2006

ROXICET 5/500

+ ROXANE 500MG;5MG A089775 001 Jan 12, 1989ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDEAB GAVIS PHARMS 650MG;EQ 25MG BASE A076202 001 Aug 02, 2002AB + WATSON LABS 650MG;EQ 25MG BASE A074699 001 Mar 24, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 7 (of 424)

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

<u>AB</u>	AMNEAL PHARMS	<u>325MG;37.5MG</u>	<u>A090485</u> <u>001</u>	Dec 09, 2009
<u>AB</u>	CARACO	<u>325MG;37.5MG</u>	<u>A077184</u> <u>001</u>	Dec 16, 2005
<u>AB</u>	MYLAN	<u>325MG;37.5MG</u>	<u>A077858</u> <u>001</u>	Sep 26, 2008
<u>AB</u>	PAR PHARM	<u>325MG;37.5MG</u>	<u>A076475</u> <u>001</u>	Apr 21, 2005
<u>AB</u>	WATSON LABS	<u>325MG;37.5MG</u>	<u>A076914</u> <u>001</u>	Jul 26, 2006
	<u>ULTRACET</u>			
<u>AB</u>	+ JANSSEN PHARMS	<u>325MG;37.5MG</u>	<u>N021123</u> <u>001</u>	Aug 15, 2001

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

ACETAZOLAMIDE

<u>AB</u>	HERITAGE PHARMS INC	<u>500MG</u>	<u>A090779</u> <u>001</u>	Jul 14, 2011
<u>AB</u>	ZYDUS PHARMS USA INC	<u>500MG</u>	<u>A040904</u> <u>001</u>	Dec 10, 2008
	<u>DIAMOX</u>			
<u>AB</u>	+ DURAMED PHARMS BARR	<u>500MG</u>	<u>N012945</u> <u>001</u>	

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	LANNETT	<u>250MG</u>	<u>A084840</u> <u>001</u>	
<u>AB</u>	MUTUAL PHARM	<u>125MG</u>	<u>A089752</u> <u>001</u>	Jun 22, 1988
<u>AB</u>	TARO	<u>125MG</u>	<u>A040195</u> <u>001</u>	May 28, 1997
<u>AB</u>	+ DURAMED PHARMS BARR	<u>250MG</u>	<u>A040195</u> <u>002</u>	May 28, 1997
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A088882</u> <u>001</u>	Oct 22, 1985

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

<u>AP</u>	BEDFORD	<u>EQ 500MG BASE/VIAL</u>	<u>A040089</u> <u>001</u>	Feb 28, 1995
<u>AP</u>	X GEN PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A040784</u> <u>001</u>	Dec 10, 2008
	<u>DIAMOX</u>			
<u>AP</u>	+ DURAMED PHARMS BARR	<u>EQ 500MG BASE/VIAL</u>	<u>N009388</u> <u>001</u>	Dec 05, 1990

ACETIC ACID, GLACIAL

SOLUTION; IRRIGATION, URETHRAL

ACETIC ACID 0.25% IN PLASTIC CONTAINER

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

SOLUTION/DROPS; OTIC

ACETIC ACID

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

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC

ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE

<u>+</u>	BAUSCH AND LOMB	<u>2%;0.79%</u>	<u>A040063</u> <u>001</u>	Feb 25, 1994
----------	-----------------	-----------------	---------------------------	--------------

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

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

PRESCRIPTION DRUG PRODUCT LIST

3 - 8 (of 424)

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

HYDROCORTISONE AND ACETIC ACID

<u>AT</u>	VINTAGE	<u>2%;1%</u>	<u>A040609</u>	<u>001</u>	Feb 06, 2006
<u>AT</u>	+ HI TECH PHARMA	<u>2%;1%</u>	<u>N012770</u>	<u>001</u>	

ACETOHEXAMIDE

TABLET; ORAL

ACETOHEXAMIDE

WATSON LABS	250MG	A071893	001	Nov 25, 1987
+	500MG	A071894	001	Nov 25, 1987

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

+ CUMBERLAND PHARMS	6GM/30ML (200MG/ML)	N021539	001	Jan 23, 2004
---------------------	---------------------	---------	-----	--------------

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

<u>AN</u>	BEDFORD	<u>10%</u>	<u>A072323</u>	<u>001</u>	Apr 30, 1992
<u>AN</u>		<u>20%</u>	<u>A072324</u>	<u>001</u>	Apr 30, 1992
<u>AN</u>	HOSPIRA	<u>10%</u>	<u>A073664</u>	<u>001</u>	Aug 30, 1994
<u>AN</u>		<u>20%</u>	<u>A074037</u>	<u>001</u>	Aug 30, 1994
<u>AN</u>	+ LUITPOLD	<u>10%</u>	<u>A072489</u>	<u>001</u>	Jul 28, 1995
<u>AN</u>	+	<u>20%</u>	<u>A072547</u>	<u>001</u>	Jul 28, 1995

ACITRETIN

CAPSULE; ORAL

SORIATANE

STIEFEL LABS INC	10MG	N019821	001	Oct 28, 1996
	17.5MG	N019821	003	Aug 06, 2009
	22.5MG	N019821	004	Aug 06, 2009
+	25MG	N019821	002	Oct 28, 1996

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

SEMPREX-D

+ UCB INC	8MG;60MG	N019806	001	Mar 25, 1994
-----------	----------	---------	-----	--------------

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

<u>AB</u>	APOTEX INC	<u>200MG</u>	<u>A075677</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>	DAVA PHARMS INC	<u>200MG</u>	<u>A074833</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A074727</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>		<u>200MG</u>	<u>A074977</u>	<u>001</u>	Apr 13, 1998
<u>AB</u>	RANBAXY	<u>200MG</u>	<u>A074975</u>	<u>001</u>	Sep 30, 1998
<u>AB</u>	STASON	<u>200MG</u>	<u>A075090</u>	<u>001</u>	Jan 26, 1999
<u>AB</u>	TEVA	<u>200MG</u>	<u>A074578</u>	<u>001</u>	Apr 22, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 9 (of 424)

ACYCLOVIR

CAPSULE; ORAL			
ACYCLOVIR			
<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>A075101 001</u> Apr 15, 1998
ZOVIRAX			
<u>AB</u>	+ GLAXOSMITHKLINE	<u>200MG</u>	<u>N018828 001</u> Jan 25, 1985
CREAM; TOPICAL			
ZOVIRAX			
+ VALEANT INTL	5%		N021478 001 Dec 30, 2002
OINTMENT; TOPICAL			
ZOVIRAX			
+ VALEANT INTL	5%		N018604 001 Mar 29, 1982
SUSPENSION; ORAL			
ACYCLOVIR			
<u>AB</u>	ACTAVIS MID ATLANTIC	<u>200MG/5ML</u>	<u>A074738 001</u> Apr 28, 1997
<u>AB</u>	HI TECH PHARMA	<u>200MG/5ML</u>	<u>A077026 001</u> Jun 07, 2005
ZOVIRAX			
<u>AB</u>	+ GLAXOSMITHKLINE	<u>200MG/5ML</u>	<u>N019909 001</u> Dec 22, 1989
TABLET; ORAL			
ACYCLOVIR			
<u>AB</u>	APOTEX INC	<u>400MG</u>	<u>A077309 001</u> Sep 29, 2005
<u>AB</u>		<u>800MG</u>	<u>A077309 002</u> Sep 29, 2005
<u>AB</u>	CARLSBAD	<u>400MG</u>	<u>A075382 001</u> Apr 30, 1999
<u>AB</u>		<u>800MG</u>	<u>A075382 002</u> Apr 30, 1999
<u>AB</u>	DAVA PHARMS INC	<u>400MG</u>	<u>A074946 001</u> Nov 19, 1997
<u>AB</u>		<u>800MG</u>	<u>A074946 002</u> Nov 19, 1997
<u>AB</u>	MYLAN	<u>400MG</u>	<u>A074976 001</u> Apr 13, 1998
<u>AB</u>		<u>400MG</u>	<u>A075211 001</u> Sep 28, 1998
<u>AB</u>		<u>800MG</u>	<u>A074976 002</u> Apr 13, 1998
<u>AB</u>		<u>800MG</u>	<u>A075211 002</u> Sep 28, 1998
<u>AB</u>	RANBAXY	<u>400MG</u>	<u>A074980 001</u> Sep 30, 1998
<u>AB</u>		<u>800MG</u>	<u>A074980 002</u> Sep 30, 1998
<u>AB</u>	TEVA	<u>400MG</u>	<u>A074556 002</u> Apr 22, 1997
<u>AB</u>		<u>800MG</u>	<u>A074556 003</u> Apr 22, 1997
ZOVIRAX			
<u>AB</u>	GLAXOSMITHKLINE	<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>	+ APP PHARMS	<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>	BAXTER HLTHCARE	<u>EQ 500MG BASE/VIAL</u>	<u>A074913 001</u> Oct 15, 1997
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A074913 002</u> Oct 15, 1997
<u>AP</u>	BEDFORD	<u>EQ 500MG BASE/VIAL</u>	<u>A074596 002</u> Apr 22, 1997
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A074596 001</u> Apr 22, 1997
ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE			
+ BAXTER HLTHCARE		EQ 500MG BASE/VIAL	A074885 001 Dec 19, 1997
		EQ 1GM BASE/VIAL	A074885 002 Dec 19, 1997

ACYCLOVIR; HYDROCORTISONE

CREAM; TOPICAL			
XERESE			
+ VALEANT INTL	5%;1%		N022436 001 Jul 31, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 10 (of 424)

ADAPALENE

CREAM; TOPICAL

ADAPALENE

<u>AB</u>	NYCOMED US	<u>0.1%</u>	<u>A090824</u> 001	Jun 30, 2010
	<u>DIFFERIN</u>			
<u>AB</u>	+ GALDERMA LABS LP	<u>0.1%</u>	<u>N020748</u> 001	May 26, 2000
	GEL; TOPICAL			
	<u>ADAPALENE</u>			
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A091314</u> 001	Jul 01, 2010
<u>AB</u>	PLIVA HRVATSKA DOO	<u>0.1%</u>	<u>A090962</u> 001	Jun 02, 2010
	<u>DIFFERIN</u>			
<u>AB</u>	+ GALDERMA LABS LP	<u>0.1%</u>	<u>N020380</u> 001	May 31, 1996
	DIFFERIN			
	+ GALDERMA LABS LP	0.3%	N021753 001	Jun 19, 2007
	LOTION; TOPICAL			
	DIFFERIN			
	+ GALDERMA R AND D	0.1%	N022502 001	Mar 17, 2010

ADAPALENE; BENZOYL PEROXIDE

GEL; TOPICAL

EPIDUO

+ GALDERMA LABS	0.1%; 2.5%	N022320 001	Dec 08, 2008
-----------------	------------	-------------	--------------

ADEFOVIR DIPIVOXIL

TABLET; ORAL

HEPSERA

+ GILEAD	10MG	N021449 001	Sep 20, 2002
----------	------	-------------	--------------

ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

<u>AP</u>	+ ASTELLAS	<u>3MG/ML</u>	<u>N019937</u> 002	Oct 30, 1989
	<u>ADENOSINE</u>			
<u>AP</u>	AKORN	<u>3MG/ML</u>	<u>A078076</u> 001	Oct 31, 2008
<u>AP</u>	APP PHARMS	<u>3MG/ML</u>	<u>A077133</u> 001	Apr 27, 2005
<u>AP</u>	BAXTER HLTHCARE	<u>3MG/ML</u>	<u>A076500</u> 001	Jun 16, 2004
<u>AP</u>	BEDFORD	<u>3MG/ML</u>	<u>A076404</u> 001	Jun 16, 2004
<u>AP</u>	GLAND PHARMA LTD	<u>3MG/ML</u>	<u>A077283</u> 001	Jun 14, 2007
<u>AP</u>	LUITPOLD	<u>3MG/ML</u>	<u>A090010</u> 001	Apr 28, 2009
<u>AP</u>	STRIDES ARCOLAB LTD	<u>3MG/ML</u>	<u>A078686</u> 001	May 13, 2009
<u>AP</u>	TEVA PARENTERAL	<u>3MG/ML</u>	<u>A076564</u> 001	Jun 16, 2004
<u>AP</u>		<u>3MG/ML</u>	<u>A078676</u> 001	Jul 31, 2008
<u>AP</u>	WOCKHARDT	<u>3MG/ML</u>	<u>A090220</u> 001	Jul 20, 2009
	ADENOSCAN			
	+ ASTELLAS	3MG/ML	N020059 001	May 18, 1995

ALBENDAZOLE

TABLET; ORAL

ALBENZA

+ COREPHARMA	200MG	N020666 001	Jun 11, 1996
--------------	-------	-------------	--------------

ALBUMIN HUMAN

INJECTABLE; INJECTION

OPTISON

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

PRESCRIPTION DRUG PRODUCT LIST

3 - 11 (of 424)

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 GLOBAL	EQ 0.09MG BASE/INH	N021457 001	Oct 29, 2004
PROVENTIL-HFA			
BX + 3M	EQ 0.09MG BASE/INH	N020503 001	Aug 15, 1996
VENTOLIN HFA			
BX + GLAXOSMITHKLINE	EQ 0.09MG BASE/INH	N020983 001	Apr 19, 2001
SOLUTION; INHALATION			
<u>ACCUNEB</u>			
AN + DEY	<u>EQ 0.021% BASE</u>	<u>N020949 002</u>	Apr 30, 2001
AN +	<u>EQ 0.042% BASE</u>	<u>N020949 001</u>	Apr 30, 2001
<u>ALBUTEROL SULFATE</u>			
AN APOTEX INC	<u>EQ 0.021% BASE</u>	<u>A078623 001</u>	Apr 05, 2010
AN	<u>EQ 0.042% BASE</u>	<u>A078623 002</u>	Apr 05, 2010
AN	<u>EQ 0.083% BASE</u>	<u>A075717 001</u>	Feb 02, 2007
AN + BAUSCH AND LOMB	<u>EQ 0.5% BASE</u>	<u>A075050 001</u>	Jun 18, 1998
AN + DEY	<u>EQ 0.083% BASE</u>	<u>A072652 001</u>	Feb 21, 1992
AN HI TECH PHARMA	<u>EQ 0.083% BASE</u>	<u>A075063 001</u>	Feb 09, 1999
AN	<u>EQ 0.5% BASE</u>	<u>A074543 001</u>	Jan 15, 1998
AN LANDELA PHARM	<u>EQ 0.083% BASE</u>	<u>A077569 001</u>	Apr 04, 2006
AN NEPHRON	<u>EQ 0.021% BASE</u>	<u>A076355 002</u>	Mar 31, 2010
AN	<u>EQ 0.042% BASE</u>	<u>A076355 001</u>	Jun 28, 2004
AN	<u>EQ 0.083% BASE</u>	<u>A074880 001</u>	Sep 17, 1997
AN	<u>EQ 0.5% BASE</u>	<u>A075664 001</u>	Jun 26, 2001
AN NOVEX	<u>EQ 0.5% BASE</u>	<u>A076391 001</u>	Apr 01, 2003
AN RITEDOSE CORP	<u>EQ 0.083% BASE</u>	<u>A077839 001</u>	Dec 16, 2008
AN TEVA PARENTERAL	<u>EQ 0.083% BASE</u>	<u>A075343 001</u>	Nov 09, 1999
AN WATSON LABS	<u>EQ 0.021% BASE</u>	<u>A077772 001</u>	Sep 25, 2007
AN	<u>EQ 0.042% BASE</u>	<u>A077772 002</u>	Sep 25, 2007
AN WATSON LABS INC	<u>EQ 0.083% BASE</u>	<u>A076370 001</u>	Nov 24, 2003
AN WOCKHARDT	<u>EQ 0.083% BASE</u>	<u>A075394 001</u>	Nov 22, 1999
SYRUP; ORAL			
<u>ALBUTEROL SULFATE</u>			
AA ACTAVIS MID ATLANTIC	<u>EQ 2MG BASE/5ML</u>	<u>A074454 001</u>	Sep 25, 1995
AA AMNEAL PHARMS	<u>EQ 2MG BASE/5ML</u>	<u>A079241 001</u>	May 12, 2010
AA HI TECH PHARMA	<u>EQ 2MG BASE/5ML</u>	<u>A074749 001</u>	Jan 30, 1998
AA + TEVA	<u>EQ 2MG BASE/5ML</u>	<u>A073419 001</u>	Mar 30, 1992
AA VINTAGE	<u>EQ 2MG BASE/5ML</u>	<u>A078105 001</u>	Dec 27, 2006
AA VISTAPHARM	<u>EQ 2MG BASE/5ML</u>	<u>A077788 001</u>	Jun 26, 2007
TABLET; ORAL			
<u>ALBUTEROL SULFATE</u>			
AB MUTUAL PHARM	<u>EQ 2MG BASE</u>	<u>A072637 002</u>	Dec 05, 1989
AB	<u>EQ 4MG BASE</u>	<u>A072637 001</u>	Dec 05, 1989
AB MYLAN	<u>EQ 2MG BASE</u>	<u>A072894 002</u>	Jan 17, 1991
AB +	<u>EQ 4MG BASE</u>	<u>A072894 001</u>	Jan 17, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 12 (of 424)

ALBUTEROL SULFATE

TABLET; ORAL

ALBUTEROL SULFATE

<u>AB</u>	WATSON LABS	<u>EQ 2MG BASE</u>	<u>A072764 001</u>	Aug 28, 1991
TABLET, EXTENDED RELEASE; ORAL				
<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
<u>VOSPIRE ER</u>				
<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

AEROSOL, 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

<u>AN</u>	APOTEX CORP	<u>EQ 0.083% BASE;0.017%</u>	<u>A077117 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>	SANDOZ	<u>EQ 0.083% BASE;0.017%</u>	<u>A076867 001</u>	Dec 21, 2006
<u>AN</u>	TEVA PARENTERAL	<u>EQ 0.083% BASE;0.017%</u>	<u>A076724 001</u>	Dec 31, 2007
<u>AN</u>	WATSON LABS	<u>EQ 0.083% BASE;0.017%</u>	<u>A077063 001</u>	Dec 31, 2007
<u>AN</u>		<u>EQ 0.083% BASE;0.017%</u>	<u>A077559 001</u>	Dec 31, 2007
<u>DUONEB</u>				
<u>AN</u> +	DEY	<u>EQ 0.083% BASE;0.017%</u>	<u>N020950 001</u>	Mar 21, 2001

SPRAY, METERED; INHALATION

COMBIVENT RESPIMAT

+ BOEHRINGER INGELHEIM EQ 0.1MG BASE/INH;0.02MG/INH

N021747 001 Oct 07, 2011

ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

LASTACAFT

+ ALLERGAN 0.25%

N022134 001 Jul 28, 2010

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ACLOVATE

<u>AB</u> +	FOUGERA PHARMS	<u>0.05%</u>	<u>N018707 001</u>	Dec 14, 1982
<u>ALCLOMETASONE DIPROPIONATE</u>				
<u>AB</u>	ALTANA	<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

ACLOVATE

<u>AB</u> +	FOUGERA PHARMS	<u>0.05%</u>	<u>N018702 001</u>	Dec 14, 1982
<u>ALCLOMETASONE DIPROPIONATE</u>				
<u>AB</u>	ALTANA	<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

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION

ALCOHOL 5% AND DEXTROSE 5%

<u>AP</u> +	B BRAUN	<u>5ML/100ML;5GM/100ML</u>	<u>N004589 004</u>
ALCOHOL 10% AND DEXTROSE 5%			
+ B BRAUN		<u>10ML/100ML;5GM/100ML</u>	<u>N004589 006</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 13 (of 424)

ALENDRONATE SODIUM

SOLUTION; ORAL

FOSAMAX

+ MERCK

EQ 70MG BASE/75ML

N021575 001 Sep 17, 2003

TABLET; ORAL

ALENDRONATE SODIUM

AB	APOTEX	<u>EQ 5MG BASE</u>	<u>A077982 001</u>	Aug 04, 2008
AB		<u>EQ 10MG BASE</u>	<u>A077982 002</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A077982 003</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A077982 004</u>	Aug 04, 2008
AB	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A090124 001</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A090124 002</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A090124 003</u>	Aug 04, 2008
AB	AUSTARPHARMA LLC	<u>EQ 5MG BASE</u>	<u>A090258 001</u>	Sep 24, 2009
AB		<u>EQ 10MG BASE</u>	<u>A090258 002</u>	Sep 24, 2009
AB		<u>EQ 35MG BASE</u>	<u>A090258 003</u>	Sep 24, 2009
AB		<u>EQ 70MG BASE</u>	<u>A090258 004</u>	Sep 24, 2009
AB	CADISTA PHARMS	<u>EQ 5MG BASE</u>	<u>A090557 001</u>	Feb 18, 2010
AB		<u>EQ 10MG BASE</u>	<u>A090557 002</u>	Feb 18, 2010
AB		<u>EQ 35MG BASE</u>	<u>A090557 003</u>	Feb 18, 2010
AB		<u>EQ 70MG BASE</u>	<u>A090557 004</u>	Feb 18, 2010
AB	DR REDDYS LABS LTD	<u>EQ 5MG BASE</u>	<u>A079109 001</u>	Aug 04, 2008
AB		<u>EQ 10MG BASE</u>	<u>A079109 002</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A079049 001</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A079049 002</u>	Aug 04, 2008
AB	MYLAN	<u>EQ 5MG BASE</u>	<u>A076584 001</u>	Aug 04, 2008
AB		<u>EQ 10MG BASE</u>	<u>A076584 002</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A076584 003</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A078638 001</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A076584 004</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A078638 002</u>	Aug 04, 2008
AB	SUN PHARMA GLOBAL	<u>EQ 5MG BASE</u>	<u>A090022 001</u>	Sep 10, 2008
AB		<u>EQ 10MG BASE</u>	<u>A090022 002</u>	Sep 10, 2008
AB		<u>EQ 35MG BASE</u>	<u>A090022 003</u>	Sep 10, 2008
AB		<u>EQ 70MG BASE</u>	<u>A090022 004</u>	Sep 10, 2008
AB	TEVA PHARMS	<u>EQ 5MG BASE</u>	<u>A075710 001</u>	Feb 06, 2008
AB		<u>EQ 10MG BASE</u>	<u>A075710 002</u>	Feb 06, 2008
AB		<u>EQ 35MG BASE</u>	<u>A075710 003</u>	Feb 06, 2008
AB		<u>EQ 40MG BASE</u>	<u>A075710 004</u>	Feb 06, 2008
AB		<u>EQ 70MG BASE</u>	<u>A075710 005</u>	Feb 06, 2008
AB	WATSON LABS	<u>EQ 5MG BASE</u>	<u>A076768 001</u>	Aug 04, 2008
AB		<u>EQ 10MG BASE</u>	<u>A076768 002</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A076768 003</u>	Aug 04, 2008
AB		<u>EQ 35MG BASE</u>	<u>A076984 001</u>	Aug 04, 2008
AB		<u>EQ 40MG BASE</u>	<u>A076768 004</u>	Aug 04, 2008
AB		<u>EQ 40MG BASE</u>	<u>A076984 002</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A076768 005</u>	Aug 04, 2008
AB		<u>EQ 70MG BASE</u>	<u>A076984 003</u>	Aug 04, 2008
<u>FOSAMAX</u>				
AB	MERCK AND CO INC	<u>EQ 5MG BASE</u>	<u>N020560 003</u>	Apr 25, 1997
AB		<u>EQ 10MG BASE</u>	<u>N020560 001</u>	Sep 29, 1995
AB		<u>EQ 35MG BASE</u>	<u>N020560 004</u>	Oct 20, 2000
AB		<u>EQ 40MG BASE</u>	<u>N020560 002</u>	Sep 29, 1995
AB	+	<u>EQ 70MG BASE</u>	<u>N020560 005</u>	Oct 20, 2000

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

PRESCRIPTION DRUG PRODUCT LIST

3 - 14 (of 424)

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> MYLAN	<u>10MG</u>	<u>A079014 001</u>	Jul 18, 2011
<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> TTorrent PHARMS	<u>10MG</u>	<u>A079054 001</u>	Jul 18, 2011
UROXATRAL			
<u>AB</u> + SANOFI AVENTIS US	<u>10MG</u>	<u>N021287 001</u>	Jun 12, 2003

ALGLUCERASE

INJECTABLE; INJECTION

CEREDASE

+ GENZYME

80 UNITS/ML

N020057 003

Apr 05, 1991

ALISKIREN HEMIFUMARATE

TABLET; ORAL

TEKTURNNA

NOVARTIS

EQ 150MG BASE

N021985 001

Mar 05, 2007

+

EQ 300MG BASE

N021985 002

Mar 05, 2007

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE

TABLET; ORAL

TEKAMLO

NOVARTIS

EQ 150MG BASE;EQ 5MG BASE

N022545 001

Aug 26, 2010

EQ 150MG BASE;EQ 10MG BASE

N022545 002

Aug 26, 2010

EQ 300MG BASE;EQ 5MG BASE

N022545 003

Aug 26, 2010

+

EQ 300MG BASE;EQ 10MG BASE

N022545 004

Aug 26, 2010

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMTURNIDE

NOVARTIS

EQ 150MG BASE;EQ 5MG BASE;12.5MG

N200045 001

Dec 21, 2010

EQ 300MG BASE;EQ 5MG BASE;12.5MG

N200045 002

Dec 21, 2010

EQ 300MG BASE;EQ 5MG BASE;25MG

N200045 003

Dec 21, 2010

EQ 300MG BASE;EQ 10MG BASE;12.5MG

N200045 004

Dec 21, 2010

+

EQ 300MG BASE;EQ 10MG BASE;25MG

N200045 005

Dec 21, 2010

ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEKTURNNA HCT

NOVARTIS

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

ALISKIREN HEMIFUMARATE; VALSARTAN

TABLET; ORAL

VALTURNA

NOVARTIS

EQ 150MG BASE;160MG

N022217 001

Sep 16, 2009

+

EQ 300MG BASE;320MG

N022217 002

Sep 16, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 15 (of 424)

ALITRETIINOIN

GEL; TOPICAL
PANRETIN
+ EISAI INC EQ 0.1% BASE N020886 001 Feb 02, 1999

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077353</u> <u>001</u>	Sep 08, 2005
<u>AB</u>		<u>300MG</u>	<u>A077353</u> <u>002</u>	Sep 08, 2005
<u>AB</u>	CARACO	<u>100MG</u>	<u>A078390</u> <u>001</u>	Aug 30, 2007
<u>AB</u>		<u>300MG</u>	<u>A078390</u> <u>002</u>	Aug 30, 2007
<u>AB</u>	IPCA LABS LTD	<u>100MG</u>	<u>A090637</u> <u>001</u>	Mar 16, 2011
<u>AB</u>		<u>300MG</u>	<u>A090637</u> <u>002</u>	Mar 16, 2011
<u>AB</u>	MUTUAL PHARM	<u>100MG</u>	<u>A071449</u> <u>001</u>	Jan 09, 1987
<u>AB</u>		<u>300MG</u>	<u>A071450</u> <u>001</u>	Jan 09, 1987
<u>AB</u>	MYLAN	<u>100MG</u>	<u>N018659</u> <u>001</u>	Oct 24, 1986
<u>AB</u>		<u>300MG</u>	<u>N018659</u> <u>002</u>	Oct 24, 1986
<u>AB</u>	NORTHSTAR HLTHCARE	<u>100MG</u>	<u>A078253</u> <u>001</u>	Sep 11, 2007
<u>AB</u>		<u>300MG</u>	<u>A078253</u> <u>002</u>	Sep 11, 2007
<u>AB</u>	VINTAGE PHARMS	<u>100MG</u>	<u>A075798</u> <u>001</u>	Jun 27, 2003
<u>AB</u>		<u>300MG</u>	<u>A075798</u> <u>002</u>	Jun 27, 2003
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N018832</u> <u>002</u>	Sep 28, 1984
<u>AB</u>		<u>300MG</u>	<u>N018877</u> <u>001</u>	Sep 28, 1984
	<u>LOPURIN</u>			
<u>AB</u>	DR REDDYS LA	<u>100MG</u>	<u>A071586</u> <u>001</u>	Apr 02, 1987
<u>AB</u>		<u>300MG</u>	<u>A071587</u> <u>001</u>	Apr 02, 1987
	<u>ZYLOPRIM</u>			
<u>AB</u>	PROMETHEUS LABS	<u>100MG</u>	<u>N016084</u> <u>001</u>	
<u>AB</u>	+	<u>300MG</u>	<u>N016084</u> <u>002</u>	

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALLOPURINOL SODIUM

<u>AP</u>	BEDFORD LABS	<u>EQ 500MG BASE/VIAL</u>	<u>A076870</u> <u>001</u>	Aug 26, 2004
<u>AP</u>	+ MYLAN INSTITUTIONAL	<u>EQ 500MG BASE/VIAL</u>	<u>N020298</u> <u>001</u>	May 17, 1996

ALMOTRIPTAN MALATE

TABLET; ORAL

AXERT

JANSSEN PHARMS	EQ 6.25MG BASE	N021001 001	May 07, 2001
+	EQ 12.5MG BASE	N021001 002	May 07, 2001

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

LOTRONEX

PROMETHEUS LABS	EQ 0.5MG BASE	N021107 002	Dec 23, 2003
+	EQ 1MG BASE	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	2 IU/ML; 40MG/ML; 12MCG/ML; 40IU/ML; 1MCG/ML; 3MG/ML; 120MCG/ML; 8MG/ML; 1.2MG/ML; 0.72MG/ML; 1.2MG/ML; 660IU/ML; 0.03MG/ML	N021163 001	May 18, 2000
----------	---	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 16 (of 424)

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; IV (INFUSION)

+ SANDOZ	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

+ ROXANE	1MG/ML	A074312 001	Oct 31, 1993
----------	--------	-------------	--------------

TABLET; ORAL

ALPRAZOLAM

AB ACTAVIS ELIZABETH	0.25MG	A074342 001	Oct 31, 1993
AB	0.5MG	A074342 002	Oct 31, 1993
AB	1MG	A074342 003	Oct 31, 1993
AB	2MG	A074342 004	Oct 31, 1993
AB ALPHAPHARM	0.25MG	A074046 001	Oct 19, 1993
AB	0.5MG	A074046 002	Oct 19, 1993
AB	1MG	A074046 003	Oct 19, 1993
AB	2MG	A074046 004	May 07, 1997
AB APOTEX INC	0.25MG	A077741 001	Jan 19, 2007
AB	0.5MG	A077741 002	Jan 19, 2007
AB	1MG	A077741 003	Jan 19, 2007
AB	2MG	A077741 004	Jan 19, 2007
AB BOCA PHARMA	0.25MG	A090248 001	Sep 17, 2010
AB	0.5MG	A090248 002	Sep 17, 2010
AB	1MG	A090248 003	Sep 17, 2010
AB	2MG	A090248 004	Sep 17, 2010
AB DAVA INTL INC	0.25MG	A074174 001	Oct 19, 1993
AB	0.5MG	A074174 002	Oct 19, 1993
AB	1MG	A074174 003	Oct 19, 1993
AB	2MG	A074174 004	Oct 19, 1993
AB MYLAN	0.25MG	A074215 001	Jan 27, 1994
AB	0.5MG	A074215 002	Jan 27, 1994
AB	1MG	A074215 003	Jan 27, 1994
AB	2MG	A074215 004	Jan 27, 1994
AB SANDOZ	0.25MG	A074112 001	Dec 29, 1995
AB	0.5MG	A074112 002	Dec 29, 1995
AB	1MG	A074112 003	Dec 29, 1995
AB	2MG	A074909 001	Mar 25, 1998
AB SUN PHARMA GLOBAL	0.25MG	A090082 001	Jun 17, 2010
AB	0.5MG	A090082 002	Jun 17, 2010
AB	1MG	A090082 003	Jun 17, 2010
AB	2MG	A090082 004	Jun 17, 2010
AB VINTAGE	0.25MG	A078491 001	Sep 25, 2008
AB	0.5MG	A078491 002	Sep 25, 2008
AB	1MG	A078491 003	Sep 25, 2008
AB	2MG	A078491 004	Dec 12, 2008

XANAX

AB PHARMACIA AND UPJOHN	0.25MG	N018276 001	
AB	0.5MG	N018276 002	
AB +	1MG	N018276 003	
AB	2MG	N018276 004	Nov 27, 1985

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

AB ACTAVIS ELIZABETH	0.5MG	A078056 001	Feb 13, 2007
AB	1MG	A078056 002	Feb 13, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 17 (of 424)

ALPRAZOLAM

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>2MG</u>	<u>A078056</u>	<u>003</u>	Feb 13, 2007
<u>AB</u>		<u>3MG</u>	<u>A078056</u>	<u>004</u>	Feb 13, 2007
<u>AB</u>	AMNEAL PHARMS NY	<u>0.5MG</u>	<u>A078387</u>	<u>001</u>	May 30, 2008
<u>AB</u>		<u>1MG</u>	<u>A078387</u>	<u>002</u>	May 30, 2008
<u>AB</u>		<u>2MG</u>	<u>A078387</u>	<u>003</u>	May 30, 2008
<u>AB</u>		<u>3MG</u>	<u>A078387</u>	<u>004</u>	May 30, 2008
<u>AB</u>	ANCHEN PHARMS	<u>0.5MG</u>	<u>A078469</u>	<u>001</u>	Sep 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A078469</u>	<u>002</u>	Sep 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A078469</u>	<u>003</u>	Sep 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A078469</u>	<u>004</u>	Sep 29, 2011
<u>AB</u>	APOTEX INC	<u>0.5MG</u>	<u>A078449</u>	<u>001</u>	Nov 12, 2008
<u>AB</u>		<u>2MG</u>	<u>A078449</u>	<u>002</u>	Nov 12, 2008
<u>AB</u>		<u>3MG</u>	<u>A078449</u>	<u>003</u>	Nov 12, 2008
<u>AB</u>	AUROBINDO PHARMA USA	<u>0.5MG</u>	<u>A090871</u>	<u>001</u>	Jun 07, 2011
<u>AB</u>		<u>1MG</u>	<u>A090871</u>	<u>002</u>	Jun 07, 2011
<u>AB</u>		<u>2MG</u>	<u>A090871</u>	<u>003</u>	Jun 07, 2011
<u>AB</u>		<u>3MG</u>	<u>A090871</u>	<u>004</u>	Jun 07, 2011
<u>AB</u>	BARR	<u>0.5MG</u>	<u>A077725</u>	<u>001</u>	Jul 31, 2006
<u>AB</u>		<u>1MG</u>	<u>A077725</u>	<u>002</u>	Jul 31, 2006
<u>AB</u>		<u>2MG</u>	<u>A077725</u>	<u>004</u>	Jul 31, 2006
<u>AB</u>		<u>3MG</u>	<u>A077725</u>	<u>003</u>	Jul 31, 2006
<u>AB</u>	COREPHARMA	<u>0.5MG</u>	<u>A077996</u>	<u>001</u>	Jan 31, 2007
<u>AB</u>		<u>1MG</u>	<u>A077996</u>	<u>002</u>	Jan 31, 2007
<u>AB</u>		<u>2MG</u>	<u>A077996</u>	<u>003</u>	Jan 31, 2007
<u>AB</u>		<u>3MG</u>	<u>A077996</u>	<u>004</u>	Jan 31, 2007
<u>AB</u>	IMPAX LABS	<u>0.5MG</u>	<u>A077968</u>	<u>004</u>	May 24, 2007
<u>AB</u>		<u>1MG</u>	<u>A077968</u>	<u>003</u>	May 24, 2007
<u>AB</u>		<u>2MG</u>	<u>A077968</u>	<u>002</u>	May 24, 2007
<u>AB</u>		<u>3MG</u>	<u>A077968</u>	<u>001</u>	May 24, 2007
<u>AB</u>	MYLAN	<u>0.5MG</u>	<u>A077391</u>	<u>002</u>	Jan 26, 2006
<u>AB</u>		<u>1MG</u>	<u>A077391</u>	<u>003</u>	Jan 26, 2006
<u>AB</u>		<u>2MG</u>	<u>A077391</u>	<u>004</u>	Jan 26, 2006
<u>AB</u>		<u>3MG</u>	<u>A077391</u>	<u>001</u>	Jan 26, 2006
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A077777</u>	<u>001</u>	Jun 30, 2006
<u>AB</u>		<u>1MG</u>	<u>A077777</u>	<u>002</u>	Jun 30, 2006
<u>AB</u>		<u>2MG</u>	<u>A077777</u>	<u>003</u>	Jun 30, 2006
<u>AB</u>		<u>3MG</u>	<u>A077777</u>	<u>004</u>	Jun 30, 2006
<u>AB</u>	TEVA PHARMS	<u>0.5MG</u>	<u>A077979</u>	<u>001</u>	Feb 28, 2007
<u>AB</u>		<u>1MG</u>	<u>A077979</u>	<u>002</u>	Feb 28, 2007
<u>AB</u>		<u>2MG</u>	<u>A077979</u>	<u>003</u>	Feb 28, 2007
<u>AB</u>		<u>3MG</u>	<u>A077979</u>	<u>004</u>	Feb 28, 2007
<u>AB</u>	VINTAGE	<u>0.5MG</u>	<u>A078442</u>	<u>001</u>	Oct 15, 2007
<u>AB</u>		<u>1MG</u>	<u>A078442</u>	<u>002</u>	Oct 15, 2007
<u>AB</u>		<u>2MG</u>	<u>A078442</u>	<u>003</u>	Oct 15, 2007
<u>AB</u>		<u>3MG</u>	<u>A078442</u>	<u>004</u>	Oct 15, 2007
<u>AB</u>	WATSON LABS FLORIDA	<u>0.5MG</u>	<u>A077198</u>	<u>001</u>	May 13, 2010
<u>AB</u>		<u>1MG</u>	<u>A077198</u>	<u>002</u>	May 13, 2010
<u>AB</u>		<u>2MG</u>	<u>A077198</u>	<u>003</u>	May 13, 2010
<u>AB</u>		<u>3MG</u>	<u>A077198</u>	<u>004</u>	May 13, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.5MG</u>	<u>A078489</u>	<u>001</u>	Oct 17, 2008
<u>AB</u>		<u>1MG</u>	<u>A078489</u>	<u>002</u>	Oct 17, 2008
<u>AB</u>		<u>2MG</u>	<u>A078489</u>	<u>003</u>	Oct 17, 2008
<u>AB</u>		<u>3MG</u>	<u>A078489</u>	<u>004</u>	Oct 17, 2008
<u>XANAX XR</u>					
<u>AB</u>	PHARMACIA AND UPJOHN	<u>0.5MG</u>	<u>N021434</u>	<u>001</u>	Jan 17, 2003
<u>AB</u>		<u>1MG</u>	<u>N021434</u>	<u>002</u>	Jan 17, 2003
<u>AB</u>		<u>2MG</u>	<u>N021434</u>	<u>003</u>	Jan 17, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 18 (of 424)

ALPRAZOLAM

TABLET, EXTENDED RELEASE; ORAL

XANAX XRAB + PHARMACIA AND UPJOHN 3MG N021434 004 Jan 17, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

ALPRAZOLAM

AB	ACTAVIS ELIZABETH	<u>0.25MG</u>	A078561 001	Mar 16, 2010
AB		<u>0.5MG</u>	A078561 002	Mar 16, 2010
AB		<u>1MG</u>	A078561 003	Mar 16, 2010
AB		<u>2MG</u>	A078561 004	Mar 16, 2010
AB	PAR PHARM	<u>0.25MG</u>	A078088 001	Jan 09, 2009
AB		<u>0.5MG</u>	A078088 002	Jan 09, 2009
AB		<u>1MG</u>	A078088 003	Jan 09, 2009
AB		<u>2MG</u>	A078088 004	Jan 09, 2009

NIRAVAM

AB	SCHWARZ PHARMA	<u>0.25MG</u>	N021726 001	Jan 19, 2005
AB		<u>0.5MG</u>	N021726 002	Jan 19, 2005
AB	+	<u>1MG</u>	N021726 003	Jan 19, 2005
AB		<u>2MG</u>	N021726 004	Jan 19, 2005

ALPROSTADIL

INJECTABLE; INJECTION

ALPROSTADIL

AP	BEDFORD	<u>0.5MG/ML</u>	A074815 001	Jan 20, 1998
AP	TEVA PARENTERAL	<u>0.5MG/ML</u>	A075196 001	Apr 30, 1999

CAVERJECT

AP	PHARMACIA AND UPJOHN	<u>0.01MG/VIAL</u>	N020379 001	Jul 06, 1995
AP	+	<u>0.02MG/VIAL</u>	N020379 002	Jul 06, 1995
AP	+	<u>0.04MG/VIAL</u>	N020379 004	May 19, 1997

EDEX

AP	SCHWARZ PHARMA	<u>0.01MG/VIAL</u>	N020649 002	Jun 12, 1997
AP		<u>0.02MG/VIAL</u>	N020649 003	Jun 12, 1997
AP	+	<u>0.04MG/VIAL</u>	N020649 004	Jun 12, 1997

PROSTIN VR PEDIATRIC

AP	+ PHARMACIA AND UPJOHN	<u>0.5MG/ML</u>	N018484 001	
----	------------------------	-----------------	-------------	--

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

+ SCHWARZ PHARMA	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

MEDA PHARMS	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
-------------	------	-------------	--------------

ALVIMOPAN

CAPSULE; ORAL

ENTEREG

+ ADOLOR	12MG	N021775 001	May 20, 2008
----------	------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 19 (of 424)

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u>	BANNER PHARMACAPS	<u>100MG</u>	<u>A078720</u>	<u>001</u>	May 29, 2008
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A071293</u>	<u>001</u>	Feb 18, 1987
<u>AB</u> + USL PHARMA		<u>100MG</u>	<u>A070589</u>	<u>001</u>	Aug 05, 1986

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

<u>AA</u> + CAROLINA MEDCL		<u>50MG/5ML</u>	<u>A075819</u>	<u>001</u>	Sep 11, 2002
<u>AA</u> + HI TECH PHARMA		<u>50MG/5ML</u>	<u>A074170</u>	<u>001</u>	Oct 28, 1994
<u>AA</u> + MIKART		<u>50MG/5ML</u>	<u>A074028</u>	<u>001</u>	Jun 28, 1993
<u>AA</u> + PHARM ASSOC		<u>50MG/5ML</u>	<u>A074509</u>	<u>001</u>	Jul 17, 1995
<u>AA</u> + SILARX		<u>50MG/5ML</u>	<u>A076352</u>	<u>001</u>	Sep 10, 2004
<u>AA</u> + VINTAGE		<u>50MG/5ML</u>	<u>A077992</u>	<u>001</u>	Dec 12, 2006
<u>AA</u> + WOCKHARDT		<u>50MG/5ML</u>	<u>A075060</u>	<u>001</u>	Dec 24, 1998

TABLET; ORAL

AMANTADINE HYDROCHLORIDE

+ USL PHARMA		100MG	A076186	001	Dec 16, 2002
--------------	--	-------	---------	-----	--------------

AMBENONIUM CHLORIDE

TABLET; ORAL

MYTELASE

+ SANOFI AVENTIS US		10MG	N010155	002	
---------------------	--	------	---------	-----	--

AMBRISENTAN

TABLET; ORAL

LETAIRIS

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

AMCINONIDE

CREAM; TOPICAL

AMCINONIDE

<u>AB</u> + ALTANA		<u>0.1%</u>	<u>A076065</u>	<u>001</u>	May 15, 2003
<u>AB</u> TARO PHARM IND		<u>0.1%</u>	<u>A076229</u>	<u>001</u>	May 31, 2002

LOTION; TOPICAL

AMCINONIDE

+ ALTANA		0.1%	A076329	001	Nov 06, 2002
----------	--	------	---------	-----	--------------

OINTMENT; TOPICAL

AMCINONIDE

<u>AB</u> + ALTANA		<u>0.1%</u>	<u>A076096</u>	<u>001</u>	Nov 19, 2002
<u>AB</u> TARO PHARM IND		<u>0.1%</u>	<u>A076367</u>	<u>001</u>	Mar 19, 2003

AMIFOSTINE

INJECTABLE; INJECTION

AMIFOSTINE

<u>AP</u> SUN PHARMA GLOBAL		<u>500MG/VIAL</u>	<u>A077126</u>	<u>001</u>	Mar 14, 2008
<u>AP</u> + ETHYOL					

<u>AP</u> + MEDIIMMUNE		<u>500MG/VIAL</u>	<u>N020221</u>	<u>001</u>	Dec 08, 1995
------------------------	--	-------------------	----------------	------------	--------------

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

<u>AP</u> + BEDFORD		<u>EQ 50MG BASE/ML</u>	<u>A063313</u>	<u>001</u>	Apr 11, 1994
<u>AP</u> +		<u>EQ 250MG BASE/ML</u>	<u>A063315</u>	<u>001</u>	Apr 11, 1994
<u>AP</u> HOSPIRA		<u>EQ 50MG BASE/ML</u>	<u>A063263</u>	<u>001</u>	Nov 30, 1994
<u>AP</u>		<u>EQ 250MG BASE/ML</u>	<u>A063264</u>	<u>001</u>	Nov 30, 1994
<u>AP</u> TEVA PARENTERAL		<u>EQ 250MG BASE/ML</u>	<u>A064045</u>	<u>002</u>	Sep 28, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 20 (of 424)

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

<u>AB</u>	+ PAR PHARM	<u>5MG</u>	<u>A070346 001</u>	Jan 22, 1986
<u>AB</u>	SIGMAPHARM LABS LLC	<u>5MG</u>	<u>A079133 001</u>	Jan 30, 2009
	<u>MIDAMOR</u>			
<u>AB</u>	PADDOCK LLC	<u>5MG</u>	<u>N018200 001</u>	

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	BARR	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>A071111 001</u>	May 10, 1988
<u>AB</u>	+ MYLAN	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>A073209 001</u>	Oct 31, 1991

AMINO ACIDS

INJECTABLE; INJECTION

AMINOSYN 10%

HOSPIRA 10% (10GM/100ML) N017673 003

AMINOSYN 10% (PH6)

HOSPIRA 10% (10GM/100ML) N017673 008 Nov 18, 1985

AMINOSYN 3.5%

HOSPIRA 3.5% (3.5GM/100ML) N017789 004

AMINOSYN 5%

HOSPIRA 5% (5GM/100ML) N017673 001

AMINOSYN 7%

HOSPIRA 7% (7GM/100ML) N017673 002

AMINOSYN 7% (PH6)

HOSPIRA 7% (7GM/100ML) N017673 006 Nov 18, 1985

AMINOSYN 8.5%

HOSPIRA 8.5% (8.5GM/100ML) N017673 004

AMINOSYN 8.5% (PH6)

HOSPIRA 8.5% (8.5GM/100ML) N017673 007 Nov 18, 1985

AMINOSYN II 10%

HOSPIRA 10% (10GM/100ML) N019438 005 Apr 03, 1986

AMINOSYN II 10% IN PLASTIC CONTAINER

HOSPIRA 10% (10GM/100ML) N020015 001 Dec 19, 1991

AMINOSYN II 15% IN PLASTIC CONTAINER

HOSPIRA 15% (15GM/100ML) N020041 001 Dec 19, 1991

AMINOSYN II 7%

HOSPIRA 7% (7GM/100ML) N019438 003 Apr 03, 1986

AMINOSYN II 8.5%

HOSPIRA 8.5% (8.5GM/100ML) N019438 004 Apr 03, 1986

AMINOSYN-HBC 7%

HOSPIRA 7% (7GM/100ML) N019374 001 Jul 12, 1985

AMINOSYN-HF 8%

HOSPIRA 8% (8GM/100ML) N020345 001 Apr 04, 1996

AMINOSYN-PF 10%

HOSPIRA 10% (10GM/100ML) N019492 002 Oct 17, 1986

AMINOSYN-PF 7%

HOSPIRA 7% (7GM/100ML) N019398 001 Sep 06, 1985

AMINOSYN-RF 5.2%

HOSPIRA 5.2% (5.2GM/100ML) N018429 001

BRANCHAMIN 4% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4% (4GM/100ML) N018684 001 Sep 28, 1984

CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER

CLINTEC NUTR 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

PRESCRIPTION DRUG PRODUCT LIST

3 - 21 (of 424)

AMINO ACIDS

INJECTABLE; INJECTION				
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
HEPATASOL 8%				
BAXTER HLTHCARE	8% (8GM/100ML)	A020360	001	Apr 04, 1996
NEPHRAMINE 5.4%				
B BRAUN	5.4% (5.4GM/100ML)	N017766	001	
PREMASOL 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	10% (10GM/100ML)	A075880	002	Jun 19, 2003
PREMASOL 6% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	6% (6GM/100ML)	A075880	001	Jun 19, 2003
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	20% (20GM/100ML)	N020849	001	Aug 26, 1998
RENAMIN W/O ELECTROLYTES				
BAXTER HLTHCARE	6.5% (6.5GM/100ML)	N017493	007	Oct 15, 1982
TRAVASOL 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	10% (10MG/100ML)	N018931	003	Aug 23, 1984
TRAVASOL 10% W/O ELECTROLYTES				
BAXTER HLTHCARE	10% (10GM/100ML)	N017493	006	
TRAVASOL 5.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5.5% (5.5GM/100ML)	N018931	001	Aug 23, 1984
TRAVASOL 5.5% W/O ELECTROLYTES				
BAXTER HLTHCARE	5.5% (5.5GM/100ML)	N017493	004	
TRAVASOL 8.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	8.5% (8.5GM/100ML)	N018931	002	Aug 23, 1984
TRAVASOL 8.5% W/O ELECTROLYTES				
BAXTER HLTHCARE	8.5% (8.5GM/100ML)	N017493	005	
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/10 OML	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;2 61MG/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

PRESCRIPTION DRUG PRODUCT LIST

3 - 22 (of 424)

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

INJECTABLE; INJECTION

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;2 61MG/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;261 MG/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;261 MG/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;261 MG/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;261 MG/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;261 MG/100ML;340MG/100ML;59MG/100ML	N020678 021	Mar 26, 1997	

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	

PRESCRIPTION DRUG PRODUCT LIST

3 - 23 (of 424)

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

INJECTABLE; INJECTION

AMINOSYN 3.5% M

HOSPIRA 3.5%;21MG/100ML;40MG/100ML;128MG/100ML; N017789 003
234MG/100MLAMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE;
SODIUM CHLORIDE

INJECTABLE; INJECTION

FREAMEINE III 3% W/ ELECTROLYTES

B BRAUN 3%;54MG/100ML;40MG/100ML;150MG/100ML;20 N016822 003
0MG/100ML;120MG/100MLAMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC;
SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 10% W/ ELECTROLYTES

HOSPIRA 10%;102MG/100ML;45MG/100ML;522MG/100ML; N019437 004 Apr 03, 1986
410MG/100ML

AMINOSYN II 8.5% W/ ELECTROLYTES

HOSPIRA 8.5%;102MG/100ML;45MG/100ML;522MG/100ML N019437 005 Apr 03, 1986
;410MG/100MLAMINO 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/100MLAMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 7% W/ ELECTROLYTES

HOSPIRA 7%;102MG/100ML;522MG/100ML;410MG/100ML N017789 002

AMINOSYN 8.5% W/ ELECTROLYTES

HOSPIRA 8.5%;102MG/100ML;522MG/100ML;410MG/100M N017673 005
LAMINOCAPROIC 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 + XANODYNE PHARM 1.25GM/5ML N015230 002**

PRESCRIPTION DRUG PRODUCT LIST

3 - 24 (of 424)

AMINOCAPROIC ACID

SYRUP; ORAL

AMINOCAPROIC ACID

<u>AA</u>	MIKART	<u>1.25GM/5ML</u>	<u>A074759 001</u>	Sep 02, 1998
TABLET; ORAL				
	<u>AMICAR</u>			
<u>AB</u>	XANODYNE PHARM	<u>500MG</u>	<u>N015197 001</u>	
	<u>AMINOCAPROIC</u>			
<u>AB</u>	MIKART	<u>500MG</u>	<u>A075602 001</u>	May 24, 2001
AMICAR				
+ XANODYNE PHARM	1GM		N015197 002	Jun 24, 2004

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION

AMINOHIPPURATE SODIUM

+ MERCK 20%

N005619 001

AMINOLEVULINIC ACID HYDROCHLORIDE

SOLUTION; TOPICAL

LEVULAN

+ DUSA 20%

N020965 001 Dec 03, 1999

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

<u>AP</u> + HOSPIRA	<u>25MG/ML</u>	<u>A087242 001</u>	Oct 26, 1983
<u>AP</u> INTL MEDICATION	<u>25MG/ML</u>	<u>A087209 001</u>	Feb 01, 1982
<u>AP</u> LUITPOLD	<u>25MG/ML</u>	<u>A087600 001</u>	
<u>AP</u> PHARMA SERVE NY	<u>25MG/ML</u>	<u>A087392 001</u>	Dec 15, 1983
TABLET; ORAL			
AMINOPHYLLINE			
+ WEST WARD	100MG	A084540 001	
+	200MG	A085003 001	

AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER

+ JACOBUS 4GM/PACKET

A074346 001 Jun 30, 1994

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

<u>AP</u> + AKORN	<u>50MG/ML</u>	<u>A076232 001</u>	Jul 05, 2006
<u>AP</u> + APP PHARMS	<u>50MG/ML</u>	<u>A075761 001</u>	Oct 15, 2002
<u>AP</u> + BEDFORD	<u>50MG/ML</u>	<u>A076018 001</u>	Oct 15, 2002
<u>AP</u> + BEDFORD LABS	<u>50MG/ML</u>	<u>A076299 001</u>	Oct 24, 2002
<u>AP</u> + BIONICHE PHARMA	<u>50MG/ML</u>	<u>A076217 001</u>	Oct 15, 2002
<u>AP</u> + CLARIS LIFESCIENCES	<u>50MG/ML</u>	<u>A076394 001</u>	Apr 25, 2003
<u>AP</u> + GLAND PHARMA LTD	<u>50MG/ML</u>	<u>A077161 001</u>	Apr 20, 2005
<u>AP</u> HIKMA FARMACEUTICA	<u>50MG/ML</u>	<u>A077234 001</u>	Feb 25, 2008
<u>AP</u> + HOSPIRA	<u>50MG/ML</u>	<u>A075955 001</u>	Oct 18, 2002
<u>AP</u> + TEVA PARENTERAL	<u>50MG/ML</u>	<u>A076163 001</u>	Sep 05, 2003
<u>AP</u> WOCKHARDT	<u>50MG/ML</u>	<u>A077610 001</u>	Oct 30, 2008
<u>AP</u>	<u>50MG/ML</u>	<u>A077834 001</u>	Oct 30, 2008
NEXTERONE			
<u>AP</u> BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N022325 001</u>	Dec 24, 2008
NEXTERONE			
+ BAXTER HLTHCARE	150MG/100ML (1.5MG/ML)	N022325 002	Nov 16, 2010
+	360MG/200ML (1.8MG/ML)	N022325 003	Nov 16, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 25 (of 424)

AMIODARONE HYDROCHLORIDE

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

<u>AB</u>	APOTEX CORP	<u>200MG</u>	<u>A078578</u>	<u>001</u>	Nov 06, 2008
<u>AB</u>	AUROSAL PHARMS	<u>200MG</u>	<u>A077069</u>	<u>001</u>	Apr 08, 2005
<u>AB</u>		<u>400MG</u>	<u>A077069</u>	<u>002</u>	Apr 08, 2005
<u>AB</u>	BARR	<u>200MG</u>	<u>A075389</u>	<u>001</u>	Jan 25, 2001
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A075188</u>	<u>001</u>	Feb 24, 1999
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A075315</u>	<u>001</u>	Dec 23, 1998
<u>AB</u>		<u>400MG</u>	<u>A075315</u>	<u>002</u>	Jun 30, 2000
<u>AB</u>	TARO	<u>100MG</u>	<u>A075424</u>	<u>002</u>	Dec 18, 2002
<u>AB</u>		<u>200MG</u>	<u>A075424</u>	<u>001</u>	Mar 30, 2001
<u>AB</u>		<u>400MG</u>	<u>A076362</u>	<u>001</u>	Nov 29, 2002
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>A074739</u>	<u>001</u>	Nov 30, 1998
<u>AB</u>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A079029</u>	<u>001</u>	Sep 16, 2008
	<u>CORDARONE</u>				
<u>AB</u>	+ WYETH PHARMS INC	<u>200MG</u>	<u>N018972</u>	<u>001</u>	Dec 24, 1985
	<u>PACERONE</u>				
<u>AB</u>	UPSHER SMITH	<u>100MG</u>	<u>A075135</u>	<u>002</u>	Apr 12, 2005
<u>AB</u>		<u>200MG</u>	<u>A075135</u>	<u>001</u>	Apr 30, 1998
	AMIODARONE HYDROCHLORIDE				
	TARO	300MG	A076362	002	Dec 02, 2003

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	CARACO	<u>10MG</u>	<u>A040815</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>		<u>25MG</u>	<u>A040816</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>		<u>50MG</u>	<u>A040817</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>		<u>75MG</u>	<u>A040818</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>		<u>100MG</u>	<u>A040819</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>		<u>150MG</u>	<u>A040820</u>	<u>001</u>	Jun 27, 2008
<u>AB</u>	MUTUAL PHARM	<u>10MG</u>	<u>A089398</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>		<u>25MG</u>	<u>A089399</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>		<u>50MG</u>	<u>A089400</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>		<u>75MG</u>	<u>A089401</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>		<u>100MG</u>	<u>A089402</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>		<u>150MG</u>	<u>A089403</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A086009</u>	<u>002</u>	
<u>AB</u>		<u>25MG</u>	<u>A086009</u>	<u>003</u>	
<u>AB</u>		<u>50MG</u>	<u>A086009</u>	<u>001</u>	
<u>AB</u>		<u>75MG</u>	<u>A086009</u>	<u>004</u>	
<u>AB</u>		<u>100MG</u>	<u>A086009</u>	<u>005</u>	
<u>AB</u>		<u>150MG</u>	<u>A086009</u>	<u>006</u>	
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A085969</u>	<u>001</u>	
<u>AB</u>	+	<u>25MG</u>	<u>A085966</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A085968</u>	<u>001</u>	
<u>AB</u>		<u>75MG</u>	<u>A085971</u>	<u>001</u>	
<u>AB</u>		<u>100MG</u>	<u>A085967</u>	<u>001</u>	
<u>AB</u>		<u>150MG</u>	<u>A085970</u>	<u>001</u>	
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A040218</u>	<u>001</u>	Sep 11, 1997
<u>AB</u>		<u>25MG</u>	<u>A040218</u>	<u>002</u>	Sep 11, 1997
<u>AB</u>		<u>50MG</u>	<u>A040218</u>	<u>003</u>	Sep 11, 1997
<u>AB</u>		<u>75MG</u>	<u>A040218</u>	<u>004</u>	Sep 11, 1997
<u>AB</u>		<u>100MG</u>	<u>A040218</u>	<u>005</u>	Sep 11, 1997
<u>AB</u>		<u>150MG</u>	<u>A040218</u>	<u>006</u>	Sep 11, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 26 (of 424)

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>EQ 12.5MG BASE;5MG</u>	<u>A071297</u> <u>002</u>	Dec 10, 1986
<u>AB</u>		<u>EQ 25MG BASE;10MG</u>	<u>A071297</u> <u>001</u>	Dec 10, 1986
	<u>LIMBITROL</u>			
<u>AB</u>	VALEANT PHARM INTL	<u>EQ 12.5MG BASE;5MG</u>	<u>N016949</u> <u>001</u>	
	<u>LIMBITROL DS</u>			
<u>AB</u>	+ VALEANT PHARM INTL	<u>EQ 25MG BASE;10MG</u>	<u>N016949</u> <u>002</u>	

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

MYLAN	10MG;2MG	<u>A071443</u> <u>002</u>	Nov 10, 1988
	10MG;4MG	<u>A071443</u> <u>003</u>	Nov 10, 1988
+	25MG;2MG	<u>A071443</u> <u>004</u>	Nov 10, 1988
+	25MG;4MG	<u>A071443</u> <u>005</u>	Nov 10, 1988
+	50MG;4MG	<u>A071443</u> <u>001</u>	Nov 10, 1988

AMLEXANOX

PASTE; DENTAL

APHTHASOL

+	ULURU	5%	<u>N020511</u> <u>001</u>	Dec 17, 1996
---	-------	----	---------------------------	--------------

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

<u>AB</u>	ALKEM	<u>EQ 2.5MG BASE</u>	<u>A078925</u> <u>001</u>	May 04, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078925</u> <u>002</u>	May 04, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078925</u> <u>003</u>	May 04, 2009
<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 2.5MG BASE</u>	<u>A078477</u> <u>001</u>	Jan 16, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078477</u> <u>002</u>	Jan 16, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078477</u> <u>003</u>	Jan 16, 2008
<u>AB</u>	APOTEX	<u>EQ 2.5MG BASE</u>	<u>A076719</u> <u>001</u>	May 23, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076719</u> <u>002</u>	May 23, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076719</u> <u>003</u>	May 23, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 2.5MG BASE</u>	<u>A078021</u> <u>001</u>	Jul 17, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078021</u> <u>002</u>	Jul 17, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078021</u> <u>003</u>	Jul 17, 2007
<u>AB</u>	CARACO	<u>EQ 2.5MG BASE</u>	<u>A078231</u> <u>001</u>	Nov 30, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078231</u> <u>002</u>	Nov 30, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078231</u> <u>003</u>	Nov 30, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 2.5MG BASE</u>	<u>A076692</u> <u>001</u>	Jul 20, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076692</u> <u>002</u>	Jul 20, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076692</u> <u>003</u>	Jul 20, 2007
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 2.5MG BASE</u>	<u>A078552</u> <u>001</u>	Apr 08, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078552</u> <u>002</u>	Apr 08, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078552</u> <u>003</u>	Apr 08, 2009
<u>AB</u>	HIKMA PHARMS	<u>EQ 2.5MG BASE</u>	<u>A077771</u> <u>001</u>	Apr 12, 2011
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077771</u> <u>002</u>	Apr 12, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077771</u> <u>003</u>	Apr 12, 2011
<u>AB</u>	INVAGEN PHARMS	<u>EQ 2.5MG BASE</u>	<u>A077955</u> <u>001</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077955</u> <u>002</u>	Aug 28, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077955</u> <u>003</u>	Aug 28, 2007
<u>AB</u>	LEK PHARMS DD	<u>EQ 2.5MG BASE</u>	<u>A076859</u> <u>001</u>	Sep 10, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A076859</u> <u>002</u>	Sep 10, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076859</u> <u>003</u>	Sep 10, 2007
<u>AB</u>	LUPIN	<u>EQ 2.5MG BASE</u>	<u>A078043</u> <u>001</u>	Jul 12, 2007
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078043</u> <u>002</u>	Jul 12, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078043</u> <u>003</u>	Jul 12, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 27 (of 424)

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

AB	MATRIX LABS LTD	<u>EQ 2.5MG BASE</u>	<u>A078224</u> <u>001</u>	Feb 27, 2008
AB		<u>EQ 5MG BASE</u>	<u>A078224</u> <u>002</u>	Feb 27, 2008
AB		<u>EQ 10MG BASE</u>	<u>A078224</u> <u>003</u>	Feb 27, 2008
AB	MYLAN	<u>EQ 2.5MG BASE</u>	<u>A076418</u> <u>001</u>	Oct 03, 2005
AB		<u>EQ 5MG BASE</u>	<u>A076418</u> <u>002</u>	Oct 03, 2005
AB		<u>EQ 10MG BASE</u>	<u>A076418</u> <u>003</u>	Oct 03, 2005
AB	ORCHID HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>A078453</u> <u>001</u>	Jul 02, 2009
AB		<u>EQ 5MG BASE</u>	<u>A078453</u> <u>002</u>	Jul 02, 2009
AB		<u>EQ 10MG BASE</u>	<u>A078453</u> <u>003</u>	Jul 02, 2009
AB	PURACAP PHARM	<u>EQ 2.5MG BASE</u>	<u>A078131</u> <u>001</u>	Sep 04, 2007
AB		<u>EQ 5MG BASE</u>	<u>A078131</u> <u>002</u>	Sep 04, 2007
AB		<u>EQ 10MG BASE</u>	<u>A078131</u> <u>003</u>	Sep 04, 2007
AB	RANBAXY	<u>EQ 2.5MG BASE</u>	<u>A077974</u> <u>001</u>	Jul 09, 2007
AB		<u>EQ 5MG BASE</u>	<u>A077974</u> <u>002</u>	Jul 09, 2007
AB		<u>EQ 10MG BASE</u>	<u>A077974</u> <u>003</u>	Jul 09, 2007
AB	ROXANE	<u>EQ 2.5MG BASE</u>	<u>A077262</u> <u>001</u>	Jul 09, 2007
AB		<u>EQ 5MG BASE</u>	<u>A077262</u> <u>002</u>	Jul 09, 2007
AB		<u>EQ 10MG BASE</u>	<u>A077262</u> <u>003</u>	Jul 09, 2007
AB	SECAN PHARMS	<u>EQ 5MG BASE</u>	<u>A090752</u> <u>001</u>	Apr 15, 2011
AB		<u>EQ 10MG BASE</u>	<u>A090752</u> <u>002</u>	Apr 15, 2011
AB	TEVA	<u>EQ 2.5MG BASE</u>	<u>A076846</u> <u>001</u>	Jun 28, 2007
AB		<u>EQ 5MG BASE</u>	<u>A076846</u> <u>002</u>	Jun 28, 2007
AB		<u>EQ 10MG BASE</u>	<u>A076846</u> <u>003</u>	Jun 28, 2007
AB	TORRENT PHARMS	<u>EQ 2.5MG BASE</u>	<u>A078573</u> <u>001</u>	Sep 22, 2008
AB		<u>EQ 5MG BASE</u>	<u>A078573</u> <u>002</u>	Sep 22, 2008
AB		<u>EQ 10MG BASE</u>	<u>A078573</u> <u>003</u>	Sep 22, 2008
AB	UPSHER SMITH	<u>EQ 2.5MG BASE</u>	<u>A077759</u> <u>001</u>	Jul 09, 2007
AB		<u>EQ 5MG BASE</u>	<u>A077759</u> <u>002</u>	Jul 09, 2007
AB		<u>EQ 10MG BASE</u>	<u>A077759</u> <u>003</u>	Jul 09, 2007
AB	VINTAGE	<u>EQ 2.5MG BASE</u>	<u>A078414</u> <u>001</u>	Apr 07, 2010
AB		<u>EQ 5MG BASE</u>	<u>A078414</u> <u>002</u>	Apr 07, 2010
AB		<u>EQ 10MG BASE</u>	<u>A078414</u> <u>003</u>	Apr 07, 2010
AB	WATSON LABS	<u>EQ 2.5MG BASE</u>	<u>A077073</u> <u>001</u>	Sep 26, 2007
AB		<u>EQ 2.5MG BASE</u>	<u>A077671</u> <u>001</u>	Jul 19, 2007
AB		<u>EQ 5MG BASE</u>	<u>A077073</u> <u>002</u>	Sep 26, 2007
AB		<u>EQ 5MG BASE</u>	<u>A077671</u> <u>002</u>	Jul 19, 2007
AB		<u>EQ 10MG BASE</u>	<u>A077073</u> <u>003</u>	Sep 26, 2007
AB		<u>EQ 10MG BASE</u>	<u>A077671</u> <u>003</u>	Jul 19, 2007
AB	WOCKHARDT	<u>EQ 2.5MG BASE</u>	<u>A078500</u> <u>001</u>	Sep 06, 2007
AB		<u>EQ 5MG BASE</u>	<u>A078500</u> <u>002</u>	Sep 06, 2007
AB		<u>EQ 10MG BASE</u>	<u>A078500</u> <u>003</u>	Sep 06, 2007
AB	WORLD GEN	<u>EQ 2.5MG BASE</u>	<u>A077516</u> <u>001</u>	Jul 11, 2007
AB		<u>EQ 5MG BASE</u>	<u>A077516</u> <u>002</u>	Jul 11, 2007
AB		<u>EQ 10MG BASE</u>	<u>A077516</u> <u>003</u>	Jul 11, 2007
AB	ZYDUS PHARMS USA	<u>EQ 2.5MG BASE</u>	<u>A078226</u> <u>001</u>	Jul 09, 2007
AB		<u>EQ 5MG BASE</u>	<u>A078226</u> <u>002</u>	Jul 09, 2007
AB		<u>EQ 10MG BASE</u>	<u>A078226</u> <u>003</u>	Jul 09, 2007
<u>NORVASC</u>				
AB	PFIZER	<u>EQ 2.5MG BASE</u>	<u>N019787</u> <u>001</u>	Jul 31, 1992
AB		<u>EQ 5MG BASE</u>	<u>N019787</u> <u>002</u>	Jul 31, 1992
AB	+	<u>EQ 10MG BASE</u>	<u>N019787</u> <u>003</u>	Jul 31, 1992

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

CADUET

PFIZER

EQ 2.5MG BASE;EQ 10MG BASE
EQ 2.5MG BASE;EQ 20MG BASEN021540 009 Jul 29, 2004
N021540 010 Jul 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 28 (of 424)

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL
CADUET

PFIZER	EQ 2.5MG BASE;EQ 40MG BASE	N021540 011	Jul 29, 2004
	EQ 5MG BASE;EQ 10MG BASE	N021540 001	Jan 30, 2004
	EQ 5MG BASE;EQ 20MG BASE	N021540 002	Jan 30, 2004
	EQ 5MG BASE;EQ 40MG BASE	N021540 003	Jan 30, 2004
	EQ 5MG BASE;EQ 80MG BASE	N021540 004	Jan 30, 2004
	EQ 10MG BASE;EQ 10MG BASE	N021540 005	Jan 30, 2004
	EQ 10MG BASE;EQ 20MG BASE	N021540 006	Jan 30, 2004
	EQ 10MG BASE;EQ 40MG BASE	N021540 007	Jan 30, 2004
+	EQ 10MG BASE;EQ 80MG BASE	N021540 008	Jan 30, 2004

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

AB	DR REDDYS LABS INC	<u>EQ 2.5MG BASE;10MG</u>	A077183 001	Apr 15, 2010
AB		<u>EQ 5MG BASE;10MG</u>	A077183 002	Apr 15, 2010
AB		<u>EQ 5MG BASE;20MG</u>	A077183 003	Apr 15, 2010
AB		<u>EQ 5MG BASE;40MG</u>	A090149 001	Jul 05, 2011
AB		<u>EQ 10MG BASE;20MG</u>	A077183 004	Apr 15, 2010
AB		<u>EQ 10MG BASE;40MG</u>	A090149 002	Jul 05, 2011
AB	LUPIN PHARMS	<u>EQ 2.5MG BASE;10MG</u>	A078466 001	Feb 05, 2010
AB		<u>EQ 5MG BASE;10MG</u>	A078466 002	Feb 05, 2010
AB		<u>EQ 5MG BASE;20MG</u>	A078466 003	Feb 05, 2010
AB		<u>EQ 5MG BASE;40MG</u>	A078466 005	Jul 05, 2011
AB		<u>EQ 10MG BASE;20MG</u>	A078466 004	Feb 05, 2010
AB		<u>EQ 10MG BASE;40MG</u>	A078466 006	Jul 05, 2011
AB	MYLAN	<u>EQ 2.5MG BASE;10MG</u>	A077375 001	May 21, 2010
AB		<u>EQ 5MG BASE;10MG</u>	A077375 002	May 21, 2010
AB		<u>EQ 5MG BASE;20MG</u>	A077375 003	May 21, 2010
AB		<u>EQ 5MG BASE;40MG</u>	A079047 001	Jul 05, 2011
AB		<u>EQ 10MG BASE;20MG</u>	A077375 004	May 21, 2010
AB		<u>EQ 10MG BASE;40MG</u>	A079047 002	Jul 05, 2011
AB	PAR PHARM	<u>EQ 2.5MG BASE;10MG</u>	A078381 001	Jul 29, 2010
AB		<u>EQ 5MG BASE;10MG</u>	A078381 002	Jul 29, 2010
AB		<u>EQ 5MG BASE;20MG</u>	A078381 003	Jul 29, 2010
AB		<u>EQ 5MG BASE;40MG</u>	A078381 005	Jul 29, 2010
AB		<u>EQ 10MG BASE;20MG</u>	A078381 004	Jul 29, 2010
AB		<u>EQ 10MG BASE;40MG</u>	A078381 006	Jul 29, 2010
AB	TEVA PHARMS	<u>EQ 2.5MG BASE;10MG</u>	A077179 001	May 18, 2007
AB		<u>EQ 5MG BASE;10MG</u>	A077179 002	May 18, 2007
AB		<u>EQ 5MG BASE;20MG</u>	A077179 003	May 18, 2007
AB		<u>EQ 5MG BASE;40MG</u>	A077179 005	Jul 05, 2011
AB		<u>EQ 10MG BASE;20MG</u>	A077179 004	May 18, 2007
AB		<u>EQ 10MG BASE;40MG</u>	A077179 006	Jul 05, 2011
AB	WATSON LABS	<u>EQ 2.5MG BASE;10MG</u>	A077890 001	Oct 14, 2010
AB		<u>EQ 5MG BASE;10MG</u>	A077890 002	Oct 14, 2010
AB		<u>EQ 5MG BASE;20MG</u>	A077890 003	Oct 14, 2010
AB		<u>EQ 10MG BASE;20MG</u>	A077890 004	Oct 14, 2010
AB	WATSON LABS INC	<u>EQ 5MG BASE;40MG</u>	A090364 001	Jul 05, 2011
AB		<u>EQ 10MG BASE;40MG</u>	A090364 002	Jul 05, 2011
	LOTREL			
AB	NOVARTIS	<u>EQ 2.5MG BASE;10MG</u>	N020364 002	Mar 03, 1995
AB		<u>EQ 5MG BASE;10MG</u>	N020364 003	Mar 03, 1995
AB		<u>EQ 5MG BASE;20MG</u>	N020364 004	Mar 03, 1995
AB		<u>EQ 5MG BASE;40MG</u>	N020364 007	Apr 11, 2006
AB		<u>EQ 10MG BASE;20MG</u>	N020364 005	Jun 20, 2002
AB	+	<u>EQ 10MG BASE;40MG</u>	N020364 006	Apr 11, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 29 (of 424)

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL TRIBENZOR			
DAIICHI SANKYO	EQ 5MG BASE;12.5MG;20MG	N200175 001	Jul 23, 2010
	EQ 5MG BASE;12.5MG;40MG	N200175 002	Jul 23, 2010
	EQ 5MG BASE;25MG;40MG	N200175 003	Jul 23, 2010
	EQ 10MG BASE;12.5MG;40MG	N200175 004	Jul 23, 2010
+	EQ 10MG BASE;25MG;40MG	N200175 005	Jul 23, 2010

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL EXFORGE HCT			
NOVARTIS	5MG;12.5MG;160MG	N022314 001	Apr 30, 2009
	5MG;25MG;160MG	N022314 002	Apr 30, 2009
	10MG;12.5MG;160MG	N022314 003	Apr 30, 2009
	10MG;25MG;160MG	N022314 004	Apr 30, 2009
+	10MG;25MG;320MG	N022314 005	Apr 30, 2009

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET; ORAL AZOR			
DAIICHI SANKYO	EQ 5MG BASE;20MG	N022100 001	Sep 26, 2007
	EQ 5MG BASE;40MG	N022100 002	Sep 26, 2007
	EQ 10MG BASE;20MG	N022100 003	Sep 26, 2007
+	EQ 10MG BASE;40MG	N022100 004	Sep 26, 2007

AMLODIPINE BESYLATE; TELMISARTAN

TABLET; ORAL TWYNSTA			
BOEHRINGER INGELHEIM	EQ 5MG BASE;40MG	N022401 001	Oct 16, 2009
	EQ 5MG BASE;80MG	N022401 003	Oct 16, 2009
	EQ 10MG BASE;40MG	N022401 002	Oct 16, 2009
+	EQ 10MG BASE;80MG	N022401 004	Oct 16, 2009

AMLODIPINE BESYLATE; VALSARTAN

TABLET; ORAL EXFORGE			
NOVARTIS	EQ 5MG BASE;160MG	N021990 002	Jun 20, 2007
	EQ 5MG BASE;320MG	N021990 004	Jun 20, 2007
+	EQ 10MG BASE;160MG	N021990 003	Jun 20, 2007
+	EQ 10MG BASE;320MG	N021990 005	Jun 20, 2007

AMMONIA, N-13

INJECTABLE; INTRAVENOUS AMMONIA N 13			
+ FEINSTEIN	30MCI-300MCI/8ML (3.75-37.5MCI/ML)	N022119 001	Aug 23, 2007

AMMONIUM CHLORIDE

INJECTABLE; INJECTION AMMONIUM CHLORIDE IN PLASTIC CONTAINER			
+ HOSPIRA	5MEQ/ML	A088366 001	Jun 13, 1984

AMMONIUM LACTATE

CREAM; TOPICAL AMMONIUM LACTATE			
AB PADDOCK LLC	EQ 12% BASE	A076829 001	Feb 07, 2006
AB PERRIGO NEW YORK	EQ 12% BASE	A075774 001	May 01, 2002
AB TARO	EQ 12% BASE	A075883 001	Apr 10, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 30 (of 424)

AMMONIUM LACTATE

CREAM; TOPICAL LAC-HYDRIN AB + RANBAXY	<u>EQ 12% BASE</u>	<u>N020508 001</u>	Aug 29, 1996
LOTION; TOPICAL AMMONIUM LACTATE			
AB PADDOCK LLC	<u>EQ 12% BASE</u>	<u>A075575 001</u>	Jun 11, 2002
AB PERRIGO NEW YORK	<u>EQ 12% BASE</u>	<u>A075570 001</u>	Jun 23, 2004
AB TARO	<u>EQ 12% BASE</u>	<u>A076216 001</u>	May 28, 2004
LAC-HYDRIN AB + RANBAXY	<u>EQ 12% BASE</u>	<u>N019155 001</u>	Apr 24, 1985

AMOXAPINE

TABLET; ORAL AMOXAPINE WATSON LABS	25MG 50MG 100MG + 150MG	A072688 001 A072689 001 A072690 001 A072691 001	Aug 28, 1992 Aug 28, 1992 Aug 28, 1992 Aug 28, 1992
--	----------------------------------	--	--

AMOXICILLIN

CAPSULE; ORAL AMOXICILLIN			
AB AM ANTIBIOTICS	<u>250MG</u>	<u>A062058 001</u>	
AB	<u>500MG</u>	<u>A062058 002</u>	
AB AUROBINDO	<u>250MG</u>	<u>A065271 001</u>	Nov 09, 2005
AB	<u>500MG</u>	<u>A065271 002</u>	Nov 09, 2005
AB DAVA PHARMS INC	<u>250MG</u>	<u>A062884 001</u>	Feb 25, 1988
AB	<u>500MG</u>	<u>A062881 001</u>	Feb 25, 1988
AB HIKMA PHARMS	<u>250MG</u>	<u>A065291 001</u>	Feb 05, 2007
AB	<u>500MG</u>	<u>A065291 002</u>	Feb 05, 2007
AB RANBAXY	<u>250MG</u>	<u>A065016 001</u>	Apr 08, 1999
AB	<u>500MG</u>	<u>A065016 002</u>	Apr 08, 1999
AB SANDOZ	<u>250MG</u>	<u>A064076 001</u>	Sep 30, 1994
AB	<u>500MG</u>	<u>A064076 002</u>	Sep 30, 1994
AB TEVA	<u>250MG</u>	<u>A061926 001</u>	
AB +	<u>500MG</u>	<u>A061926 003</u>	
AMOXIL			
AB DR REDDYS LABS INC	<u>250MG</u>	<u>A062216 001</u>	
AB	<u>500MG</u>	<u>A062216 004</u>	

FOR SUSPENSION; ORAL

AMOXICILLIN			
AB AUROBINDO	<u>200MG/5ML</u>	<u>A065334 001</u>	Dec 28, 2006
AB	<u>400MG/5ML</u>	<u>A065334 002</u>	Dec 28, 2006
AB DAVA PHARMS INC	<u>125MG/5ML</u>	<u>A062927 001</u>	Nov 25, 1988
AB	<u>250MG/5ML</u>	<u>A062927 002</u>	Nov 25, 1988
AB HIKMA	<u>125MG/5ML</u>	<u>A065322 002</u>	Jun 19, 2006
AB	<u>200MG/5ML</u>	<u>A065325 002</u>	Jun 19, 2006
AB	<u>250MG/5ML</u>	<u>A065322 001</u>	Jun 19, 2006
AB	<u>400MG/5ML</u>	<u>A065325 001</u>	Jun 19, 2006
AB RANBAXY	<u>200MG/5ML</u>	<u>A065113 001</u>	Nov 29, 2002
AB	<u>400MG/5ML</u>	<u>A065113 002</u>	Nov 29, 2002
AB SANDOZ	<u>125MG/5ML</u>	<u>A065387 001</u>	Mar 26, 2007
AB	<u>200MG/5ML</u>	<u>A065378 001</u>	Mar 26, 2007
AB	<u>250MG/5ML</u>	<u>A065387 002</u>	Mar 26, 2007
AB	<u>400MG/5ML</u>	<u>A065378 002</u>	Mar 26, 2007
AB TEVA	<u>125MG/5ML</u>	<u>A061931 001</u>	
AB	<u>200MG/5ML</u>	<u>A065119 001</u>	Dec 04, 2002
AB +	<u>250MG/5ML</u>	<u>A061931 002</u>	
AB +	<u>400MG/5ML</u>	<u>A065119 002</u>	Dec 04, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 31 (of 424)

AMOXICILLIN

FOR SUSPENSION; ORAL

AMOXICILLINAB WOCKHARDT 400MG/5ML A065319 002 Jun 18, 2007AMOXICILLIN PEDIATRICAB TEVA 50MG/ML A061931 003 Dec 01, 1982AMOXILAB DR REDDYS LABS INC 50MG/ML A062226 005AB 125MG/5ML A062226 001AB 200MG/5ML N050760 001 Apr 15, 1999AB 250MG/5ML A062226 002AB 400MG/5ML N050760 002 Apr 15, 1999LAROTIDAB DR REDDYS LABS INC 125MG/5ML A062226 003AB 250MG/5ML A062226 004

TABLET; ORAL

AMOXICILLINAB AUROBINDO 500MG A065256 001 Nov 09, 2005AB 875MG A065256 002 Nov 09, 2005AB DAVA PHARMS INC 875MG A065344 001 Jan 15, 2009AB HIKMA 875MG A065255 001 Mar 29, 2006AB RANBAXY 500MG A065059 001 Nov 24, 2000AB 875MG A065059 002 Nov 24, 2000AB SANDOZ 500MG A065228 001 Jul 13, 2005AB 875MG A065228 002 Jul 13, 2005AB TEVA 500MG A065056 001 Sep 18, 2000AB + 875MG A065056 002 Sep 18, 2000AMOXILAB DR REDDYS LABS INC 500MG N050754 002 Jul 10, 1998AB 875MG N050754 001 Jul 10, 1998

TABLET, CHEWABLE; ORAL

AMOXICILLINAB DAVA PHARMS INC 125MG A064139 001 Jan 29, 1996AB 250MG A064139 002 Jan 29, 1996AB RANBAXY 125MG A065021 001 Dec 23, 1999AB 250MG A065021 002 Dec 23, 1999AB TEVA 125MG A064013 002 Sep 11, 1995AB + 250MG A064013 001 Dec 22, 1992AMOXILAB DR REDDYS LABS INC 125MG N050542 002AB 200MG N050761 001 Apr 15, 1999AB 250MG N050542 001AB 400MG N050761 002 Apr 15, 1999

AMOXICILLIN

RANBAXY 200MG A065060 001 Nov 29, 2000RANBAXY 400MG A065060 002 Nov 29, 2000

TABLET, EXTENDED RELEASE; ORAL

MOXATAG

+ SHIONOGI INC 775MG N050813 001 Jan 23, 2008

TABLET, FOR SUSPENSION; ORAL

AMOXICILLINAB AUROBINDO PHARMA 200MG A065324 001 Jan 17, 2007AB 400MG A065324 002 Jan 17, 2007

DISPERMOX

+ RANBAXY 600MG A065159 001 Dec 04, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 32 (of 424)

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS; ORAL
 PREVPAC
 + TAKEDA PHARMS NA 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30M N050757 001 Dec 02, 1997
 G

AMOXICILLIN; CLARITHROMYCIN; OMEPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED RELEASE, TABLET; ORAL
 OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN
 + GASTROENTERO 500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,20M N050824 001 Feb 08, 2011
 G

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<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>	LEK PHARMS	<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>	LEK PHARMS DD	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065358 001</u>	Aug 13, 2007	
<u>AB</u>	RANBAXY	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065132 001</u>	Mar 19, 2003	
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065132 002</u>	Mar 19, 2003	
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065207 002</u>	Jan 30, 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>	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>	WOCHARDT	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A065431 001</u>	Nov 25, 2008	
<u>AUGMENTIN '125'</u>					
<u>AB</u>	DR REDDYS LABS INC	<u>125MG/5ML;EQ 31.25MG BASE/5ML</u>	<u>N050575 001</u>	Aug 06, 1984	
<u>AUGMENTIN '200'</u>					
<u>AB</u>	DR REDDYS LABS INC	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N050725 001</u>	May 31, 1996	
<u>AUGMENTIN '250'</u>					
<u>AB</u>	+	<u>DR REDDYS LABS INC</u>	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>N050575 002</u>	Aug 06, 1984
<u>AUGMENTIN '400'</u>					
<u>AB</u>	DR REDDYS LABS INC	<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N050725 002</u>	May 31, 1996	
<u>AUGMENTIN ES-600</u>					
<u>AB</u>	DR REDDYS LABS INC	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>N050755 001</u>	Jun 22, 2001	

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

TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	APOTEX	<u>250MG;EQ 125MG BASE</u>	<u>A065333 001</u>	Feb 24, 2009
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A065333 002</u>	Feb 24, 2009
<u>AB</u>	APOTEX INC	<u>875MG;EQ 125MG BASE</u>	<u>A065317 003</u>	Oct 20, 2008
<u>AB</u>	LEK PHARMS	<u>500MG;EQ 125MG BASE</u>	<u>A065117 001</u>	Nov 27, 2002
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>A065093 001</u>	Nov 21, 2002
<u>AB</u>	RANBAXY	<u>500MG;EQ 125MG BASE</u>	<u>A065109 001</u>	Nov 04, 2002
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>A065102 001</u>	Sep 17, 2002
<u>AB</u>	SANDOZ	<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>	TEVA	<u>500MG;EQ 125MG BASE</u>	<u>A065101 001</u>	Oct 30, 2002
<u>AB</u>	+	<u>875MG;EQ 125MG BASE</u>	<u>A065096 001</u>	Oct 29, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 33 (of 424)

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET; ORAL

AUGMENTIN '250'

<u>AB</u>	DR REDDYS LABS INC	<u>250MG;EQ 125MG BASE</u>	<u>N050564 001</u>	Aug 06, 1984
<u>AB</u>	DR REDDYS LABS INC	<u>500MG;EQ 125MG BASE</u>	<u>N050564 002</u>	Aug 06, 1984
<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

<u>AB</u>	RANBAXY	<u>200MG;EQ 28.5MG BASE</u>	<u>A065161 001</u>	Dec 03, 2003
<u>AB</u>		<u>400MG;EQ 57MG BASE</u>	<u>A065161 002</u>	Dec 03, 2003
<u>AB</u>	SANDOZ	<u>200MG;EQ 28.5MG BASE</u>	<u>A065065 001</u>	Apr 18, 2002
<u>AB</u>		<u>400MG;EQ 57MG BASE</u>	<u>A065065 002</u>	Apr 18, 2002
<u>AB</u>	TEVA	<u>200MG;EQ 28.5MG BASE</u>	<u>A065205 001</u>	Feb 09, 2005
<u>AB</u>	+	<u>400MG;EQ 57MG BASE</u>	<u>A065205 002</u>	Feb 09, 2005
<u>AB</u>		<u>AUGMENTIN '125'</u>		
<u>AB</u>	DR REDDYS LABS INC	<u>125MG;EQ 31.25MG BASE</u>	<u>N050597 001</u>	Jul 22, 1985
<u>AB</u>		<u>AUGMENTIN '200'</u>		
<u>AB</u>	DR REDDYS LABS INC	<u>200MG;EQ 28.5MG BASE</u>	<u>N050726 001</u>	May 31, 1996
<u>AB</u>		<u>AUGMENTIN '250'</u>		
<u>AB</u>	DR REDDYS LABS INC	<u>250MG;EQ 62.5MG BASE</u>	<u>N050597 002</u>	Jul 22, 1985
<u>AB</u>		<u>AUGMENTIN '400'</u>		
<u>AB</u>	DR REDDYS LABS INC	<u>400MG;EQ 57MG BASE</u>	<u>N050726 002</u>	May 31, 1996

TABLET, EXTENDED RELEASE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	SANDOZ	<u>1GM;EQ 62.5MG BASE</u>	<u>A090227 001</u>	Apr 21, 2010	
<u>AB</u>	+	<u>DR REDDYS LABS INC</u>	<u>1GM;EQ 62.5MG BASE</u>	<u>N050785 001</u>	Sep 25, 2002

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

ADDERALL XR 10

	SHIRE	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N021303 001</u>	Oct 11, 2001
	ADDERALL XR 15	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N021303 006</u>	May 22, 2002
	ADDERALL XR 20	<u>5MG;5MG;5MG;5MG</u>	<u>N021303 002</u>	Oct 11, 2001
	ADDERALL XR 25	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>N021303 004</u>	May 22, 2002
	ADDERALL XR 30	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N021303 003</u>	Oct 11, 2001
+	SHIRE	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N021303 005</u>	May 22, 2002
	ADDERALL XR 5			
	SHIRE			

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<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>	COREPHARMA	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040444 001</u>	Jun 19, 2002
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040444 002</u>	Jun 19, 2002
<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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 34 (of 424)

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE;
DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u>	SANDOZ	<u>5MG; 5MG; 5MG; 5MG</u>	<u>A040439</u>	<u>002</u>	Jun 14, 2002
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A040439</u>	<u>003</u>	Jun 14, 2002
<u>AB</u>	TEVA PHARMS	<u>1.25MG; 1.25MG; 1.25MG; 1.25MG</u>	<u>A040472</u>	<u>001</u>	Sep 30, 2003
<u>AB</u>		<u>2.5MG; 2.5MG; 2.5MG; 2.5MG</u>	<u>A040472</u>	<u>002</u>	Sep 30, 2003
<u>AB</u>		<u>5MG; 5MG; 5MG; 5MG</u>	<u>A040472</u>	<u>003</u>	Sep 30, 2003
<u>AB</u>		<u>7.5MG; 7.5MG; 7.5MG; 7.5MG</u>	<u>A040472</u>	<u>004</u>	Sep 30, 2003

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

<u>AP</u>	+ X GEN PHARMS	<u>50MG/VIAL</u>	<u>A063206</u>	<u>001</u>	Apr 29, 1992
INJECTABLE, LIPID COMPLEX; INJECTION					
ABELCET					
+ SIGMA TAU		5MG/ML	N050724	001	Nov 20, 1995
AMPHOTEC					
+ ALDOPHARMA USA		50MG/VIAL	N050729	001	Nov 22, 1996
+ ASTELLAS		100MG/VIAL	N050729	002	Nov 22, 1996
INJECTABLE, LIPOSOMAL; INJECTION					
AMBISOME					
+ ASTELLAS		50MG/VIAL	N050740	001	Aug 11, 1997

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

<u>AP</u>	ACIC FINE CHEMS	<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>	HANFORD GC	<u>EQ 10GM BASE/VIAL</u>	<u>A065493</u>	<u>001</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A063145</u>	<u>001</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A063146</u>	<u>001</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>A063140</u>	<u>001</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>	IBI	<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>	INSTITUTO BIOCHEMICO	<u>EQ 125MG BASE/VIAL</u>	<u>A062797</u>	<u>001</u>	Jul 12, 1993
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062797</u>	<u>002</u>	Jul 12, 1993
<u>AP</u>	+ SANDOZ	<u>EQ 125MG BASE/VIAL</u>	<u>A061395</u>	<u>001</u>	
<u>AP</u>	+	<u>EQ 250MG BASE/VIAL</u>	<u>A061395</u>	<u>002</u>	
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>A061395</u>	<u>003</u>	
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>A061395</u>	<u>004</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062738</u>	<u>001</u>	Feb 19, 1987
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>A061395</u>	<u>005</u>	
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>A062738</u>	<u>002</u>	Feb 19, 1987
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>A061395</u>	<u>006</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 35 (of 424)

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

AP	ACS DOBFAR	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065406	001	Dec 22, 2009
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065406	002	Dec 22, 2009
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A065403	001	Dec 23, 2009
AP	AUROBINDO PHARMA	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090340	001	Sep 20, 2010
AP		EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090349	001	Sep 20, 2010
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090340	002	Sep 20, 2010
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090349	002	Sep 20, 2010
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A090339	001	Sep 20, 2010
AP	BAXTER HLTHCARE	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065074	001	Mar 19, 2002
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065074	002	Mar 19, 2002
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A065076	001	Mar 19, 2002
AP	BIONICHE PHARMA	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065316	001	Jun 29, 2007
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065316	002	Jun 29, 2007
AP	HANFORD GC	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065176	001	Nov 30, 2005
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065176	002	Nov 30, 2005
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A065188	001	Nov 25, 2005
AP	HOSPIRA INC	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090375	001	Dec 21, 2011
AP		EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A090653	001	Dec 21, 2011
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090375	002	Dec 21, 2011
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A090653	002	Dec 21, 2011
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A090646	001	Dec 21, 2011
AP	INSTITUTO BIOCHIMICO	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065222	001	Nov 29, 2005
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065222	002	Nov 29, 2005
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A065314	001	Nov 27, 2006
AP	SANDOZ	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065241	001	Jul 25, 2006
AP		EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A065310	001	Jul 25, 2006
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065241	002	Jul 25, 2006
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	A065310	002	Jul 25, 2006
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	A065240	001	Jul 25, 2006
<u>UNASYN</u>					
AP	PFIZER	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	A062901	001	Nov 23, 1988
AP	+	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	N050608	002	Dec 31, 1986
AP	+	EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	N050608	001	Dec 31, 1986
AP	+	EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	N050608	005	Dec 10, 1993

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

AB	DAVA PHARMS INC	EQ 250MG BASE	A062883	001	Feb 25, 1988
AB	+	EQ 500MG BASE	A062882	001	Feb 25, 1988
AB	SANDOZ	EQ 250MG BASE	A064082	001	Aug 29, 1995
AB		EQ 500MG BASE	A064082	002	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

AB	SHIRE	EQ 0.5MG BASE	N020333	001	Mar 14, 1997
<u>ANAGRELIDE HYDROCHLORIDE</u>					
AB	ALPHAPHARM	EQ 0.5MG BASE	A077613	001	Jun 27, 2006
AB		EQ 1MG BASE	A077613	002	Jun 27, 2006
AB	BARR	EQ 0.5MG BASE	A076530	001	Apr 18, 2005
AB		EQ 1MG BASE	A076530	002	Apr 18, 2005
AB	IMPAX LABS	EQ 0.5MG BASE	A076910	001	Apr 18, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 36 (of 424)

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

ANAGRELIDE HYDROCHLORIDE

<u>AB</u>	IMPAX LABS	<u>EQ 1MG BASE</u>	<u>A076910</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 0.5MG BASE</u>	<u>A076468</u>	<u>001</u>	Apr 18, 2005
<u>AB</u> +		<u>EQ 1MG BASE</u>	<u>A076468</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	MYLAN	<u>EQ 0.5MG BASE</u>	<u>A076811</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076811</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	SANDOZ	<u>EQ 0.5MG BASE</u>	<u>A076683</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076683</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	WATSON LABS	<u>EQ 0.5MG BASE</u>	<u>A076417</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076417</u>	<u>002</u>	Apr 18, 2005

ANASTROZOLE

TABLET; ORAL

ANASTROZOLE

<u>AB</u>	ACCORD HLTHCARE INC	<u>1MG</u>	<u>A090568</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A090732</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	FRESENIUS KABI ONCOL	<u>1MG</u>	<u>A090088</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	KUDCO IRELAND	<u>1MG</u>	<u>A091331</u>	<u>001</u>	Jan 05, 2011
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A091051</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	NATCO PHARMA LTD	<u>1MG</u>	<u>A079220</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	ROXANE	<u>1MG</u>	<u>A078485</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	SANDOZ	<u>1MG</u>	<u>A079007</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	SANTOS BIOTECH	<u>1MG</u>	<u>A078944</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	SUN PHARM IND LTD	<u>1MG</u>	<u>A091177</u>	<u>001</u>	Jul 15, 2011
<u>AB</u>	TEVA PHARMS	<u>1MG</u>	<u>A078058</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	THREE RIVERS PHARMS	<u>1MG</u>	<u>A091164</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>A078984</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>1MG</u>	<u>A078921</u>	<u>001</u>	Jun 28, 2010
	<u>ARIMIDEX</u>				
<u>AB</u> +	ASTRAZENECA	<u>1MG</u>	<u>N020541</u>	<u>001</u>	Dec 27, 1995

ANIDULAFUNGIN

INJECTABLE; IV (INFUSION)

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

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

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

APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLONIDINE HYDROCHLORIDE

<u>AT</u>	AKORN INC	<u>EQ 0.5% BASE</u>	<u>A077764</u>	<u>001</u>	Mar 12, 2009
	<u>IOPIDINE</u>				
<u>AT</u> +	ALCON	<u>EQ 0.5% BASE</u>	<u>N020258</u>	<u>001</u>	Jul 30, 1993
	IOPIDINE				
+ ALCON		<u>EQ 1% BASE</u>	<u>N019779</u>	<u>001</u>	Dec 31, 1987

APREPITANT

CAPSULE; ORAL

EMEND

MERCK	40MG	N021549	003	Jun 30, 2006
	80MG	N021549	001	Mar 26, 2003
+	125MG	N021549	002	Mar 26, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 37 (of 424)

APROTININ

INJECTABLE; INJECTION
 TRASYLOL
 + BAYER HLTHCARE 10,000KIU/ML N020304 001 Dec 29, 1993

ARFORMOTEROL TARTRATE

SOLUTION; INHALATION
 BROVANA
 + SUNOVION EQ 0.015MG BASE/2ML N021912 001 Oct 06, 2006

ARGATROBAN

INJECTABLE; INJECTION
 ARGATROBAN
 + PFIZER 100MG/ML N020883 001 Jun 30, 2000

INJECTABLE; IV (INFUSION)

+ EAGLE PHARMS 50MG/50ML (1MG/ML) N022434 001 Jun 29, 2011
 + SANDOZ 125MG/125ML (1MG/ML) N022485 001 May 09, 2011

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION
 R-GENE 10
 + PHARMACIA AND UPJOHN 10GM/100ML N016931 001

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; ORAL

ABILIFY
 OTSUKA 2MG N021436 006 Nov 15, 2002
 + 5MG N021436 005 Nov 15, 2002
 + 10MG N021436 001 Nov 15, 2002
 15MG N021436 002 Nov 15, 2002
 20MG N021436 003 Nov 15, 2002
 30MG N021436 004 Nov 15, 2002

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY
 + OTSUKA 10MG N021729 002 Jun 07, 2006
 15MG N021729 003 Jun 07, 2006

ARMODAFINIL

TABLET; ORAL
 NUvigil
 CEPHALON 50MG N021875 001 Jun 15, 2007
 150MG N021875 003 Jun 15, 2007
 + 250MG N021875 004 Jun 15, 2007

ARSENIC TRIOXIDE

INJECTABLE; INJECTION
 TRISENOX
 + CEPHALON 1MG/ML N021248 001 Sep 25, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 38 (of 424)

ARTEMETHER; LUMEFANTRINE

TABLET; ORAL
 COARTEM
 + NOVARTIS 20MG;120MG N022268 001 Apr 07, 2009

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

<u>AP</u>	HOSPIRA	<u>4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)</u>	<u>A079138 001</u>	Jun 18, 2010
<u>AP</u>	+ DEPROCO	<u>4%;EQ 0.017MG BASE/1.7ML (4%;EQ 0.01MG BASE/ML)</u>	<u>N020971 001</u>	Apr 03, 2000
ARTICAINE HYDROCHLORIDE WITH EPINEPHRINE				
	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
SEPTOCAIN				
	+ DEPROCO	4%;EQ 0.0085MG BASE/1.7ML (4%;EQ 0.005MG BASE/ML)	N022010 001	Mar 30, 2006

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

INJECTABLE; IV (INFUSION)

INFUVITE PEDIATRIC

+ SANDOZ	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
+ SANDOZ	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

INJECTABLE; INJECTION

M.V.I.-12 (WITHOUT VITAMIN K)

+ HOSPIRA	20MG/ML;0.006MG/ML;0.05MCG/ML;1.5MG/ML; 0.0005MG/ML;0.06MG/ML;4MG/ML;0.6MG/ML;0 .36MG/ML;0.6MG/ML;0.1MG/ML;1MG/ML	N008809 006	Sep 09, 2004
-----------	---	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 39 (of 424)

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;15MG G/VIAL;0.005MG/VIAL;0.6MG/VIAL;40MG/VIAL;6MG/VIAL;3.6MG/VIAL;6MG/VIAL;1MG/VIAL;10MG/VIAL;0.15MG/VIAL

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

+ HOSPIRA 200MG/5ML;0.06MG/5ML;0.005MG/5ML;15MG/5ML;0.005MG/5ML;0.6MG/5ML;40MG/5ML;6MG/5ML;3.6MG/5ML;6MG/5ML;1MG/5ML;10MG/5ML;0.15MG/5ML

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 MALEATETABLET; SUBLINGUAL
SAPHRIS

ORGANON USA INC EQ 5MG BASE N022117 001 Aug 13, 2009
+ EQ 10MG BASE N022117 002 Aug 13, 2009

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINAL

AA + WATSON LABS INC 325MG;50MG;40MG N017534 005 Apr 16, 1986
LANORINAL

AA LANNETT 325MG;50MG;40MG A086996 002 Oct 11, 1985

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

AA + WEST WARD 325MG;50MG;40MG A086162 002 Feb 16, 1984

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

<u>AB</u>	NEXGEN PHARMA INC	<u>325MG;50MG;40MG;30MG</u>	A075231 001	Nov 30, 2001
<u>AB</u>	STEVENS J	<u>325MG;50MG;40MG;30MG</u>	A074951 001	Aug 31, 1998
<u>AB</u>	WATSON LABS	<u>325MG;50MG;40MG;30MG</u>	A074359 001	Aug 31, 1995
<u>FIORINAL W/CODEINE</u>				
<u>AB</u>	+ WATSON LABS INC	<u>325MG;50MG;40MG;30MG</u>	N019429 003	Oct 26, 1990

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

SYNALGOS-DC

+ CARACO 356.4MG;30MG;16MG N011483 004 Sep 06, 1983

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

INVAGESIC

AB SANDOZ 385MG;30MG;25MG A074817 001 Nov 27, 1996

INVAGESIC FORTE

AB SANDOZ 770MG;60MG;50MG A074817 002 Nov 27, 1996

NORGESIC

AB MEDICIS 385MG;30MG;25MG N013416 003 Oct 27, 1982

NORGESIC FORTE

AB + MEDICIS 770MG;60MG;50MG N013416 004 Oct 27, 1982

PRESCRIPTION DRUG PRODUCT LIST

3 - 40 (of 424)

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

<u>AB</u>	SANDOZ	<u>385MG;30MG;25MG</u>	<u>A074654</u> <u>001</u>	Dec 31, 1996
<u>AB</u>		<u>770MG;60MG;50MG</u>	<u>A074654</u> <u>002</u>	Dec 31, 1996
<u>AB</u>	STEVENS J	<u>385MG;30MG;25MG</u>	<u>A074988</u> <u>001</u>	Apr 30, 1999
<u>AB</u>		<u>770MG;60MG;50MG</u>	<u>A074988</u> <u>002</u>	Apr 30, 1999

ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

<u>AB</u>	HERITAGE PHARMS INC	<u>325MG;200MG</u>	<u>A089594</u> <u>001</u>	Mar 31, 1989
<u>AB</u>	MIRROR PHARMS	<u>325MG;200MG</u>	<u>A040832</u> <u>001</u>	Jan 07, 2010
<u>AB</u>	PROSAM LABS	<u>325MG;200MG</u>	<u>A040252</u> <u>001</u>	Dec 10, 1997
<u>AB</u>	SANDOZ	<u>325MG;200MG</u>	<u>A040116</u> <u>001</u>	Apr 25, 1996
	<u>SOMA COMPOUND</u>			
<u>AB</u>	+ MEDA PHARMS	<u>325MG;200MG</u>	<u>N012365</u> <u>005</u>	Jul 11, 1983

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

<u>AB</u>	MIRROR PHARMS	<u>325MG;200MG;16MG</u>	<u>A040860</u> <u>001</u>	Jan 07, 2010
<u>AB</u>	PROSAM LABS	<u>325MG;200MG;16MG</u>	<u>A040283</u> <u>001</u>	Dec 29, 1998
<u>AB</u>	SANDOZ	<u>325MG;200MG;16MG</u>	<u>A040118</u> <u>001</u>	Apr 16, 1996
	<u>SOMA COMPOUND W/ CODEINE</u>			
<u>AB</u>	+ MEDA PHARMS	<u>325MG;200MG;16MG</u>	<u>N012366</u> <u>002</u>	Jul 11, 1983

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOX

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>25MG;200MG</u>	<u>N020884</u> <u>001</u>	Nov 22, 1999
<u>AB</u>	<u>ASPIRIN AND DIPYRIDAMOLE</u>		<u>A078804</u> <u>001</u>	Aug 14, 2009

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL AND ASPIRIN

<u>AB</u>	STEVENS J	<u>325MG;400MG</u>	<u>A081145</u> <u>001</u>	Jan 31, 1995
-----------	-----------	--------------------	---------------------------	--------------

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE AND ASPIRIN

<u>AA</u>	COASTAL PHARMS	<u>325MG;4.8355MG</u>	<u>A091670</u> <u>001</u>	Mar 16, 2011
<u>AA</u>	WATSON LABS	<u>325MG;4.8355MG</u>	<u>A090084</u> <u>001</u>	Mar 22, 2011
	<u>PERCODAN</u>			
<u>AA</u>	+ ENDO PHARMS	<u>325MG;4.8355MG</u>	<u>N007337</u> <u>007</u>	Aug 05, 2005

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

OXYCODONE AND ASPIRIN

<u>WATSON LABS</u>	<u>325MG;4.5MG;0.38MG</u>	<u>A040255</u> <u>001</u>	Feb 27, 1998
--------------------	---------------------------	---------------------------	--------------

ATAZANAVIR SULFATE

CAPSULE; ORAL

REYATAZ

<u>BRISTOL MYERS SQUIBB</u>	<u>EQ 100MG BASE</u>	<u>N021567</u> <u>001</u>	Jun 20, 2003
	<u>EQ 150MG BASE</u>	<u>N021567</u> <u>002</u>	Jun 20, 2003
	<u>EQ 200MG BASE</u>	<u>N021567</u> <u>003</u>	Jun 20, 2003
<u>+</u>	<u>EQ 300MG BASE</u>	<u>N021567</u> <u>004</u>	Oct 16, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 41 (of 424)

ATENOLOL

TABLET; ORAL

ATENOLOL

<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A078512</u>	<u>001</u>	Oct 31, 2007
<u>AB</u>		<u>50MG</u>	<u>A078512</u>	<u>002</u>	Oct 31, 2007
<u>AB</u>		<u>100MG</u>	<u>A078512</u>	<u>003</u>	Oct 31, 2007
<u>AB</u>	CARACO	<u>25MG</u>	<u>A078210</u>	<u>001</u>	Jul 10, 2007
<u>AB</u>		<u>50MG</u>	<u>A078210</u>	<u>002</u>	Jul 10, 2007
<u>AB</u>		<u>100MG</u>	<u>A078210</u>	<u>003</u>	Jul 10, 2007
<u>AB</u>	DAVA PHARMS INC	<u>25MG</u>	<u>A074099</u>	<u>001</u>	Apr 28, 1992
<u>AB</u>		<u>50MG</u>	<u>A073542</u>	<u>001</u>	Dec 19, 1991
<u>AB</u>		<u>100MG</u>	<u>A073543</u>	<u>001</u>	Dec 19, 1991
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A077877</u>	<u>001</u>	Dec 27, 2006
<u>AB</u>		<u>50MG</u>	<u>A077877</u>	<u>002</u>	Dec 27, 2006
<u>AB</u>		<u>100MG</u>	<u>A077877</u>	<u>003</u>	Dec 27, 2006
<u>AB</u>	IPR	<u>25MG</u>	<u>A073646</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>		<u>50MG</u>	<u>A072303</u>	<u>001</u>	Jul 15, 1988
<u>AB</u>		<u>100MG</u>	<u>A072304</u>	<u>001</u>	Jul 15, 1988
<u>AB</u>	MUTUAL PHARM	<u>25MG</u>	<u>A074499</u>	<u>001</u>	Jul 30, 1997
<u>AB</u>		<u>50MG</u>	<u>A073475</u>	<u>001</u>	Mar 30, 1993
<u>AB</u>		<u>100MG</u>	<u>A073476</u>	<u>001</u>	Mar 30, 1993
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A073457</u>	<u>002</u>	Apr 26, 1999
<u>AB</u>		<u>50MG</u>	<u>A074126</u>	<u>003</u>	Aug 26, 1998
<u>AB</u>		<u>100MG</u>	<u>A073456</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>		<u>50MG</u>	<u>A074126</u>	<u>001</u>	Mar 23, 1994
<u>AB</u>		<u>100MG</u>	<u>A073457</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>		<u>100MG</u>	<u>A074126</u>	<u>002</u>	Mar 23, 1994
<u>AB</u>	NORTHSTAR HLTHCARE	<u>25MG</u>	<u>A078254</u>	<u>001</u>	Sep 25, 2009
<u>AB</u>		<u>50MG</u>	<u>A078254</u>	<u>002</u>	Sep 25, 2009
<u>AB</u>		<u>100MG</u>	<u>A078254</u>	<u>003</u>	Sep 25, 2009
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A074052</u>	<u>001</u>	May 01, 1992
<u>AB</u>		<u>50MG</u>	<u>A073025</u>	<u>001</u>	Sep 17, 1991
<u>AB</u>		<u>100MG</u>	<u>A073026</u>	<u>001</u>	Sep 17, 1991
<u>AB</u>	TEVA	<u>25MG</u>	<u>A074056</u>	<u>003</u>	Jul 19, 2004
<u>AB</u>		<u>50MG</u>	<u>A074056</u>	<u>001</u>	Jan 18, 1995
<u>AB</u>		<u>100MG</u>	<u>A074056</u>	<u>002</u>	Jan 18, 1995
<u>AB</u>	UNIQUE PHARM LABS	<u>25MG</u>	<u>A077443</u>	<u>001</u>	Sep 13, 2006
<u>AB</u>		<u>50MG</u>	<u>A077443</u>	<u>002</u>	Sep 13, 2006
<u>AB</u>		<u>100MG</u>	<u>A077443</u>	<u>003</u>	Sep 13, 2006
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>A073352</u>	<u>001</u>	Dec 27, 1991
<u>AB</u>		<u>100MG</u>	<u>A073353</u>	<u>001</u>	Dec 27, 1991
<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
	<u>TENORMIN</u>				
<u>AB</u>	ASTRAZENECA	<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>	IPR	<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>	MUTUAL PHARM	<u>50MG;25MG</u>	<u>A073581</u>	<u>001</u>	Apr 29, 1993
<u>AB</u>		<u>100MG;25MG</u>	<u>A073582</u>	<u>001</u>	Apr 29, 1993
<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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 42 (of 424)

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

TENORETIC 100

<u>AB</u> + ASTRazeneca	<u>100MG;25MG</u>	<u>N018760 001</u>	Jun 08, 1984
<u>AB</u> ASTRazeneca	<u>50MG;25MG</u>	<u>N018760 002</u>	Jun 08, 1984

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

ATOMOXETINE HYDROCHLORIDE

<u>AB</u> AUROBINDO PHARM	<u>10MG</u>	<u>A079016 001</u>	Sep 13, 2010	
<u>AB</u>	<u>18MG</u>	<u>A079016 002</u>	Sep 13, 2010	
<u>AB</u>	<u>25MG</u>	<u>A079016 003</u>	Sep 13, 2010	
<u>AB</u>	<u>40MG</u>	<u>A079016 004</u>	Sep 13, 2010	
<u>AB</u>	<u>60MG</u>	<u>A079016 005</u>	Sep 13, 2010	
<u>AB</u>	<u>80MG</u>	<u>A079016 006</u>	Sep 13, 2010	
<u>AB</u>	<u>100MG</u>	<u>A079016 007</u>	Sep 13, 2010	
<u>AB</u> MYLAN	<u>10MG</u>	<u>A079021 001</u>	Aug 30, 2010	
<u>AB</u>	<u>18MG</u>	<u>A079021 002</u>	Aug 30, 2010	
<u>AB</u>	<u>25MG</u>	<u>A079021 003</u>	Aug 30, 2010	
<u>AB</u>	<u>40MG</u>	<u>A079021 004</u>	Aug 30, 2010	
<u>AB</u>	<u>60MG</u>	<u>A079021 005</u>	Aug 30, 2010	
<u>AB</u>	<u>80MG</u>	<u>A079021 006</u>	Aug 30, 2010	
<u>AB</u>	<u>100MG</u>	<u>A079021 007</u>	Aug 30, 2010	
<u>AB</u> SANDOZ	<u>10MG</u>	<u>A079018 001</u>	Sep 13, 2010	
<u>AB</u>	<u>18MG</u>	<u>A079018 002</u>	Sep 13, 2010	
<u>AB</u>	<u>25MG</u>	<u>A079018 003</u>	Sep 13, 2010	
<u>AB</u>	<u>40MG</u>	<u>A079018 004</u>	Sep 13, 2010	
<u>AB</u>	<u>60MG</u>	<u>A079018 005</u>	Sep 13, 2010	
<u>AB</u>	<u>80MG</u>	<u>A079018 006</u>	Sep 13, 2010	
<u>AB</u>	<u>100MG</u>	<u>A079018 007</u>	Sep 13, 2010	
<u>AB</u> SUN PHARMA GLOBAL	<u>10MG</u>	<u>A079020 001</u>	Aug 30, 2010	
<u>AB</u>	<u>18MG</u>	<u>A079020 002</u>	Aug 30, 2010	
<u>AB</u>	<u>25MG</u>	<u>A079020 003</u>	Aug 30, 2010	
<u>AB</u>	<u>40MG</u>	<u>A079020 004</u>	Aug 30, 2010	
<u>AB</u>	<u>60MG</u>	<u>A079020 005</u>	Aug 30, 2010	
<u>AB</u>	<u>80MG</u>	<u>A079020 006</u>	Aug 30, 2010	
<u>AB</u>	<u>100MG</u>	<u>A079020 007</u>	Aug 30, 2010	
<u>AB</u> TEVA PHARMS	<u>10MG</u>	<u>A079022 001</u>	Oct 01, 2010	
<u>AB</u>	<u>18MG</u>	<u>A079022 002</u>	Oct 01, 2010	
<u>AB</u>	<u>25MG</u>	<u>A079022 003</u>	Oct 01, 2010	
<u>AB</u>	<u>40MG</u>	<u>A079022 004</u>	Oct 01, 2010	
<u>AB</u>	<u>60MG</u>	<u>A079022 005</u>	Oct 01, 2010	
<u>AB</u>	<u>80MG</u>	<u>A079022 006</u>	Oct 01, 2010	
<u>AB</u>	<u>100MG</u>	<u>A079022 007</u>	Oct 01, 2010	
<u>AB</u> ZYDUS PHARMS USA INC	<u>18MG</u>	<u>A079017 001</u>	Sep 17, 2010	
<u>AB</u>	<u>25MG</u>	<u>A079017 002</u>	Sep 17, 2010	
<u>AB</u>	<u>40MG</u>	<u>A079017 003</u>	Sep 17, 2010	
<u>AB</u>	<u>60MG</u>	<u>A079017 004</u>	Sep 17, 2010	
<u>AB</u>	<u>80MG</u>	<u>A079017 005</u>	Sep 17, 2010	
<u>AB</u>	<u>100MG</u>	<u>A079017 006</u>	Sep 17, 2010	
<u>AB</u> STRATTERA	<u>LILLY</u>	<u>10MG</u>	<u>N021411 002</u>	Nov 26, 2002
<u>AB</u>	<u>18MG</u>		<u>N021411 003</u>	Nov 26, 2002
<u>AB</u>	<u>25MG</u>		<u>N021411 004</u>	Nov 26, 2002
<u>AB</u>	<u>40MG</u>		<u>N021411 005</u>	Nov 26, 2002
<u>AB</u> +	<u>60MG</u>		<u>N021411 006</u>	Nov 26, 2002
<u>AB</u>	<u>80MG</u>		<u>N021411 007</u>	Feb 14, 2005
<u>AB</u>	<u>100MG</u>		<u>N021411 008</u>	Feb 14, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 43 (of 424)

ATORVASTATIN CALCIUM

TABLET; ORAL

ATORVASTATIN CALCIUM

<u>AB</u>	RANBAXY LABS LTD	<u>EQ 10MG BASE</u>	<u>A076477 001</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076477 002</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076477 003</u>	Nov 30, 2011
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A076477 004</u>	Nov 30, 2011
	<u>LIPITOR</u>			
<u>AB</u>	PFIZER	<u>EQ 10MG BASE</u>	<u>N020702 001</u>	Dec 17, 1996
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N020702 002</u>	Dec 17, 1996
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N020702 003</u>	Dec 17, 1996
<u>AB</u> +		<u>EQ 80MG BASE</u>	<u>N020702 004</u>	Apr 07, 2000

ATOVAQUONE

SUSPENSION; ORAL

MEPRON

+ GLAXOSMITHKLINE LLC 750MG/5ML

N020500 001 Feb 08, 1995

ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

ATOVAQUONE AND PROGUANIL HYDROCHLORIDE

<u>AB</u>	GLENMARK GENERICS	<u>250MG;100MG</u>	<u>A091211 001</u>	Jan 12, 2011
<u>AB</u>	<u>MALARONE</u>		<u>N021078 001</u>	Jul 14, 2000
<u>AB</u> +	GLAXOSMITHKLINE	<u>250MG;100MG</u>		
	MALARONE PEDIATRIC			
	GLAXOSMITHKLINE	62.5MG;25MG	N021078 002	Jul 14, 2000

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

+ BEDFORD 10MG/ML
ATRACURIUM BESYLATE PRESERVATIVE FREE
+ BEDFORD 10MG/MLA074901 001 Jul 18, 1997
A074900 001 Jul 18, 1997ATROPINE

INJECTABLE; INJECTION

ATROPEN

+ MERIDIAN MEDCL TECHN EQ 0.25MG SULFATE/0.3ML
+ EQ 0.5MG SULFATE/0.7ML
+ EQ 1MG SULFATE/0.7ML
+ EQ 2MG SULFATE/0.7MLN017106 004 Sep 17, 2004
N017106 003 Jun 19, 2003
N017106 002 Jun 19, 2003
N017106 001ATROPINE SULFATE

INJECTABLE; IM-IV-SC

ATROPINE SULFATE ANSYR PLASTIC SYRINGE
HOSPIRA 0.05MG/ML
+ 0.1MG/MLN021146 002 Jul 09, 2001
N021146 001 Jul 09, 2001ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

+ VALEANT 0.025MG;1MG

N017744 002

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

<u>AA</u>	ROXANE	<u>0.025MG/5ML;2.5MG/5ML</u>	<u>A087708 001</u>	May 03, 1982
<u>AA</u>	<u>LOMOTIL</u>			
<u>AA</u> +	GD SEARLE LLC	<u>0.025MG/5ML;2.5MG/5ML</u>	<u>N012699 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 44 (of 424)

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

<u>AA</u>	LANNETT	<u>0.025MG;2.5MG</u>	<u>A085372</u>	<u>001</u>	
<u>AA</u>	MYLAN	<u>0.025MG;2.5MG</u>	<u>A085762</u>	<u>001</u>	
<u>AA</u>	PAR PHARM	<u>0.025MG;2.5MG</u>	<u>A040357</u>	<u>001</u>	May 02, 2000
	<u>LOMOTIL</u>				
<u>AA</u>	+ GD SEARLE LLC	<u>0.025MG;2.5MG</u>	<u>N012462</u>	<u>001</u>	
	<u>LONOX</u>				
<u>AA</u>	SANDOZ	<u>0.025MG;2.5MG</u>	<u>A085311</u>	<u>002</u>	

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON-PLUS

+ MYLAN INSTITUTIONAL 0.14MG/ML;10MG/ML

N019678 001 Nov 06, 1991

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

DUODOTE

+ MERIDIAN MEDCL 2.1MG/0.7ML;600MG/2ML

N021983 001 Sep 28, 2006

AURANOFIN

CAPSULE; ORAL

RIDAURA

+ PROMETHEUS LABS 3MG

N018689 001 May 24, 1985

AZACITIDINE

INJECTABLE; IV-SC

VIDAZA

+ CELGENE 100MG/VIAL

N050794 001 May 19, 2004

AZATHIOPRINE

TABLET; ORAL

AZASAN

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

AZATHIOPRINE

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

IMURAN

<u>AB</u>	+ PROMETHEUS LABS	<u>50MG</u>	<u>N016324</u>	<u>001</u>	
-----------	-------------------	-------------	----------------	------------	--

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

+ BEDFORD EQ 100MG BASE/VIAL

A074419 001 Mar 31, 1995

AZELAIC ACID

CREAM; TOPICAL

AZELEX

+ ALLERGAN 20%

N020428 001 Sep 13, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 45 (of 424)

AZELAIC ACID

GEL; TOPICAL
FINACEA
+ INTENDIS 15% N021470 001 Dec 24, 2002

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
AZELASTINE HYDROCHLORIDE
AT APOTEX INC 0.05% A078621 001 Aug 03, 2009
AT SUN PHARMA GLOBAL 0.05% A078738 001 Jun 21, 2010
OPTIVAR
AT + MEDA PHARMS 0.05% N021127 001 May 22, 2000
SPRAY, METERED; NASAL
ASTELIN
AB + MEDA PHARMS EQ 0.125MG BASE/SPRAY N020114 001 Nov 01, 1996
AZELASTINE HYDROCHLORIDE
AB APOTEX INC EQ 0.125MG BASE/SPRAY A077954 001 Apr 30, 2009
ASTEPRO
+ MEDA PHARMS EQ 0.1876MG BASE/SPRAY N022371 001 Aug 31, 2009

AZILSARTAN KAMEDOXOMIL

TABLET; ORAL
EDARBI
TAKEDA PHARMS EQ 40MG MEDOXOMIL N200796 001 Feb 25, 2011
+ EQ 80MG MEDOXOMIL N200796 002 Feb 25, 2011

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET; ORAL
EDARBYCLOR
TAKEDA PHARMS EQ 40MG MEDOXOMIL;12.5MG N202331 001 Dec 20, 2011
+ EQ 40MG MEDOXOMIL;25MG N202331 002 Dec 20, 2011

AZITHROMYCIN

FOR SUSPENSION; ORAL
AZITHROMYCIN
AB PLIVA EQ 100MG BASE/5ML A065246 002 Jul 05, 2006
AB EQ 200MG BASE/5ML A065246 001 Jul 05, 2006
AB SANDOZ EQ 100MG BASE/5ML A065297 001 Sep 18, 2006
AB EQ 200MG BASE/5ML A065297 002 Sep 18, 2006
AB TEVA PHARMS EQ 100MG BASE/5ML A065419 001 Jun 24, 2008
AB EQ 200MG BASE/5ML A065419 002 Jun 24, 2008
ZITHROMAX
AB PFIZER EQ 100MG BASE/5ML N050710 001 Oct 19, 1995
AB + EQ 200MG BASE/5ML N050710 002 Oct 19, 1995
ZITHROMAX
+ PFIZER EQ 1GM BASE/PACKET N050693 001 Sep 28, 1994

FOR SUSPENSION, EXTENDED RELEASE; ORAL

ZMAX
+ PFIZER GLOBAL EQ 2GM BASE/BOT N050797 001 Jun 10, 2005

INJECTABLE; INJECTION

AZITHROMYCIN
AP APP PHARMS EQ 500MG BASE/VIAL A065179 001 Dec 13, 2005
AP GLAND PHARMA LTD EQ 500MG BASE/VIAL A065501 001 Nov 09, 2009
AP HOSPIRA EQ 500MG BASE/VIAL A065500 001 Jun 26, 2009
AP EQ 500MG BASE/VIAL A065511 001 Jun 26, 2009
AP PLIVA HRVATSKA DOO EQ 500MG BASE/VIAL A065265 001 Jan 18, 2007
AP SAGENT STRIDES EQ 500MG BASE/VIAL A065506 001 Mar 24, 2009
AP + TEVA PARENTERAL EQ 500MG BASE/VIAL N050809 001 Dec 19, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 46 (of 424)

AZITHROMYCIN

INJECTABLE; INJECTION

ZITHROMAX

<u>AP</u>	+ PFIZER	<u>EQ 500MG BASE/VIAL</u>	<u>N050733 001</u>	Jan 30, 1997
	AZITHROMYCIN			
	+ TEVA PARENTERAL	EQ 2.5GM BASE/VIAL	N050809 002	Dec 19, 2006

SOLUTION/DROPS; OPHTHALMIC

+ INSPIRE	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>	MYLAN	<u>EQ 250MG BASE</u>	<u>A065365 001</u>	May 30, 2007
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065366 001</u>	May 30, 2007
<u>AB</u>		<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>AB</u>	<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

AZTREONAM

<u>AP</u>	APP PHARMS	<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
<u>AP</u>	BEDFORD	<u>1GM/VIAL</u>	<u>A065286 001</u>	Mar 23, 2011
<u>AP</u>		<u>2GM/VIAL</u>	<u>A065286 002</u>	Mar 23, 2011

AZACTAM IN PLASTIC CONTAINER

+ BRISTOL MYERS SQUIBB	20MG/ML	N050632 002	May 24, 1989
+	40MG/ML	N050632 001	May 24, 1989

AZTREONAM

APP PHARMS	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>AP</u>	<u>BACITRACIN</u>		<u>A065116 001</u>	Dec 03, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 47 (of 424)

BACITRACIN

INJECTABLE; INJECTION

BACITRACIN

<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
	BACITRACIN			
	PHARMACIA AND UPJOHN	10,000 UNITS/VIAL	A060733 001	
	OINTMENT; OPHTHALMIC			
	BACITRACIN			
	+ FERA PHARMS	500 UNITS/GM	A061212 001	
	POWDER; FOR RX COMPOUNDING			
	BACI-RX			
	X GEN PHARMS	5,000,000 UNITS/BOT	A061580 001	

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

+ BAUSCH AND LOMB	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A064068 001	Oct 30, 1995
+ MONARCH PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM	N050168 002	May 04, 1984

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>	FERA PHARMS	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A060764 002</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>	FERA PHARMS	<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

+ FERA PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062166 002
---------------	---	-------------

BACLOFEN

INJECTABLE; INTRATHECAL

GABLOFEN

<u>AP</u>	CNS THERAPS INC	<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
<u>AP</u>	+ MEDTRONIC	<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

TABLET; ORAL

BACLOFEN

<u>AB</u>	CARACO	<u>10MG</u>	<u>A077984 001</u>	Aug 14, 2006
-----------	--------	-------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 48 (of 424)

BACLOFEN

TABLET; ORAL

BACLOFEN

<u>AB</u>	CARACO	<u>20MG</u>	<u>A077862</u>	<u>002</u>	Aug 14, 2006
<u>AB</u>	IMPAX LABS	<u>10MG</u>	<u>A078146</u>	<u>001</u>	Oct 26, 2007
<u>AB</u>		<u>20MG</u>	<u>A077971</u>	<u>002</u>	Oct 26, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>10MG</u>	<u>A072234</u>	<u>001</u>	Jul 21, 1988
<u>AB</u>	+ LANNETT	<u>20MG</u>	<u>A072235</u>	<u>001</u>	Jul 21, 1988
<u>AB</u>	LANNETT	<u>10MG</u>	<u>A078220</u>	<u>001</u>	Jul 06, 2007
<u>AB</u>		<u>20MG</u>	<u>A077241</u>	<u>001</u>	Dec 20, 2005
<u>AB</u>	MATRIX LABS LTD	<u>10MG</u>	<u>A090334</u>	<u>001</u>	Feb 18, 2010
<u>AB</u>		<u>20MG</u>	<u>A090334</u>	<u>002</u>	Feb 18, 2010
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A077181</u>	<u>001</u>	Jul 29, 2005
<u>AB</u>		<u>20MG</u>	<u>A077121</u>	<u>002</u>	Jul 29, 2005
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078504</u>	<u>001</u>	Sep 18, 2009
<u>AB</u>		<u>20MG</u>	<u>A078401</u>	<u>001</u>	Sep 18, 2009
<u>AB</u>	PROSAM LABS	<u>10MG</u>	<u>A077089</u>	<u>001</u>	Oct 31, 2007
<u>AB</u>		<u>20MG</u>	<u>A077088</u>	<u>001</u>	Oct 31, 2007
<u>AB</u>	USL PHARMA	<u>10MG</u>	<u>A074584</u>	<u>001</u>	Aug 19, 1996
<u>AB</u>		<u>20MG</u>	<u>A074584</u>	<u>002</u>	Aug 19, 1996
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A077156</u>	<u>001</u>	Aug 30, 2005
<u>AB</u>		<u>20MG</u>	<u>A077068</u>	<u>001</u>	Aug 30, 2005
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A072824</u>	<u>001</u>	Sep 18, 1991
<u>AB</u>		<u>20MG</u>	<u>A072825</u>	<u>001</u>	Sep 18, 1991

BALSALAZIDE DISODIUM

CAPSULE; ORAL

BALSALAZIDE DISODIUM

<u>AB</u>	APOTEX INC	<u>750MG</u>	<u>A077883</u>	<u>001</u>	Dec 28, 2007
<u>AB</u>	MYLAN	<u>750MG</u>	<u>A077807</u>	<u>001</u>	Dec 28, 2007
<u>AB</u>	ROXANE	<u>750MG</u>	<u>A077806</u>	<u>001</u>	Dec 28, 2007
<u>AB</u>	<u>COLAZAL</u>				
<u>AB</u>	+ SALIX PHARMS	<u>750MG</u>	<u>N020610</u>	<u>001</u>	Jul 18, 2000

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

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED; NASAL

BECONASE AQ

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

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

<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>	BIOKEY	<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

PRESCRIPTION DRUG PRODUCT LIST

3 - 49 (of 424)

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

<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>	RANBAXY	<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>	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>	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>	WATSON LABS FLORIDA	<u>5MG</u>	<u>A076267</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076267</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076267</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076267</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
<u>LOTENSIN</u>					
<u>AB</u>	NOVARTIS	<u>5MG</u>	<u>N019851</u>	<u>001</u>	Jun 25, 1991
<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; HYDROCHLORTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLORTHIAZIDE

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>5MG; 6.25MG</u>	<u>A076348</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076348</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076348</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076348</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	MYLAN	<u>5MG; 6.25MG</u>	<u>A076612</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>A076688</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076612</u>	<u>002</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>A076612</u>	<u>003</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>A076612</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076688</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	RANBAXY	<u>5MG; 6.25MG</u>	<u>A077483</u>	<u>001</u>	Sep 08, 2005
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A077483</u>	<u>002</u>	Sep 08, 2005
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A077483</u>	<u>003</u>	Sep 08, 2005
<u>AB</u>		<u>20MG; 25MG</u>	<u>A077483</u>	<u>004</u>	Sep 08, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 50 (of 424)

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<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
<u>AB</u>	WATSON LABS FLORIDA	<u>5MG; 6.25MG</u>	<u>A076342</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>A076342</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>A076342</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>A076342</u>	<u>004</u>	Feb 11, 2004
	<u>LOTENSIN HCT</u>				
<u>AB</u>	NOVARTIS	<u>5MG; 6.25MG</u>	<u>N020033</u>	<u>001</u>	May 19, 1992
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N020033</u>	<u>002</u>	May 19, 1992
<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

BENDAMUSTINE HYDROCHLORIDE

POWDER; IV (INFUSION)

+ CEPHALON	25MG/VIAL	N022249	002	May 01, 2009
+ +	100MG/VIAL	N022249	001	Mar 20, 2008

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

CORZIDE

<u>AB</u>	KING PHARMS	<u>5MG; 40MG</u>	<u>N018647</u>	<u>001</u>	May 25, 1983
<u>AB</u>	+ +	<u>5MG; 80MG</u>	<u>N018647</u>	<u>002</u>	May 25, 1983
	<u>NADOLOL AND BENDROFLUMETHIAZIDE</u>				
<u>AB</u>	MYLAN	<u>5MG; 40MG</u>	<u>A078688</u>	<u>001</u>	Feb 15, 2008
<u>AB</u>		<u>5MG; 80MG</u>	<u>A078688</u>	<u>002</u>	Feb 15, 2008
	<u>NADOLOL AND BENDROFLUMETHIAZIDE</u>				
<u>AB</u>	IMPAK LABS	<u>5MG; 40MG</u>	<u>A077833</u>	<u>001</u>	Mar 30, 2007
<u>AB</u>		<u>5MG; 80MG</u>	<u>A077833</u>	<u>002</u>	Mar 30, 2007

BENZONATATE

CAPSULE; ORAL

BENZONATATE

<u>AA</u>	BANNER PHARMACAPS	<u>100MG</u>	<u>A081297</u>	<u>001</u>	Jan 29, 1993
<u>AA</u>		<u>200MG</u>	<u>A081297</u>	<u>002</u>	Oct 30, 2007
<u>AA</u>	MIKART	<u>100MG</u>	<u>A040851</u>	<u>001</u>	Nov 09, 2009
<u>AA</u>		<u>200MG</u>	<u>A040851</u>	<u>003</u>	Nov 09, 2009
<u>AA</u>	NESHER PHARMS	<u>100MG</u>	<u>A040795</u>	<u>001</u>	Oct 31, 2007
<u>AA</u>		<u>200MG</u>	<u>A040795</u>	<u>002</u>	Oct 31, 2007
<u>AA</u>	ORIT LABS LLC	<u>100MG</u>	<u>A040682</u>	<u>001</u>	Jul 30, 2007
<u>AA</u>		<u>200MG</u>	<u>A040682</u>	<u>002</u>	Jul 30, 2007
<u>AA</u>	SUN PHARM INDs INC	<u>100MG</u>	<u>A040587</u>	<u>001</u>	Mar 19, 2008
<u>AA</u>		<u>200MG</u>	<u>A040587</u>	<u>002</u>	Mar 19, 2008
<u>AA</u>	THE PHARMA NETWORK	<u>100MG</u>	<u>A040627</u>	<u>001</u>	Mar 30, 2007
<u>AA</u>		<u>200MG</u>	<u>A040749</u>	<u>001</u>	Jul 25, 2007
<u>AA</u>	ZYDUS PHARMS USA	<u>100MG</u>	<u>A040597</u>	<u>001</u>	Jun 08, 2007
<u>AA</u>		<u>200MG</u>	<u>A040597</u>	<u>002</u>	Jun 08, 2007
	<u>TESSALON</u>				
<u>AA</u>	+ FOREST LABS	<u>100MG</u>	<u>N011210</u>	<u>001</u>	
<u>AA</u>	+ +	<u>200MG</u>	<u>N011210</u>	<u>003</u>	Jun 25, 1999
	<u>BENZONATATE</u>				
	MIKART	150MG	A040851	002	Nov 09, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 51 (of 424)

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

<u>AB</u>	SANOFI AVENTIS US	<u>5%;EQ 1% BASE</u>	<u>N050756 001</u>	Dec 21, 2000
<u>AB</u>	<u>CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE</u>			
<u>AB</u>	DOW PHARM SCIENCES	<u>5%;EQ 1% BASE</u>	<u>A065443 001</u>	Aug 11, 2009
	BENZACLIN			
BT	+ SANOFI AVENTIS US	5%;EQ 1% BASE	N050756 002	Apr 20, 2007
	DUAC			
BT	+ STIEFEL	5%;EQ 1% BASE	N050741 001	Aug 26, 2002
	ACANYA			
	+ DOW PHARM SCI	2.5%;1.2%	N050819 001	Oct 23, 2008

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL

BENZAMYCIN

<u>AB</u>	+ SANOFI AVENTIS US	<u>5%;3%</u>	<u>N050557 001</u>	Oct 26, 1984
<u>AB</u>	<u>ERYTHROMYCIN AND BENZOYL PEROXIDE</u>			
<u>AB</u>	TOLMAR	<u>5%;3%</u>	<u>A065112 001</u>	Mar 29, 2004
	BENZAMYCIN PAK			
	+ SANOFI AVENTIS US	5%;3%	N050769 001	Nov 27, 2000

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

<u>AA</u>	COREPHARMA	<u>50MG</u>	<u>A040714 001</u>	Oct 29, 2007
<u>AA</u>	IMPAX LABS	<u>50MG</u>	<u>A040845 001</u>	Nov 18, 2008
<u>AA</u>	KVK TECH	<u>50MG</u>	<u>A090968 001</u>	Jul 20, 2010
<u>AA</u>	MALLINCKRODT INC	<u>50MG</u>	<u>A040773 001</u>	Apr 25, 2007
<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>	PADDOCK	<u>50MG</u>	<u>A040578 001</u>	Apr 17, 2006
<u>AA</u>	TEDOR PHARM	<u>50MG</u>	<u>A040747 001</u>	Mar 30, 2007
	<u>DIDREX</u>			
<u>AA</u>	+ PHARMACIA AND UPJOHN	<u>50MG</u>	<u>N012427 002</u>	

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

<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>	NEXUS PHARMS	<u>1MG/ML</u>	<u>A090233 001</u>	Jul 28, 2009
	<u>COGENTIN</u>			
<u>AP</u>	+ LUNDBECK INC	<u>1MG/ML</u>	<u>N012015 001</u>	

TABLET; ORAL

BENZTROPINE MESYLATE

<u>AA</u>	COREPHARMA	<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>	+ LANNETT	<u>0 .5MG</u>	<u>A088877 001</u>	Apr 11, 1985
<u>AA</u>	+	<u>1MG</u>	<u>A088894 001</u>	Apr 11, 1985
<u>AA</u>	+	<u>2MG</u>	<u>A088895 001</u>	Apr 11, 1985
<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>	USL PHARMA	<u>0 .5MG</u>	<u>A040103 001</u>	Dec 12, 1996
<u>AA</u>		<u>1MG</u>	<u>A040103 002</u>	Dec 12, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 52 (of 424)

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

<u>AA</u>	USL PHARMA	<u>2MG</u>	<u>A040103</u>	<u>003</u>	Dec 12, 1996
<u>AA</u>	VINTAGE	<u>0.5MG</u>	<u>A040738</u>	<u>001</u>	Aug 27, 2007
<u>AA</u>		<u>1MG</u>	<u>A040742</u>	<u>001</u>	Aug 27, 2007
<u>AA</u>		<u>2MG</u>	<u>A040715</u>	<u>003</u>	Aug 27, 2007

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

+ ISTA PHARMS 1.5%

N022288 001 Sep 08, 2009

BERACTANT

SUSPENSION; INTRATRACHEAL

SURVANTA

+ ROSS LABS 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

+ RARE DIS THERAP 1GM/SCOOPFUL

N020576 001 Oct 25, 1996

BETAMETHASONE

SYRUP; ORAL

CELESTONE

+ SCHERING 0.6MG/5ML

N014215 002

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE

<u>AB</u>	LUITPOLD	<u>3MG/ML;EQ 3MG BASE/ML</u>	<u>A090747</u>	<u>001</u>	Jul 31, 2009
<u>AB</u>	+ SCHERING	<u>3MG/ML;EQ 3MG BASE/ML</u>	<u>N014602</u>	<u>001</u>	

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.05% BASE</u>	<u>A070885</u>	<u>001</u>	Feb 03, 1987
<u>AB</u>	+ FOUGERA	<u>EQ 0.05% BASE</u>	<u>N019137</u>	<u>001</u>	Jun 26, 1984
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A073552</u>	<u>001</u>	Apr 30, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 53 (of 424)

BETAMETHASONE DIPROPIONATE

CREAM, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALTANA	<u>EQ 0.05% BASE</u>	<u>A076215</u>	<u>001</u>	Dec 09, 2003
<u>AB</u>	GLENMARK GENERICS	<u>EQ 0.05% BASE</u>	<u>A078930</u>	<u>001</u>	Sep 23, 2008
<u>AB</u>	PERRIGO NEW YORK	<u>EQ 0.05% BASE</u>	<u>A076592</u>	<u>001</u>	Dec 09, 2003
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A076543</u>	<u>001</u>	Dec 09, 2003
<u>AB</u>	TOLMAR	<u>EQ 0.05% BASE</u>	<u>A076603</u>	<u>001</u>	Jan 23, 2004

DIPROLENE AF

<u>AB</u>	+ SCHERING	<u>EQ 0.05% BASE</u>	<u>N019555</u>	<u>001</u>	Apr 27, 1987
-----------	------------	----------------------	----------------	------------	--------------

GEL, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	+ ALTANA	<u>EQ 0.05% BASE</u>	<u>A075276</u>	<u>001</u>	May 13, 2003
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A076508</u>	<u>001</u>	Dec 02, 2003

LOTION; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.05% BASE</u>	<u>A070281</u>	<u>001</u>	Jul 31, 1985
<u>AB</u>	+ FOUGERA	<u>EQ 0.05% BASE</u>	<u>A070275</u>	<u>001</u>	Aug 12, 1985
<u>AB</u>	PERRIGO NEW YORK	<u>EQ 0.05% BASE</u>	<u>A072538</u>	<u>001</u>	Jan 31, 1990
<u>AB</u>	TEVA	<u>EQ 0.05% BASE</u>	<u>A071467</u>	<u>001</u>	Aug 10, 1987

LOTION, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALTANA	<u>EQ 0.05% BASE</u>	<u>A077111</u>	<u>001</u>	May 21, 2007
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A077477</u>	<u>001</u>	May 21, 2007
<u>AB</u>	+ SCHERING	<u>EQ 0.05% BASE</u>	<u>N019716</u>	<u>001</u>	Aug 01, 1988

OINTMENT; TOPICAL

BETAMETHASONE DIPROPIONATE

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

OINTMENT, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.05% BASE</u>	<u>A074304</u>	<u>001</u>	Aug 31, 1995
<u>AB</u>	ALTANA	<u>EQ 0.05% BASE</u>	<u>A075373</u>	<u>001</u>	Jun 22, 1999
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A076753</u>	<u>001</u>	Oct 12, 2004
<u>AB</u>	+ SCHERING	<u>EQ 0.05% BASE</u>	<u>N018741</u>	<u>001</u>	Jul 27, 1983

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE

OINTMENT; TOPICAL

TACLONEX

+ LEO PHARM	0.064%;0.005%	<u>N021852</u>	<u>001</u>	Jan 09, 2006
-------------	---------------	----------------	------------	--------------

SUSPENSION; TOPICAL

TACLONEX SCALP

+ LEO PHARM PRODS	0.064%;0.005%	<u>N022185</u>	<u>001</u>	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</u>	<u>001</u>	Aug 02, 2002
<u>AB</u>	ALTANA	<u>EQ 0.05% BASE;1%</u>	<u>A075502</u>	<u>001</u>	Jun 05, 2001
<u>AB</u>	TARO	<u>EQ 0.05% BASE;1%</u>	<u>A075673</u>	<u>001</u>	May 29, 2001

LOTRISONE

<u>AB</u>	+ SCHERING	<u>EQ 0.05% BASE;1%</u>	<u>N018827</u>	<u>001</u>	Jul 10, 1984
-----------	------------	-------------------------	----------------	------------	--------------

LOTION; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALTANA PHARMA	<u>EQ 0.05% BASE;1%</u>	<u>A076516</u>	<u>001</u>	Jun 16, 2005
-----------	---------------	-------------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 54 (of 424)

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

LOTION; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	TARO	<u>EQ 0.05% BASE;1%</u>	<u>A076493 001</u>	Jul 28, 2004
<u>AB</u>	+ SCHERING PLOUGH RES	<u>EQ 0.05% BASE;1%</u>	<u>N020010 001</u>	Dec 08, 2000

BETAMETHASONE VALERATE

AEROSOL, FOAM; TOPICAL

LUXIQ

+ CONNECTICS	<u>EQ 0.12% BASE</u>	<u>N020934 001</u>	Feb 28, 1999
--------------	----------------------	--------------------	--------------

CREAM; TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	+ FOUGERA	<u>EQ 0.1% BASE</u>	<u>N018861 001</u>	Aug 31, 1983
<u>AB</u>	<u>BETA-VAL</u>	<u>EQ 0.1% BASE</u>	<u>N018642 001</u>	Mar 24, 1983
<u>AB</u>	TEVA	<u>EQ 0.1% BASE</u>	<u>A072041 001</u>	Jan 06, 1988
<u>AB</u>	<u>DERMABET</u>	<u>EQ 0.1% BASE</u>	<u>A070050 001</u>	Oct 10, 1984
<u>AB</u>	TARO	<u>EQ 0.1% BASE</u>		
<u>AB</u>	<u>VALNAC</u>	<u>EQ 0.1% BASE</u>		
<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.1% BASE</u>		

LOTION; TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	+ FOUGERA	<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
<u>AB</u>	<u>BETA-VAL</u>	<u>EQ 0.1% BASE</u>	<u>A070072 001</u>	Jun 27, 1985

OINTMENT; TOPICAL

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	<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>	BAUSCH AND LOMB	<u>EQ 0.5% BASE</u>	<u>A075630 001</u>	Apr 12, 2001
<u>AT</u>	NOVEX	<u>EQ 0.5% BASE</u>	<u>A075446 001</u>	Sep 28, 2000
<u>AT</u>	WOCKHARDT	<u>EQ 0.5% BASE</u>	<u>A078694 001</u>	Nov 16, 2009
<u>AT</u>	<u>BETOPTIC</u>			
<u>AT</u>	+ ALCON	<u>EQ 0.5% BASE</u>	<u>N019270 001</u>	Aug 30, 1985

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC S

+ ALCON	<u>EQ 0.25% BASE</u>	<u>N019845 001</u>	Dec 29, 1989
---------	----------------------	--------------------	--------------

TABLET; ORAL

BETAXOLOL HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA	<u>10MG</u>	<u>A075541 001</u>	Oct 22, 1999
<u>AB</u>	+ KVK TECH	<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>	EMCURE USA	<u>5MG</u>	<u>A091256 001</u>	May 04, 2010
<u>AA</u>		<u>10MG</u>	<u>A091256 002</u>	May 04, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 55 (of 424)

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

<u>AA</u>	EMCURE USA	<u>25MG</u>	<u>A091256</u> <u>003</u>	May 04, 2010
<u>AA</u>		<u>50MG</u>	<u>A091256</u> <u>004</u>	May 04, 2010
<u>AA</u>	IMPAK LABS	<u>5MG</u>	<u>A040739</u> <u>001</u>	Nov 01, 2006
<u>AA</u>		<u>10MG</u>	<u>A040741</u> <u>001</u>	Nov 01, 2006
<u>AA</u>		<u>25MG</u>	<u>A040740</u> <u>001</u>	Nov 01, 2006
<u>AA</u>		<u>50MG</u>	<u>A040721</u> <u>004</u>	Nov 01, 2006
<u>AA</u>	LANNETT	<u>5MG</u>	<u>A040703</u> <u>001</u>	Mar 27, 2008
<u>AA</u>		<u>10MG</u>	<u>A040704</u> <u>001</u>	Mar 27, 2008
<u>AA</u>		<u>25MG</u>	<u>A040678</u> <u>003</u>	Mar 27, 2008
<u>AA</u>		<u>50MG</u>	<u>A040677</u> <u>001</u>	Mar 27, 2008
<u>AA</u>	PHARMAX	<u>5MG</u>	<u>A040725</u> <u>001</u>	Oct 26, 2007
<u>AA</u>		<u>10MG</u>	<u>A040726</u> <u>001</u>	Oct 26, 2007
<u>AA</u>		<u>25MG</u>	<u>A040727</u> <u>001</u>	Oct 26, 2007
<u>AA</u>		<u>50MG</u>	<u>A040728</u> <u>001</u>	Oct 26, 2007
<u>AA</u>	SUN PHARM INDs INC	<u>5MG</u>	<u>A040897</u> <u>001</u>	Apr 22, 2009
<u>AA</u>		<u>10MG</u>	<u>A040897</u> <u>002</u>	Apr 22, 2009
<u>AA</u>		<u>25MG</u>	<u>A040897</u> <u>003</u>	Apr 22, 2009
<u>AA</u>		<u>50MG</u>	<u>A040897</u> <u>004</u>	Apr 22, 2009
<u>AA</u>	UPSHER SMITH	<u>5MG</u>	<u>A040633</u> <u>001</u>	Jun 01, 2005
<u>AA</u>		<u>10MG</u>	<u>A040634</u> <u>001</u>	Jun 01, 2005
<u>AA</u>		<u>25MG</u>	<u>A040635</u> <u>001</u>	Jun 01, 2005
<u>AA</u>		<u>50MG</u>	<u>A040636</u> <u>001</u>	Jun 01, 2005
<u>AA</u>	WOCKHARDT	<u>5MG</u>	<u>A040532</u> <u>001</u>	Sep 29, 2003
<u>AA</u>		<u>10MG</u>	<u>A040533</u> <u>001</u>	Sep 29, 2003
<u>AA</u>		<u>25MG</u>	<u>A040534</u> <u>001</u>	Sep 29, 2003
<u>AA</u>		<u>50MG</u>	<u>A040518</u> <u>001</u>	Sep 29, 2003
<u>DUVOID</u>				
<u>AA</u>	WELSPRING PHARM	<u>10MG</u>	<u>A086262</u> <u>001</u>	
<u>AA</u>		<u>25MG</u>	<u>A086263</u> <u>001</u>	
<u>AA</u>		<u>50MG</u>	<u>A085882</u> <u>003</u>	
<u>URECHOLINE</u>				
<u>AA</u>	+ ODYSSEY PHARMS	<u>5MG</u>	<u>A089095</u> <u>001</u>	Dec 19, 1985
<u>AA</u>	+	<u>10MG</u>	<u>A088440</u> <u>001</u>	May 29, 1984
<u>AA</u>	+	<u>25MG</u>	<u>A088441</u> <u>001</u>	May 29, 1984
<u>AA</u>	+	<u>50MG</u>	<u>A089096</u> <u>001</u>	Dec 19, 1985

BEXAROTENE

CAPSULE; ORAL

TARGRETIN

+ EISAI INC

75MG

N021055 001 Dec 29, 1999

GEL; TOPICAL

TARGRETIN

+ EISAI INC

1%

N021056 001 Jun 28, 2000

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

<u>AB</u>	ACCORD HLTHCARE INC	<u>50MG</u>	<u>A078917</u> <u>001</u>	Jul 06, 2009
<u>AB</u>	ACTAVIS TOTOWA	<u>50MG</u>	<u>A078634</u> <u>001</u>	Aug 28, 2009
<u>AB</u>	FRESENIUS KABI ONCOL	<u>50MG</u>	<u>A079045</u> <u>001</u>	May 13, 2010
<u>AB</u>	KUDCO IRELAND	<u>50MG</u>	<u>A077995</u> <u>001</u>	Jul 06, 2009
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A079185</u> <u>001</u>	Jul 06, 2009
<u>AB</u>	ROXANE	<u>50MG</u>	<u>A078285</u> <u>001</u>	Mar 24, 2011
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>A078575</u> <u>001</u>	Jul 06, 2009
<u>AB</u>	SUN PHARMA GLOBAL	<u>50MG</u>	<u>A079110</u> <u>001</u>	Jul 06, 2009
<u>AB</u>	SYNTHON PHARMS	<u>50MG</u>	<u>A077973</u> <u>001</u>	Jul 06, 2009
<u>AB</u>	TEVA	<u>50MG</u>	<u>A076932</u> <u>001</u>	Jul 06, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 56 (of 424)

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

<u>AB</u>	ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A079089</u>	<u>001</u>	Jul 06, 2009
	CASODEX				

<u>AB</u>	ASTRAZENECA	<u>50MG</u>	<u>N020498</u>	<u>001</u>	Oct 04, 1995
-----------	-------------	-------------	----------------	------------	--------------

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC

LUMIGAN

+ ALLERGAN	0.01%	N022184	001	Aug 31, 2010
+	0.03%	N021275	001	Mar 16, 2001

SOLUTION/DROPS; TOPICAL

+ ALLERGAN	0.03%	N022369	001	Dec 24, 2008
------------	-------	---------	-----	--------------

BIPERIDEN HYDROCHLORIDE

TABLET; ORAL

AKINETON

+ ABBOTT	2MG	N012003	001
----------	-----	---------	-----

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

FOR SOLUTION; TABLET, DELAYED RELEASE; ORAL

HALFLYTETLY

+ 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 SUBLITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE

CAPSULE; ORAL

PYLERA

+ APTALIS PHARMA US	140MG;125MG;125MG	N050786	001	Sep 28, 2006
---------------------	-------------------	---------	-----	--------------

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL

HEЛИДАК

+ PROMETHEUS LABS	262.4MG,N/A,N/A;N/A,250MG,N/A;N/A,N/A,5 00MG	N050719	001	Aug 15, 1996
-------------------	---	---------	-----	--------------

BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A077910</u>	<u>001</u>	Dec 27, 2006
<u>AB</u>		<u>10MG</u>	<u>A077910</u>	<u>002</u>	Dec 27, 2006
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A075831</u>	<u>001</u>	Dec 14, 2005
<u>AB</u>		<u>10MG</u>	<u>A075831</u>	<u>002</u>	Dec 14, 2005
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A075643</u>	<u>001</u>	Nov 16, 2000
<u>AB</u>		<u>10MG</u>	<u>A075643</u>	<u>002</u>	Nov 16, 2000
<u>AB</u>	TEVA PHARMS	<u>5MG</u>	<u>A075644</u>	<u>001</u>	Jun 26, 2001
<u>AB</u>		<u>10MG</u>	<u>A075644</u>	<u>002</u>	Jun 26, 2001
<u>AB</u>	UNICHEM PHARMS (USA)	<u>5MG</u>	<u>A078635</u>	<u>001</u>	Aug 18, 2009
<u>AB</u>		<u>10MG</u>	<u>A078635</u>	<u>002</u>	Aug 18, 2009
	ZEBETA				
<u>AB</u>	DURAMED PHARMS BARR	<u>5MG</u>	<u>N019982</u>	<u>002</u>	Jul 31, 1992
<u>AB</u>	+	<u>10MG</u>	<u>N019982</u>	<u>001</u>	Jul 31, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 57 (of 424)

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	MYLAN	<u>2.5MG;6.25MG</u>	<u>A075768</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG;6.25MG</u>	<u>A075768</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG;6.25MG</u>	<u>A075768</u>	<u>003</u>	Sep 25, 2000
<u>AB</u>	SANDOZ	<u>2.5MG;6.25MG</u>	<u>A075579</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG;6.25MG</u>	<u>A075579</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG;6.25MG</u>	<u>A075579</u>	<u>003</u>	Sep 25, 2000
<u>AB</u>	UNICHEM	<u>2.5MG;6.25MG</u>	<u>A079106</u>	<u>001</u>	Jul 28, 2010
<u>AB</u>		<u>5MG;6.25MG</u>	<u>A079106</u>	<u>002</u>	Jul 28, 2010
<u>AB</u>		<u>10MG;6.25MG</u>	<u>A079106</u>	<u>003</u>	Jul 28, 2010
<u>AB</u>	WATSON LABS	<u>2.5MG;6.25MG</u>	<u>A075469</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG;6.25MG</u>	<u>A075469</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG;6.25MG</u>	<u>A075469</u>	<u>003</u>	Sep 25, 2000
<u>ZIAC</u>					
<u>AB</u>	DURAMED PHARMS BARR	<u>2.5MG;6.25MG</u>	<u>N020186</u>	<u>003</u>	Mar 26, 1993
<u>AB</u>		<u>5MG;6.25MG</u>	<u>N020186</u>	<u>001</u>	Mar 26, 1993
<u>AB</u>	+	<u>10MG;6.25MG</u>	<u>N020186</u>	<u>002</u>	Mar 26, 1993

BIVALIRUDIN

INJECTABLE; INTRAVENOUS

ANGIOMAX

+ MEDICINES CO

250MG/VIAL

N020873 001 Dec 15, 2000

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN SULFATE

<u>AP</u>	+ APP PHARMS	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065185</u>	<u>001</u>	Jan 28, 2008
<u>AP</u>	+	<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065185</u>	<u>002</u>	Jan 28, 2008
<u>AP</u>	BEDFORD	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065042</u>	<u>002</u>	Oct 17, 2001
<u>AP</u>		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065042</u>	<u>001</u>	Oct 17, 2001
<u>AP</u>	HOSPIRA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065031</u>	<u>001</u>	Mar 10, 2000
<u>AP</u>		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065031</u>	<u>002</u>	Mar 10, 2000
<u>AP</u>	PHARMACHEMIE BV	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065201</u>	<u>001</u>	Dec 13, 2007
<u>AP</u>	TEVA PARENTERAL	<u>EQ 15 UNITS BASE/VIAL</u>	<u>A065033</u>	<u>001</u>	Jun 27, 2000
<u>AP</u>		<u>EQ 30 UNITS BASE/VIAL</u>	<u>A065033</u>	<u>002</u>	Jun 27, 2000

BOCEPREVIR

CAPSULE; ORAL

VICTRELIS

+ SCHERING

200MG

N202258 001 May 13, 2011

BORTEZOMIB

INJECTABLE; INTRAVENOUS

VELCADE

+ MILLENNIUM PHARMS

3.5MG/VIAL

N021602 001 May 13, 2003

BOSENTAN

TABLET; ORAL

TRACLEER

ACTELION

62.5MG

N021290 001 Nov 20, 2001

+

125MG

N021290 002 Nov 20, 2001

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ B BRAUN

100MG/100ML

N019121 001 Apr 29, 1986

+

200MG/100ML

N019121 002 Apr 29, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 58 (of 424)

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER
+ B BRAUN 400MG/100ML

N019121 003 Apr 29, 1986

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

<u>ALPHAGAN P</u>				
<u>AT + ALLERGAN</u>	<u>0.15%</u>		<u>N021262 001</u>	Mar 16, 2001
<u>BRIMONIDINE TARTRATE</u>				
<u>AT AKORN</u>	<u>0.2%</u>		<u>A076439 001</u>	Mar 14, 2006
<u>AT ALCON PHARMS LTD</u>	<u>0.15%</u>		<u>N021764 001</u>	May 22, 2006
<u>AT</u>	<u>0.2%</u>		<u>A076254 001</u>	Sep 16, 2003
<u>AT + BAUSCH AND LOMB</u>	<u>0.2%</u>		<u>A076260 001</u>	May 28, 2003
<u>AT SANDOZ</u>	<u>0.2%</u>		<u>A078075 001</u>	Jan 30, 2008
ALPHAGAN P				
+ ALLERGAN	0.1%		N021770 001	Aug 19, 2005

BRIMONIDINE TARTRATE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COMBIGAN

+ ALLERGAN 0.2%; 0.5%

N021398 001 Oct 30, 2007

BRINZOLAMIDE

SUSPENSION/DROPS; OPHTHALMIC

AZOPT

+ ALCON PHARMS LTD 1%

N020816 001 Apr 01, 1998

BROMFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC

BROMDAY

+ ISTA PHARMS INC 0.09%
BROMFENAC SODIUM
COASTAL PHARMS 0.09%N021664 002 Oct 16, 2010
A201211 001 May 11, 2011BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

<u>BROMOCRIPTINE MESYLATE</u>				
<u>AB MYLAN</u>	<u>EQ 5MG BASE</u>		<u>A077226 001</u>	Apr 04, 2005
<u>AB ZYDUS PHARMS USA INC</u>	<u>EQ 5MG BASE</u>		<u>A078899 001</u>	Jul 30, 2008
<u>PARLODEL</u>				
<u>AB + NOVARTIS</u>	<u>EQ 5MG BASE</u>		<u>N017962 002</u>	Mar 01, 1982
TABLET; ORAL				
<u>BROMOCRIPTINE MESYLATE</u>				
<u>AB LEK PHARMS</u>	<u>EQ 2.5MG BASE</u>		<u>A074631 001</u>	Jan 13, 1998
<u>AB MYLAN</u>	<u>EQ 2.5MG BASE</u>		<u>A076962 001</u>	Sep 24, 2004
<u>AB PADDOCK LLC</u>	<u>EQ 2.5MG BASE</u>		<u>A077646 001</u>	Oct 01, 2008
<u>PARLODEL</u>				
<u>AB + NOVARTIS</u>	<u>EQ 2.5MG BASE</u>		<u>N017962 001</u>	
CYCLOSET				
+ VEROSCIENCE	EQ 0.8MG BASE		N020866 001	May 05, 2009

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMFED-DM

+ WOCKHARDT 2MG/5ML; 10MG/5ML; 30MG/5ML

A088811 001 Jun 07, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 59 (of 424)

BUDESONIDE

CAPSULE; ORAL

BUDESONIDE

<u>AB</u>	MYLAN	<u>3MG</u>	<u>A090410</u>	<u>001</u>	May 16, 2011
		<u>ENTOCORT EC</u>			
<u>AB</u>	+ ASTRazeneca	<u>3MG</u>	<u>N021324</u>	<u>001</u>	Oct 02, 2001
	POWDER, METERED; INHALATION				
	PULMICORT				
	+ ASTRazeneca	0.16MG/INH	N020441	002	Jun 24, 1997
	PULMICORT FLEXHALER				
	ASTRazeneca	0.08MG/INH	N021949	001	Jul 12, 2006
	+	0.16MG/INH	N021949	002	Jul 12, 2006
	SPRAY, METERED; NASAL				
	RHINOCORT				
	+ ASTRazeneca	0.032MG/INH	N020746	001	Oct 01, 1999
	SUSPENSION; INHALATION				
	<u>BUDESONIDE</u>				
<u>AN</u>	APOTEX	<u>0.25MG/2ML</u>	<u>A078202</u>	<u>001</u>	Mar 30, 2009
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A078202</u>	<u>002</u>	Mar 30, 2009
<u>AN</u>	TEVA PARENTERAL	<u>0.25MG/2ML</u>	<u>A077519</u>	<u>001</u>	Nov 18, 2008
<u>AN</u>		<u>0.5MG/2ML</u>	<u>A077519</u>	<u>002</u>	Nov 18, 2008
	<u>PULMICORT RESPULES</u>				
<u>AN</u>	ASTRazeneca	<u>0.25MG/2ML</u>	<u>N020929</u>	<u>001</u>	Aug 08, 2000
<u>AN</u>		<u>0.5MG/2ML</u>	<u>N020929</u>	<u>002</u>	Aug 08, 2000
	PULMICORT RESPULES				
	+ ASTRazeneca	1MG/2ML	N020929	003	Aug 08, 2000

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

SPRAY, METERED; INHALATION

SYMBICORT

+ ASTRazeneca	0.08MG/INH; 0.0045MG/INH	N021929	001	Jul 21, 2006
+	0.16MG/INH; 0.0045MG/INH	N021929	002	Jul 21, 2006

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

<u>AP</u>	BAXTER HLTHCARE	<u>0.25MG/ML</u>	<u>A079196</u>	<u>001</u>	Apr 30, 2008
<u>AP</u>	+ BEDFORD	<u>0.25MG/ML</u>	<u>A074441</u>	<u>001</u>	Jan 27, 1995
<u>AP</u>	HOSPIRA	<u>0.25MG/ML</u>	<u>A074332</u>	<u>001</u>	Oct 31, 1994

TABLET; ORAL

BUMETANIDE

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>0.5MG</u>	<u>A074225</u>	<u>001</u>	Apr 24, 1995
<u>AB</u>		<u>1MG</u>	<u>A074225</u>	<u>002</u>	Apr 24, 1995
<u>AB</u>		<u>2MG</u>	<u>A074225</u>	<u>003</u>	Apr 24, 1995
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A074700</u>	<u>001</u>	Nov 21, 1996
<u>AB</u>		<u>1MG</u>	<u>A074700</u>	<u>002</u>	Nov 21, 1996
<u>AB</u>	+	<u>2MG</u>	<u>A074700</u>	<u>003</u>	Nov 21, 1996

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION

EXPAREL

+ PACIRA PHARMS INC	133MG/10ML (13.3MG/ML)	N022496	001	Oct 28, 2011
+	266MG/20ML (13.3MG/ML)	N022496	002	Oct 28, 2011

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>0.25%</u>	<u>A070583</u>	<u>001</u>	Feb 17, 1987
<u>AP</u>		<u>0.25%</u>	<u>A070586</u>	<u>001</u>	Mar 03, 1987

PREScription DRUG PRODUCT LIST

3 - 60 (of 424)

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

<u>BUPIVACAINE HYDROCHLORIDE</u>			
<u>AP</u>	HOSPIRA	<u>0.25%</u>	A070590 001 Feb 17, 1987
<u>AP</u>		<u>0.25%</u>	N018053 002
<u>AP</u>		<u>0.5%</u>	A070584 001 Feb 17, 1986
<u>AP</u>		<u>0.5%</u>	A070597 001 Mar 03, 1987
<u>AP</u>		<u>0.5%</u>	A070609 001 Mar 03, 1987
<u>AP</u>		<u>0.5%</u>	N018053 001
<u>AP</u>		<u>0.75%</u>	A070585 001 Mar 03, 1987
<u>AP</u>		<u>0.75%</u>	A070587 001 Mar 03, 1987
<u>AP</u>		<u>0.75%</u>	N018053 003
<u>AP</u>	SAGENT STRIDES	<u>0.25%</u>	A091503 001 Oct 18, 2011
<u>AP</u>		<u>0.5%</u>	A091503 002 Oct 18, 2011

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>BUTYVALINE HYDROCHLORIDE PRESERVATIVE FREE</u>			
<u>AP</u>	INTL MEDICATED	<u>0.25%</u>	A076012 001 Jan 09, 2002
<u>AP</u>		<u>0.5%</u>	A076012 002 Jan 09, 2002
<u>AP</u>		<u>0.75%</u>	A076012 003 Jan 09, 2002
<u>AP</u>	SAGENT STRIDES	<u>0.25%</u>	A091487 002 Oct 18, 2011
<u>AP</u>		<u>0.5%</u>	A091487 001 Oct 18, 2011
<u>AP</u>		<u>0.75%</u>	A091487 003 Oct 18, 2011

MARCATINE: HYDROCHLORIDE

AP + HOSPIRA 0.25% N016964 001
AP + 0.5% N016964 006

MARCAINE HYDROCHLORIDE PRESERVATIVE FREE

AP + HOSPIRA 0.25% N016964 012
AP + 0.5% N016964 005
AP + 0.75% N016964 009

SENSORCAINE

<u>AP</u>	APP PHARMS	<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: SRTNAT

BUTYVACATNE HYDROCHLORIDE

AP HOSPIRA 0.75% A071810 001 Dec 11, 1987
MARCAINE

+ HOSPIRA

AP APP PHARMS **0.75%** **A071202 001** Apr 15, 1987

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>	+ HOSPIRA	<u>0.5%;0.005MG/ML</u>	<u>A071168</u>	<u>001</u>	Jun 16, 1988
<u>AP</u>		<u>0.5%;0.005MG/ML</u>	<u>A071170</u>	<u>001</u>	Jun 16, 1988
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE					
+ HOSPIRA		0.25%;0.005MG/ML	A071165	001	Jun 16, 1988
		0.25%;0.005MG/ML	A071167	001	Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

<u>AP</u>	SEPTODONT	<u>0.5%;0.0091MG/ML</u>	<u>A077250</u>	<u>001</u>	Sep 27, 2006
	<u>BUPIVACAIN</u>	<u>E HYDROCHLORIDE W/EPINEPHRINE</u>			
<u>AP</u>	+ HOSPIRA	<u>0.5%;0.0091MG/ML</u>	<u>N022046</u>	<u>001</u>	Jul 13, 1983
		<u>MARCAINE HYDROCHLORIDE W/ EPINEPHRINE</u>			
<u>AP</u>	+ HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N016964</u>	<u>004</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 61 (of 424)

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

MARCaine HYDROCHLORIDE W/ EPINEPHRINE

<u>AP</u>	+ HOSPIRA	<u>0.5%;0.0091MG/ML</u>	<u>N016964</u>	<u>008</u>	
		<u>MARCaine HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE</u>			
<u>AP</u>	+ HOSPIRA	<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>	APP PHARMS	<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

BUPRENORPHINEFILM, EXTENDED RELEASE; TRANSDERMAL
BUTRANS

PURDUE PHARMA LP	5MCG/HR	N021306	001	Jun 30, 2010
	10MCG/HR	N021306	002	Jun 30, 2010
+	20MCG/HR	N021306	003	Jun 30, 2010

BUPRENORPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPRENEK

<u>AP</u>	+ RECKITT BENCKISER	<u>EQ 0.3MG BASE/ML</u>	<u>N018401</u>	<u>001</u>	
		<u>BUPRENORPHINE HYDROCHLORIDE</u>			
<u>AP</u>	BEDFORD	<u>EQ 0.3MG BASE/ML</u>	<u>A076931</u>	<u>001</u>	Mar 02, 2005
<u>AP</u>	HOSPIRA	<u>EQ 0.3MG BASE/ML</u>	<u>A074137</u>	<u>001</u>	Jun 03, 1996
<u>AP</u>	LUITPOLD	<u>EQ 0.3MG BASE/ML</u>	<u>A078331</u>	<u>001</u>	Mar 27, 2007

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>EQ 2MG BASE</u>	<u>A090360</u>	<u>001</u>	May 07, 2010
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090360</u>	<u>002</u>	May 07, 2010
<u>AB</u>	ETHYPHARM	<u>EQ 2MG BASE</u>	<u>A090622</u>	<u>001</u>	Sep 24, 2010
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A090622</u>	<u>002</u>	Sep 24, 2010
<u>AB</u>	ROXANE	<u>EQ 2MG BASE</u>	<u>A078633</u>	<u>001</u>	Oct 08, 2009
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078633</u>	<u>002</u>	Oct 08, 2009
		<u>SUBUTEX</u>			
<u>AB</u>	RECKITT BENCKISER	<u>EQ 2MG BASE</u>	<u>N020732</u>	<u>002</u>	Oct 08, 2002
<u>AB</u>	+	<u>EQ 8MG BASE</u>	<u>N020732</u>	<u>003</u>	Oct 08, 2002

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

TABLET; SUBLINGUAL

SUBOXONE

RECKITT BENCKISER	2MG; 0.5MG	N020733	001	Oct 08, 2002
+	8MG; 2MG	N020733	002	Oct 08, 2002

BUPRENORPHINE; NALOXONE

FILM; SUBLINGUAL

SUBOXONE

RECKITT BENCKISER	2MG; 0.5MG	N022410	001	Aug 30, 2010
+	8MG; 2MG	N022410	002	Aug 30, 2010

BUPROPION HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

APLENZIN

VALEANT INTL	174MG	N022108	001	Apr 23, 2008
+	348MG	N022108	002	Apr 23, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 62 (of 424)

BUPROPION HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL
 APLENZIN
 VALEANT INTL 522MG

N022108 003 Apr 23, 2008

BUPROPION HYDROCHLORIDE

TABLET; ORAL

BUPROPION HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>75MG</u>	<u>A076143</u> <u>001</u>	Jan 17, 2006
<u>AB</u>		<u>100MG</u>	<u>A076143</u> <u>002</u>	Jan 17, 2006
<u>AB</u>	MYLAN	<u>75MG</u>	<u>A075491</u> <u>001</u>	Apr 17, 2000
<u>AB</u>		<u>100MG</u>	<u>A075491</u> <u>002</u>	Apr 17, 2000
<u>AB</u>	SANDOZ	<u>75MG</u>	<u>A075584</u> <u>001</u>	Feb 07, 2000
<u>AB</u>		<u>100MG</u>	<u>A075584</u> <u>002</u>	Feb 07, 2000
<u>AB</u>	TEVA	<u>75MG</u>	<u>A075310</u> <u>001</u>	Nov 29, 1999
<u>AB</u>		<u>100MG</u>	<u>A075310</u> <u>002</u>	Nov 29, 1999
<u>WELLBUTRIN</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>75MG</u>	<u>N018644</u> <u>002</u>	Dec 30, 1985
<u>AB</u> +		<u>100MG</u>	<u>N018644</u> <u>003</u>	Dec 30, 1985

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

<u>AB1</u>	ACTAVIS	<u>100MG</u>	<u>A077455</u> <u>001</u>	Jul 19, 2010
<u>AB1</u>		<u>150MG</u>	<u>A077455</u> <u>002</u>	Mar 12, 2008
<u>AB1</u>		<u>200MG</u>	<u>A077455</u> <u>003</u>	Jul 19, 2010
<u>AB1</u>	ANCHEN PHARMS	<u>100MG</u>	<u>A091459</u> <u>001</u>	Jun 09, 2011
<u>AB1</u>		<u>150MG</u>	<u>A091459</u> <u>002</u>	Jun 09, 2011
<u>AB1</u>		<u>200MG</u>	<u>A091459</u> <u>003</u>	Jun 09, 2011
<u>AB1</u>	IMPAK LABS	<u>100MG</u>	<u>A075913</u> <u>001</u>	Jan 28, 2004
<u>AB1</u>		<u>150MG</u>	<u>A075913</u> <u>002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A076711</u> <u>001</u>	Dec 03, 2004
<u>AB1</u>	MYLAN	<u>100MG</u>	<u>A090325</u> <u>001</u>	Apr 08, 2010
<u>AB1</u>		<u>150MG</u>	<u>A090325</u> <u>002</u>	Apr 08, 2010
<u>AB1</u>		<u>200MG</u>	<u>A090325</u> <u>003</u>	Apr 08, 2010
<u>AB1</u>	SANDOZ	<u>100MG</u>	<u>A075932</u> <u>001</u>	Nov 25, 2003
<u>AB1</u>		<u>150MG</u>	<u>A075932</u> <u>002</u>	Mar 22, 2004
<u>AB1</u>		<u>200MG</u>	<u>A075932</u> <u>003</u>	Jun 22, 2005
<u>AB1</u>	SUN PHARMA GLOBAL	<u>100MG</u>	<u>A078866</u> <u>001</u>	Apr 06, 2010
<u>AB1</u>		<u>150MG</u>	<u>A078866</u> <u>002</u>	Apr 06, 2010
<u>AB1</u>		<u>200MG</u>	<u>A078866</u> <u>003</u>	Apr 06, 2010
<u>AB1</u>	WATSON LABS FLORIDA	<u>100MG</u>	<u>A079095</u> <u>001</u>	Mar 24, 2009
<u>AB1</u>		<u>150MG</u>	<u>A079095</u> <u>002</u>	Mar 24, 2009
<u>AB1</u>		<u>200MG</u>	<u>A079095</u> <u>003</u>	Mar 24, 2009
<u>WELLBUTRIN SR</u>				
<u>AB1</u>	GLAXOSMITHKLINE	<u>100MG</u>	<u>N020358</u> <u>002</u>	Oct 04, 1996
<u>AB1</u> +		<u>150MG</u>	<u>N020358</u> <u>003</u>	Oct 04, 1996
<u>AB1</u>		<u>200MG</u>	<u>N020358</u> <u>004</u>	Jun 14, 2002

BUPROPION HYDROCHLORIDE

<u>AB2</u>	ACTAVIS	<u>150MG</u>	<u>A077475</u> <u>001</u>	Mar 12, 2008
<u>AB2</u>	ANCHEN PHARMS	<u>150MG</u>	<u>A091520</u> <u>001</u>	Jun 09, 2011
<u>AB2</u>	IMPAK LABS	<u>150MG</u>	<u>A075914</u> <u>001</u>	May 27, 2004
<u>AB2</u>	MYLAN	<u>150MG</u>	<u>A090941</u> <u>001</u>	May 03, 2010
<u>AB2</u>	WATSON LABS FLORIDA	<u>150MG</u>	<u>A079094</u> <u>001</u>	Mar 24, 2009

ZYBAN

<u>AB2</u> +	GLAXOSMITHKLINE	<u>150MG</u>	<u>N020711</u> <u>003</u>	May 14, 1997
--------------	-----------------	--------------	---------------------------	--------------

BUPROPION HYDROCHLORIDE

<u>AB3</u>	ACTAVIS	<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>	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>	IMPAK LABS	<u>150MG</u>	<u>A077415</u> <u>001</u>	Nov 26, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 63 (of 424)

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

<u>AB3</u>	IMPAX LABS	<u>300MG</u>	<u>A077415</u> <u>002</u>	Dec 15, 2006
<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>	WATSON LABS	<u>150MG</u>	<u>A077715</u> <u>001</u>	Nov 26, 2008
<u>AB3</u>		<u>300MG</u>	<u>A077715</u> <u>002</u>	Jun 13, 2007
		<u>WELLBUTRIN XL</u>		
<u>AB3</u> +	VALEANT INTL	<u>150MG</u>	<u>N021515</u> <u>001</u>	Aug 28, 2003
<u>AB3</u>		<u>300MG</u>	<u>N021515</u> <u>002</u>	Aug 28, 2003
		<u>FORFIVO XL</u>		
+ INTELGENX CORP		450MG	N022497 001	Nov 10, 2011

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>5MG</u>	<u>A075521</u> <u>001</u>	Apr 05, 2002
<u>AB</u>		<u>10MG</u>	<u>A075521</u> <u>002</u>	Apr 05, 2002
<u>AB</u>		<u>15MG</u>	<u>A075521</u> <u>003</u>	Apr 05, 2002
<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>	EGIS	<u>5MG</u>	<u>A075119</u> <u>001</u>	Mar 14, 2002
<u>AB</u>		<u>10MG</u>	<u>A075119</u> <u>002</u>	Mar 14, 2002
<u>AB</u>		<u>15MG</u>	<u>A075119</u> <u>003</u>	Jan 23, 2003
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A075272</u> <u>001</u>	Mar 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075467</u> <u>001</u>	Feb 28, 2002
<u>AB</u>		<u>7.5MG</u>	<u>A075467</u> <u>002</u>	Mar 28, 2001
<u>AB</u>		<u>10MG</u>	<u>A075272</u> <u>002</u>	Mar 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075467</u> <u>003</u>	Feb 28, 2002
<u>AB</u>		<u>15MG</u>	<u>A075272</u> <u>003</u>	Mar 28, 2001
<u>AB</u>		<u>15MG</u>	<u>A075467</u> <u>004</u>	Feb 28, 2002
<u>AB</u>		<u>30MG</u>	<u>A076008</u> <u>001</u>	Jun 28, 2001
<u>AB</u>	NESHER PHARMS	<u>5MG</u>	<u>A075572</u> <u>001</u>	Feb 27, 2002
<u>AB</u>		<u>10MG</u>	<u>A075572</u> <u>002</u>	Feb 27, 2002
<u>AB</u>		<u>15MG</u>	<u>A075572</u> <u>003</u>	Feb 27, 2002
<u>AB</u>	TEVA	<u>5MG</u>	<u>A075022</u> <u>001</u>	Feb 28, 2002
<u>AB</u>		<u>10MG</u>	<u>A075022</u> <u>002</u>	Feb 28, 2002
<u>AB</u> +		<u>15MG</u>	<u>A075022</u> <u>003</u>	Feb 28, 2002
<u>AB</u>		<u>30MG</u>	<u>A075022</u> <u>004</u>	Mar 25, 2004
<u>AB</u>	WATSON LABS	<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

BUSULFAN

INJECTABLE; INJECTION

BUSULFEX

+ OTSUKA PHARM 6MG/ML N020954 001 Feb 04, 1999

TABLET; ORAL

MYLERAN

+ ASPEN GLOBAL 2MG N009386 001

BUTABARBITAL SODIUM

ELIXIR; ORAL

BUTISOL SODIUM

+ MEDA PHARMS 30MG/5ML A085380 001

PRESCRIPTION DRUG PRODUCT LIST

3 - 64 (of 424)

BUTABARBITAL SODIUM

TABLET; ORAL			
BUTISOL SODIUM			
+ MEDA PHARMS	30MG	N000793 004	
+	50MG	N000793 003	

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL			
MENTAX			
+ MYLAN	1%	N020524 001	Oct 18, 1996
MENTAX-TC			
+ MYLAN	1%	N021408 001	Oct 17, 2002

BUTOCONAZOLE NITRATE

CREAM; VAGINAL			
GYNAZOLE-1			
+ KV PHARM	2%	N019881 001	Feb 07, 1997

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

<u>AP</u>	BEDFORD	<u>2MG/ML</u>	<u>A075046 001</u>	Aug 12, 1998
<u>AP</u>	CLARIS LIFESCIENCES	<u>2MG/ML</u>	<u>A075697 001</u>	Oct 23, 2001
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A078400 001</u>	May 01, 2009
<u>AP</u>		<u>2MG/ML</u>	<u>A078247 001</u>	Apr 29, 2009
<u>AP</u>		<u>2MG/ML</u>	<u>A078400 002</u>	May 01, 2009

BUTORPHANOL TARTRATE PRESERVATIVE FREE

<u>AP</u>	BEDFORD	<u>1MG/ML</u>	<u>A075045 001</u>	Aug 12, 1998
<u>AP</u>		<u>2MG/ML</u>	<u>A075045 002</u>	Aug 12, 1998
<u>AP</u>	CLARIS LIFESCIENCES	<u>1MG/ML</u>	<u>A075695 001</u>	Oct 23, 2001
<u>AP</u>		<u>2MG/ML</u>	<u>A075695 002</u>	Oct 23, 2001
<u>AP</u>	HOSPIRA	<u>1MG/ML</u>	<u>A074626 001</u>	Jan 23, 1997
<u>AP</u>		<u>2MG/ML</u>	<u>A074626 002</u>	Jan 23, 1997

STADOL

<u>AP</u>	+ APOTHECON	<u>2MG/ML</u>	<u>N017857 004</u>	
<u>AP</u>	<u>STADOL PRESERVATIVE FREE</u>			

<u>AP</u>	+ APOTHECON	<u>1MG/ML</u>	<u>N017857 001</u>	
<u>AP</u>	+	<u>2MG/ML</u>	<u>N017857 002</u>	

SPRAY, METERED; NASAL

BUTORPHANOL TARTRATE

<u>AB</u>	+ MYLAN	<u>1MG/SPRAY</u>	<u>A075759 001</u>	Aug 08, 2001
<u>AB</u>	NOVEX	<u>1MG/SPRAY</u>	<u>A075499 001</u>	Dec 04, 2002
<u>AB</u>	ROXANE	<u>1MG/SPRAY</u>	<u>A075824 001</u>	Mar 12, 2002

CABAZITAXEL

SOLUTION; IV (INFUSION)

+ SANOFI AVENTIS US	60MG/1.5ML (40MG/ML)	N201023 001	Jun 17, 2010
---------------------	----------------------	-------------	--------------

CABERGOLINE

TABLET; ORAL

CABERGOLINE

<u>AB</u>	IMPAK LABS INC	<u>0.5MG</u>	<u>A077843 001</u>	Jul 03, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>0.5MG</u>	<u>A077750 001</u>	Mar 07, 2007
<u>AB</u>	+ PAR PHARM	<u>0.5MG</u>	<u>A076310 001</u>	Dec 29, 2005
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>A078035 001</u>	Apr 21, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 65 (of 424)

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAF'CIT

<u>AP</u> + MEAD JOHNSON	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N020793 001</u>	Sep 21, 1999
<u>CAFFEINE CITRATE</u>			
<u>AP</u> APP PHARMS	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077997 001</u>	Jul 20, 2007
<u>AP</u> LUITPOLD	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077906 001</u>	May 15, 2007
<u>AP</u> PADDOCK LLC	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077233 001</u>	Sep 21, 2006
<u>AP</u> SUN PHARMA GLOBAL	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A090077 001</u>	Sep 30, 2009

SOLUTION; ORAL

CAF'CIT

<u>AA</u> + MEAD JOHNSON	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>N020793 002</u>	Apr 12, 2000
<u>CAFFEINE CITRATE</u>			
<u>AA</u> APP PHARMS	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A078002 001</u>	Jan 31, 2008
<u>AA</u> LUITPOLD	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A090064 001</u>	Nov 20, 2009
<u>AA</u> PADDOCK LLC	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A077304 001</u>	Sep 21, 2006
<u>AA</u> SUN PHARMA GLOBAL	<u>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</u>	<u>A090357 001</u>	Sep 30, 2009

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL

MIGERGOT

+ G AND W LABS	100MG;2MG	<u>A086557 001</u>	Oct 04, 1983
----------------	-----------	--------------------	--------------

TABLET; ORAL

CAF'ERGOT

<u>AA</u> + SANDOZ	<u>100MG;1MG</u>	<u>A084294 001</u>	
<u>ERGOTAMINE TARTRATE AND CAFFEINE</u>			
<u>AA</u> MIKART	<u>100MG;1MG</u>	<u>A040590 001</u>	Sep 16, 2005
<u>AA</u> WEST WARD	<u>100MG;1MG</u>	<u>A040510 001</u>	Sep 17, 2004

CALCIPIOTRIENE

AEROSOL, FOAM; TOPICAL

SORILUX

+ STIEFEL LABS INC	0.005%	<u>N022563 001</u>	Oct 06, 2010
--------------------	--------	--------------------	--------------

CREAM; TOPICAL

DOVONEX

+ LEO PHARM	0.005%	<u>N020554 001</u>	Jul 22, 1996
-------------	--------	--------------------	--------------

OINTMENT; TOPICAL

CALCIPIOTRIENE

+ GLENMARK GENERICS	0.005%	<u>A090633 001</u>	Mar 24, 2010
---------------------	--------	--------------------	--------------

SOLUTION; TOPICAL

CALCIPIOTRIENE

<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> NYCOMED US	<u>0.005%</u>	<u>A078305 001</u>	May 06, 2008
<u>AT</u> TOLMAR	<u>0.005%</u>	<u>A077029 001</u>	Nov 20, 2009
<u>DOVONEX</u>			
<u>AT</u> + LEO PHARM	<u>0.005%</u>	<u>N020611 001</u>	Mar 03, 1997

CALCITONIN SALMON

INJECTABLE; INJECTION

MIACALCIN

+ NOVARTIS	200 IU/ML	<u>N017808 002</u>	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
<u>MIACALCIN</u>			
<u>AB</u> + NOVARTIS	<u>200 IU/SPRAY</u>	<u>N020313 002</u>	Aug 17, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 66 (of 424)

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED; NASAL
 FORTICAL
 + UPSHER SMITH 200 IU/SPRAY N021406 001 Aug 12, 2005

CALCITRIOL

CAPSULE; ORAL

CALCITRIOL

<u>AB</u>	ROXANE	<u>0.25MCG</u>	<u>A076917</u> <u>001</u>	Mar 27, 2006
<u>AB</u>	TEVA	<u>0.25MCG</u>	<u>A075765</u> <u>001</u>	Oct 12, 2001
<u>AB</u>		<u>0.5MCG</u>	<u>A075765</u> <u>002</u>	Oct 12, 2001

ROCALTROL

<u>AB</u>	VALIDUS PHARMS	<u>0.25MCG</u>	<u>N018044</u> <u>001</u>
<u>AB</u>	+	<u>0.5MCG</u>	<u>N018044</u> <u>002</u>

INJECTABLE; INJECTION

CALCIJEX

<u>AP</u>	+	ABBOTT	<u>0.001MG/ML</u>	<u>N018874</u> <u>001</u>	Sep 25, 1986
<u>AP</u>	+		<u>0.002MG/ML</u>	<u>N018874</u> <u>002</u>	Sep 25, 1986

CALCITRIOL

<u>AP</u>	AKORN	<u>0.001MG/ML</u>	<u>A078066</u> <u>001</u>	Jan 29, 2008
<u>AP</u>		<u>0.002MG/ML</u>	<u>A078066</u> <u>002</u>	Jan 29, 2008
<u>AP</u>	APP PHARMS	<u>0.001MG/ML</u>	<u>A075836</u> <u>001</u>	Dec 31, 2002
<u>AP</u>		<u>0.002MG/ML</u>	<u>A075836</u> <u>002</u>	Dec 31, 2002
<u>AP</u>	FRESENIUS MEDCL	<u>0.001MG/ML</u>	<u>A075766</u> <u>001</u>	Feb 20, 2003
<u>AP</u>		<u>0.002MG/ML</u>	<u>A075766</u> <u>002</u>	Feb 20, 2003
<u>AP</u>	LUITPOLD	<u>0.001MG/ML</u>	<u>A075746</u> <u>001</u>	Sep 26, 2003
<u>AP</u>		<u>0.002MG/ML</u>	<u>A075746</u> <u>002</u>	Sep 26, 2003
<u>AP</u>	ROCKWELL MEDCL	<u>0.001MG/ML</u>	<u>A076206</u> <u>001</u>	Sep 17, 2003
<u>AP</u>	SAGENT PHARMS	<u>0.001MG/ML</u>	<u>A077102</u> <u>001</u>	Feb 08, 2006
<u>AP</u>	TEVA PARENTERAL	<u>0.002MG/ML</u>	<u>A075823</u> <u>002</u>	Mar 31, 2003

OINTMENT; TOPICAL

VECTICAL

+ GALDERMA LABS LP 3MCG/GM N022087 001 Jan 23, 2009

SOLUTION; ORAL

CALCITRIOL

<u>AA</u>	ROXANE	<u>1MCG/ML</u>	<u>A076242</u> <u>001</u>	Jul 18, 2003
<u>AA</u>	+	<u>VALIDUS PHARMS</u>	<u>1MCG/ML</u>	<u>N021068</u> <u>001</u>

CALCIUM ACETATE

CAPSULE; ORAL

CALCIUM ACETATE

<u>AB</u>	ROXANE	<u>EQ 169MG CALCIUM</u>	<u>A077728</u> <u>001</u>	Feb 26, 2008
<u>AB</u>	+	<u>FRESENIUS MEDCL</u>	<u>EQ 169MG CALCIUM</u>	<u>N021160</u> <u>003</u>

SOLUTION; ORAL

PHOSLYRA

+ FRESENIUS MEDCL EQ 169MG CALCIUM/5ML N022581 001 Apr 18, 2011

TABLET; ORAL

CALCIUM ACETATE

<u>AB</u>	PADDICK LLC	<u>EQ 169MG CALCIUM</u>	<u>A091561</u> <u>001</u>	Apr 13, 2011
<u>AB</u>	+	<u>ELIPHOS</u>	<u>EQ 169MG CALCIUM</u>	<u>A078502</u> <u>001</u>

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

+ HOSPIRA 100MG/ML N021117 001 Jan 28, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 67 (of 424)

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

BSS PLUS

<u>AT</u>	<u>+</u>	<u>ALCON</u>	<u>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</u>	<u>N018469 001</u>	
<u>AT</u>	<u>+</u>	<u>AKORN</u>	<u>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</u>	<u>N020079 001</u>	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

+ GAMBRO RENAL PRODS	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.0 5GM/1000ML; 0.157GM/1000ML; 2.21GM/1000ML ; 7.07GM/1000ML	N021703 010	Oct 10, 2008
----------------------	--	-------------	--------------

PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.0 5GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML ; 7.07GM/1000ML	N021703 011	Oct 10, 2008
----------------------	--	-------------	--------------

PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 2.21GM/1000ML ; 7.07GM/1000ML	N021703 013	Oct 10, 2008
----------------------	--	-------------	--------------

PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46GM/1000ML	N021703 006	Oct 25, 2006
----------------------	---	-------------	--------------

PRISMASOL BGK 2/0 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.0 3GM/1000ML; 0.157GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML	N021703 002	Oct 25, 2006
----------------------	--	-------------	--------------

PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	5.15GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.03GM/1000ML; 0.157GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML	N021703 003	Oct 25, 2006
----------------------	--	-------------	--------------

PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	N/A/1000ML; 20GM/1000ML; 5.4GM/1000ML; 2.4 4GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML	N021703 015	Oct 10, 2008
----------------------	--	-------------	--------------

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	3.68GM/1000ML; 20GM/1000ML; 5.4GM/1000ML; 3.05GM/1000ML; 0.314GM/1000ML; 3.09GM/1000ML ; 6.46GM/1000ML	N021703 004	Oct 25, 2006
----------------------	--	-------------	--------------

PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	N/A/1000ML; N/A/1000ML; 5.4GM/1000ML; 2.44 GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6.46 GM/1000ML	N021703 014	Oct 10, 2008
----------------------	---	-------------	--------------

PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER

+ GAMBRO RENAL PRODS	5.15GM/1000ML; N/A/1000ML; 5.4GM/1000ML; 2 .03GM/1000ML; N/A/1000ML; 3.09GM/1000ML; 6 .46GM/1000ML	N021703 001	Oct 25, 2006
----------------------	--	-------------	--------------

CALCIUM 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, 2008
--------------------	--	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 68 (of 424)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER

+ B BRAUN 37MG/100ML;5GM/100ML;31MG/100ML;120MG/1 N019864 001 Jun 10, 1993
00ML;330MG/100ML;88MG/100ML

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER

+ B BRAUN 35MG/100ML;5GM/100ML;30MG/100ML;74MG/10 N019867 001 Dec 20, 1993
0ML;640MG/100ML;500MG/100ML;74MG/100ML

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/1 N017390 001
00ML;161MG/100ML;94MG/100ML;138MG/100ML

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;9.4G N018807 003 Aug 26, 1983
M/100ML;11GM/100ML

+ 510MG/100ML;30GM/100ML;200MG/100ML;9.2G N018807 001 Aug 26, 1983
M/100ML;9.6GM/100ML

DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER

+ B BRAUN 510MG/100ML;50GM/100ML;200MG/100ML;9.2G N018807 002 Aug 26, 1983
M/100ML;9.6GM/100ML

+ 510MG/100ML;50GM/100ML;200MG/100ML;9.4G N018807 004 Aug 26, 1983
M/100ML;11GM/100ML

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 N018379 002
67MG/100ML;392MG/100ML

AT FRESENIUS MEDCL 25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5 N018883 001
67MG/100ML;392MG/100ML

DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5 N018883 004
38MG/100ML;448MG/100ML

DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5 N020171 001
38MG/100ML;448MG/100ML

DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5 N018379 003
67MG/100ML;392MG/100ML

AT FRESENIUS MEDCL 25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5 N018883 002
67MG/100ML;392MG/100ML

DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5 N018883 005
38MG/100ML;448MG/100ML

DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5 N020171 002
38MG/100ML;448MG/100ML

DELFLEX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5 N018379 007
67MG/100ML;392MG/100ML

PRESCRIPTION DRUG PRODUCT LIST

3 - 69 (of 424)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N018379 001

AT FRESENIUS MEDCL 25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N018883 003 Nov 30, 1984

DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N018883 006 Nov 30, 1984

DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N020171 003 Aug 19, 1992

DELFLEX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N018379 004 Jul 07, 1982

DELFLEX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N018379 005 Jul 07, 1982

DELFLEX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N018379 008 Jun 24, 1988

DELFLEX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT FRESENIUS MEDCL 25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N018379 006 Jul 07, 1982

DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 001

DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 003

DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 002

DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N020183 001 Dec 04, 1992

DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 007 Jul 09, 1984

DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 008 Jul 09, 1984

DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 010 Nov 18, 1985

DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;567MG/100ML;392MG/100ML N017512 009 Jul 09, 1984

DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N017512 004

AT BAXTER HLTHCARE 25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N020163 001 Dec 04, 1992

DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N017512 005

AT BAXTER HLTHCARE 25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N020163 002 Dec 04, 1992

DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N017512 011 Nov 18, 1985

DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML N017512 006

PRESCRIPTION DRUG PRODUCT LIST

3 - 70 (of 424)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;
538MG/100ML;448MG/100ML N020163 003 Dec 04, 1992

INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT FRESENIUS 18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5
38MG/100ML;448MG/100ML N020374 001 Jun 13, 1994

INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT FRESENIUS 18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5
38MG/100ML;448MG/100ML N020374 002 Jun 13, 1994

INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT FRESENIUS 18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;
538MG/100ML;448MG/100ML N020374 004 Jun 13, 1994

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

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

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

DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 18.3MG/100ML;2.5GM/100ML;5.08MG/100ML;5
38MG/100ML;448MG/100ML N020183 002 Dec 04, 1992

DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;5
38MG/100ML;448MG/100ML N020183 003 Dec 04, 1992

DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 18.3MG/100ML;4.25GM/100ML;5.08MG/100ML;
538MG/100ML;448MG/100ML N020183 004 Dec 04, 1992

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 N020374 003 Jun 13, 1994

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INTRATHECAL

ELLIOTTS B SOLUTION

+ QOL MEDCL 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 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

AP HOSPIRA 33MG/100ML;5GM/100ML;30MG/100ML;860MG/1
00ML N018254 001

DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER

AP B BRAUN 33MG/100ML;5GM/100ML;30MG/100ML;860MG/1
00ML N020000 001 Apr 17, 1992

AP BAXTER HLTHCARE 33MG/100ML;5GM/100ML;30MG/100ML;860MG/1
00ML N016695 001

PRESCRIPTION DRUG PRODUCT LIST

3 - 72 (of 424)

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

SOLUTION; IRRIGATION

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

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; N017438 001
640MG/100ML; 496MG/100ML; 89.6MG/100ML

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

SOLUTION; PERfusion, CARDIAC

CARDIOPLEGIC IN PLASTIC CONTAINER

AT BAXTER HLTHCARE 17.6MG/100ML; 325.3MG/100ML; 119.3MG/100M L; 643MG/100ML A075323 001 Apr 21, 2000

PLEGISOL IN PLASTIC CONTAINER

AT + HOSPIRA 17.6MG/100ML; 325.3MG/100ML; 119.3MG/100M L; 643MG/100ML N018608 001 Feb 26, 1982

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N020002 001	Apr 17, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N016693 001	
<u>AP</u>	HOSPIRA	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N018251 001	

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N018156 001	
<u>AT</u>	BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N018495 001	Feb 19, 1982
<u>AT</u>	HOSPIRA	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	N017635 001	

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N019632 001	Feb 29, 1988
<u>AP</u>	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N016682 001	
<u>AP</u>	HOSPIRA	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N017641 001	

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N018681 001	Dec 27, 1982
<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N018494 001	Feb 19, 1982
<u>AT</u>		<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N018921 001	Apr 03, 1984
<u>AT</u>		<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N019933 001	Aug 29, 1989
<u>AT</u>	HOSPIRA	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG /100ML</u>	N019416 001	Jan 17, 1986

CALFACTANT

SUSPENSION; INTRATRACHEAL

INFASURF PRESERVATIVE FREE

+ ONY 35MG/ML N020521 001 Jul 01, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 73 (of 424)

CANDESARTAN CILEXETIL

TABLET; ORAL ATACAND ASTRAZENECA	4MG 8MG 16MG + 32MG	N020838 001 N020838 002 N020838 003 N020838 004	Jun 04, 1998 Jun 04, 1998 Jun 04, 1998 Jun 04, 1998
--	------------------------------	--	--

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL ATACAND HCT ASTRAZENECA	16MG;12.5MG 32MG;12.5MG + 32MG;25MG	N021093 001 N021093 002 N021093 003	Sep 05, 2000 Sep 05, 2000 May 16, 2008
--	---	---	--

CAPECITABINE

TABLET; ORAL XELODA HOFFMANN LA ROCHE	150MG 500MG	N020896 001 N020896 002	Apr 30, 1998 Apr 30, 1998
---	----------------	----------------------------	------------------------------

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION CAPASTAT SULFATE + AKORN	EQ 1GM BASE/VIAL	N050095 001
--	------------------	-------------

CAPSAICIN

PATCH; TOPICAL QUTENZA + NEUROGESX	8%	N022395 001	Nov 16, 2009
--	----	-------------	--------------

CAPTOPRIL

TABLET; ORAL <u>CAPOTEN</u> AB PAR PHARM	<u>12.5MG</u> <u>25MG</u> <u>50MG</u> + <u>100MG</u>	N018343 005 N018343 002 N018343 001 N018343 003	Jan 17, 1985
<u>CAPTOPRIL</u> AB APOTEX	<u>12.5MG</u> <u>25MG</u> <u>50MG</u> <u>100MG</u>	A074737 001 A074737 002 A074737 003 A074737 004	Oct 28, 1998
AB MYLAN	<u>12.5MG</u> <u>25MG</u> <u>50MG</u> <u>100MG</u>	A074434 001 A074434 002 A074434 003 A074434 004	Feb 13, 1996
AB PRINSTON INC	<u>12.5MG</u> <u>25MG</u> <u>50MG</u> <u>100MG</u>	A074477 001 A074477 002 A074477 003 A074477 004	Feb 13, 1996
AB SANDOZ	<u>12.5MG</u> <u>25MG</u> <u>50MG</u> <u>100MG</u>	A074363 001 A074363 002 A074363 003 A074363 004	Nov 09, 1995
AB STASON	<u>12.5MG</u> <u>25MG</u> <u>50MG</u> <u>100MG</u>	A074677 004 A074677 002 A074677 001 A074677 003	May 30, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 74 (of 424)

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

<u>AB</u>	TEVA	<u>12.5MG</u>	<u>A074322</u> <u>001</u>	Feb 13, 1996
<u>AB</u>		<u>12.5MG</u>	<u>A074483</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>25MG</u>	<u>A074483</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>50MG</u>	<u>A074483</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>		<u>100MG</u>	<u>A074483</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>12.5MG</u>	<u>A074451</u> <u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074386</u> <u>002</u>	May 23, 1996
<u>AB</u>		<u>25MG</u>	<u>A074451</u> <u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074386</u> <u>003</u>	May 23, 1996
<u>AB</u>		<u>50MG</u>	<u>A074451</u> <u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>A074386</u> <u>004</u>	May 23, 1996
<u>AB</u>		<u>100MG</u>	<u>A074451</u> <u>004</u>	Feb 13, 1996
<u>AB</u>	WEST WARD	<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>	WOCKHARDT	<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

CAPPOZIDE 25/15

<u>AB</u>	APOTHECON	<u>25MG;15MG</u>	<u>N018709</u> <u>001</u>	Oct 12, 1984
<u>AB</u>	<u>CAPPOZIDE 25/25</u>	<u>25MG;25MG</u>	<u>N018709</u> <u>002</u>	Oct 12, 1984
<u>AB</u>	+ APOTHECON	<u>25MG;25MG</u>	<u>N018709</u> <u>002</u>	Oct 12, 1984
<u>AB</u>	<u>CAPPOZIDE 50/15</u>	<u>50MG;15MG</u>	<u>N018709</u> <u>004</u>	Oct 12, 1984
<u>AB</u>	+ APOTHECON	<u>50MG;15MG</u>	<u>N018709</u> <u>004</u>	Oct 12, 1984
<u>AB</u>	<u>CAPPOZIDE 50/25</u>	<u>50MG;25MG</u>	<u>N018709</u> <u>003</u>	Oct 12, 1984
<u>AB</u>	APOTHECON	<u>50MG;25MG</u>	<u>N018709</u> <u>003</u>	Oct 12, 1984
	<u>CAPTOPRIL AND HYDROCHLOROTHIAZIDE</u>			
<u>AB</u>	MYLAN	<u>25MG;15MG</u>	<u>A074896</u> <u>001</u>	Dec 29, 1997
<u>AB</u>		<u>25MG;25MG</u>	<u>A074896</u> <u>002</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;15MG</u>	<u>A074896</u> <u>004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>A074896</u> <u>003</u>	Dec 29, 1997
<u>AB</u>	TEVA	<u>25MG;15MG</u>	<u>A074827</u> <u>001</u>	Dec 29, 1997
<u>AB</u>		<u>25MG;25MG</u>	<u>A074827</u> <u>002</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;15MG</u>	<u>A074827</u> <u>004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>A074827</u> <u>003</u>	Dec 29, 1997

CARBACHOL

SOLUTION; INTRAOCULAR

MIOSTAT

+ ALCON

0.01%

N016968 001

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A078986</u> <u>001</u>	Nov 25, 2011
<u>AB</u>		<u>200MG</u>	<u>A078986</u> <u>002</u>	Nov 25, 2011
<u>AB</u>		<u>300MG</u>	<u>A078986</u> <u>003</u>	Nov 25, 2011
<u>AB</u>	NOSTRUM	<u>100MG</u>	<u>A076697</u> <u>001</u>	May 20, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 75 (of 424)

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

<u>AB</u>	NOSTRUM	<u>200MG</u>	<u>A076697</u> <u>002</u>	May 20, 2011
<u>AB</u>		<u>300MG</u>	<u>A076697</u> <u>003</u>	May 20, 2011
	<u>CARBATROL</u>			
<u>AB</u>	SHIRE	<u>100MG</u>	<u>N020712</u> <u>003</u>	Sep 30, 1997
<u>AB</u>		<u>200MG</u>	<u>N020712</u> <u>001</u>	Sep 30, 1997
<u>AB</u> +		<u>300MG</u>	<u>N020712</u> <u>002</u>	Sep 30, 1997
	EQUETRO			
	VALIDUS PHARMS INC	100MG	N021710 001	Dec 10, 2004
		200MG	N021710 002	Dec 10, 2004
	+	300MG	N021710 003	Dec 10, 2004
	SUSPENSION; ORAL			
	<u>CARBAMAZEPINE</u>			
<u>AB</u>	WOCKHARDT	<u>100MG/5ML</u>	<u>A075714</u> <u>001</u>	Jun 05, 2002
	<u>TEGRETOL</u>			
<u>AB</u> +	NOVARTIS	<u>100MG/5ML</u>	<u>N018927</u> <u>001</u>	Dec 18, 1987
	<u>TERIL</u>			
<u>AB</u>	TARO	<u>100MG/5ML</u>	<u>A076729</u> <u>001</u>	Sep 20, 2004
	TABLET; ORAL			
	<u>CARBAMAZEPINE</u>			
<u>AB</u>	APOTEX INC	<u>200MG</u>	<u>A075948</u> <u>001</u>	Feb 27, 2002
<u>AB</u>	TARO	<u>200MG</u>	<u>A074649</u> <u>001</u>	Oct 03, 1996
<u>AB</u>	TORRENT PHARMS	<u>200MG</u>	<u>A077272</u> <u>002</u>	Dec 07, 2005
	<u>EPITOL</u>			
<u>AB</u>	TEVA	<u>200MG</u>	<u>A070541</u> <u>001</u>	Sep 17, 1986
	<u>TEGRETOL</u>			
<u>AB</u> +	NOVARTIS	<u>200MG</u>	<u>N016608</u> <u>001</u>	
	CARBAMAZEPINE			
	TORRENT PHARMS	100MG	A077272 001	Dec 07, 2005
		300MG	A077272 003	Dec 07, 2005
		400MG	A077272 004	Dec 07, 2005
	TABLET, CHEWABLE; ORAL			
	<u>CARBAMAZEPINE</u>			
<u>AB</u>	TARO PHARM INDs	<u>100MG</u>	<u>A075687</u> <u>001</u>	Oct 24, 2000
<u>AB</u>	TORRENT PHARMS	<u>100MG</u>	<u>A075712</u> <u>001</u>	Jul 05, 2001
	<u>EPITOL</u>			
<u>AB</u>	TEVA	<u>100MG</u>	<u>A073524</u> <u>001</u>	Jul 29, 1992
	<u>TEGRETOL</u>			
<u>AB</u> +	NOVARTIS	<u>100MG</u>	<u>N018281</u> <u>001</u>	
	CARBAMAZEPINE			
	+ TARO PHARM INDs	200MG	A075687 002	Jul 29, 2002
	TABLET, EXTENDED RELEASE; ORAL			
	<u>CARBAMAZEPINE</u>			
<u>AB</u>	TARO	<u>100MG</u>	<u>A078115</u> <u>001</u>	Mar 31, 2009
<u>AB</u>		<u>200MG</u>	<u>A078115</u> <u>002</u>	Mar 31, 2009
<u>AB</u>		<u>400MG</u>	<u>A078115</u> <u>003</u>	Mar 31, 2009
	<u>TEGRETOL-XR</u>			
<u>AB</u>	NOVARTIS	<u>100MG</u>	<u>N020234</u> <u>001</u>	Mar 25, 1996
<u>AB</u>		<u>200MG</u>	<u>N020234</u> <u>002</u>	Mar 25, 1996
<u>AB</u> +		<u>400MG</u>	<u>N020234</u> <u>003</u>	Mar 25, 1996

CARBIDOPA

TABLET; ORAL

LODOSYN

+ ATON

25MG

N017830 001

PRESCRIPTION DRUG PRODUCT LIST

3 - 76 (of 424)

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL				
STALEVO 100				
ORION	25MG;200MG;100MG		N021485 002	Jun 11, 2003
STALEVO 125	31.25MG;200MG;125MG		N021485 006	Aug 29, 2008
STALEVO 150	37.5MG;200MG;150MG		N021485 003	Jun 11, 2003
STALEVO 200	50MG;200MG;200MG		N021485 004	Aug 02, 2007
+ ORION	12.5MG;200MG;50MG		N021485 001	Jun 11, 2003
STALEVO 75	18.75MG;200MG;75MG		N021485 005	Aug 29, 2008
ORION				

CARBIDOPA; LEVODOPA

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>	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
<u>AB</u>	TEVA	<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>SINEMET</u>		
<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>	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>	IMPAX 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
		<u>SINEMET CR</u>		
<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>	IMPAX LABS	<u>10MG;100MG</u>	<u>A090631 001</u>	Jun 08, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A090631 002</u>	Jun 08, 2010
<u>AB</u>		<u>25MG;250MG</u>	<u>A090631 003</u>	Jun 08, 2010
<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 PHARM INDs	<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

PRESCRIPTION DRUG PRODUCT LIST

3 - 77 (of 424)

CARBIDOPA; LEVODOPA

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	SUN PHARM INDS	<u>25MG;250MG</u>	<u>A078690</u>	<u>003</u>	Jul 31, 2009
	<u>PARCOPA</u>				
<u>AB</u>	SCHWARZ PHARMA	<u>10MG;100MG</u>	<u>A076699</u>	<u>001</u>	Aug 27, 2004
<u>AB</u>		<u>25MG;100MG</u>	<u>A076699</u>	<u>002</u>	Aug 27, 2004
<u>AB</u> +		<u>25MG;250MG</u>	<u>A076699</u>	<u>003</u>	Aug 27, 2004

CARBINOXAMINE MALEATE

SOLUTION; ORAL

CARBINOXAMINE MALEATE

<u>AA</u>	BOCA PHARMA	<u>4MG/5ML</u>	<u>A040814</u>	<u>001</u>	Feb 26, 2008
<u>AA</u>	CYPRESS PHARM	<u>4MG/5ML</u>	<u>A090418</u>	<u>001</u>	May 04, 2010
<u>AA</u> +	MIKART	<u>4MG/5ML</u>	<u>A040458</u>	<u>001</u>	Apr 25, 2003

TABLET; ORAL

CARBINOXAMINE MALEATE

<u>AA</u>	BOCA PHARMA	<u>4MG</u>	<u>A040639</u>	<u>002</u>	May 30, 2008
<u>AA</u>	CYPRESS PHARM	<u>4MG</u>	<u>A090417</u>	<u>001</u>	Aug 23, 2010
<u>AA</u>	INVAGEN PHARMS	<u>4MG</u>	<u>A090435</u>	<u>001</u>	Apr 15, 2010
<u>AA</u>	LYNROSE LABS	<u>4MG</u>	<u>A090756</u>	<u>001</u>	May 27, 2011
<u>AA</u> +	MIKART	<u>4MG</u>	<u>A040442</u>	<u>001</u>	Mar 19, 2003

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

<u>AP</u>	APP PHARMS	<u>50MG/VIAL</u>	<u>A076235</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>A076235</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>A076235</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	BEDFORD	<u>50MG/VIAL</u>	<u>A076099</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>A076099</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>A076099</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	PLIVA	<u>50MG/VIAL</u>	<u>A076602</u>	<u>001</u>	Nov 16, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>A076602</u>	<u>002</u>	Nov 16, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>A076602</u>	<u>003</u>	Nov 16, 2004
<u>AP</u>	SANDOZ	<u>50MG/VIAL</u>	<u>A076959</u>	<u>001</u>	Mar 18, 2005
<u>AP</u>		<u>150MG/VIAL</u>	<u>A076959</u>	<u>002</u>	Mar 18, 2005
<u>AP</u>		<u>450MG/VIAL</u>	<u>A076959</u>	<u>003</u>	Mar 18, 2005
<u>AP</u> +	WATSON LABS	<u>50MG/VIAL</u>	<u>A076162</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>A077383</u>	<u>001</u>	Jan 27, 2006
<u>AP</u> +		<u>150MG/VIAL</u>	<u>A076162</u>	<u>002</u>	Oct 14, 2004
<u>AP</u> +		<u>450MG/VIAL</u>	<u>A077383</u>	<u>002</u>	Jan 27, 2006
<u>AP</u> +		<u>450MG/VIAL</u>	<u>A076162</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>A077383</u>	<u>003</u>	Jan 27, 2006

INJECTABLE; IV (INFUSION)

CARBOPLATIN

<u>AP</u>	AKORN	<u>50MG/5ML (10MG/ML)</u>	<u>A090475</u>	<u>001</u>	Jul 29, 2009
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A090475</u>	<u>002</u>	Jul 29, 2009
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A090475</u>	<u>003</u>	Jul 29, 2009
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A091268</u>	<u>002</u>	Jul 28, 2010
<u>AP</u>	APP PHARMS	<u>50MG/5ML (10MG/ML)</u>	<u>A077266</u>	<u>001</u>	Feb 15, 2006
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077266</u>	<u>002</u>	Feb 15, 2006
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077247</u>	<u>003</u>	Oct 21, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077266</u>	<u>003</u>	Feb 15, 2006
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077266</u>	<u>004</u>	Feb 15, 2006
<u>AP</u>	BEDFORD LABS	<u>50MG/5ML (10MG/ML)</u>	<u>A077244</u>	<u>001</u>	Oct 15, 2004
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077244</u>	<u>002</u>	Oct 15, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077244</u>	<u>003</u>	Oct 15, 2004
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077244</u>	<u>004</u>	Jan 20, 2006
<u>AP</u>	BIONICHE PHARMA USA	<u>50MG/5ML (10MG/ML)</u>	<u>A077998</u>	<u>001</u>	Apr 24, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 78 (of 424)

CARBOPLATIN

INJECTABLE; IV (INFUSION)

CARBOPLATIN

<u>AP</u>	BIONICHE PHARMA USA	<u>150MG/15ML (10MG/ML)</u>	<u>A077998</u> <u>002</u>	Apr 24, 2007
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077998</u> <u>003</u>	Apr 24, 2007
<u>AP</u>	EBEWE PHARMA	<u>50MG/5ML (10MG/ML)</u>	<u>A078280</u> <u>001</u>	May 08, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A078280</u> <u>002</u>	May 08, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078280</u> <u>003</u>	May 08, 2008
<u>AP</u>	FRESENIUS KABI ONCOL	<u>450MG/45ML (10MG/ML)</u>	<u>A077432</u> <u>003</u>	Sep 29, 2006
<u>AP</u>		<u>50MG/5ML (10MG/ML)</u>	<u>A077432</u> <u>001</u>	Sep 29, 2006
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077432</u> <u>002</u>	Sep 29, 2006
<u>AP</u>	HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A076517</u> <u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A076517</u> <u>002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A076517</u> <u>003</u>	Oct 14, 2004
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077059</u> <u>001</u>	Nov 23, 2004
<u>AP</u>	ONCO THERAPIES LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A091063</u> <u>001</u>	Nov 09, 2011
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A091063</u> <u>002</u>	Nov 09, 2011
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A091063</u> <u>003</u>	Nov 09, 2011
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A091063</u> <u>004</u>	Nov 09, 2011
<u>AP</u>	+ PHARMACHEMIE	<u>50MG/5ML (10MG/ML)</u>	<u>A077269</u> <u>001</u>	Oct 14, 2004
<u>AP</u>	+	<u>150MG/15ML (10MG/ML)</u>	<u>A077269</u> <u>002</u>	Oct 14, 2004
<u>AP</u>	+	<u>450MG/45ML (10MG/ML)</u>	<u>A077269</u> <u>003</u>	Oct 14, 2004
<u>AP</u>	+	<u>600MG/60ML (10MG/ML)</u>	<u>A077269</u> <u>004</u>	Dec 28, 2007
<u>AP</u>	PHARMACHEMIE BV	<u>50MG/5ML (10MG/ML)</u>	<u>A077679</u> <u>001</u>	Feb 25, 2009
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077679</u> <u>002</u>	Feb 25, 2009
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077679</u> <u>003</u>	Feb 25, 2009
<u>AP</u>	PLIVA LACHEMA	<u>50MG/5ML (10MG/ML)</u>	<u>A078631</u> <u>001</u>	Dec 02, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A078631</u> <u>002</u>	Dec 02, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078631</u> <u>003</u>	Dec 02, 2008
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A078631</u> <u>004</u>	Dec 02, 2008
<u>AP</u>	SUN PHARMA GLOBAL	<u>50MG/5ML (10MG/ML)</u>	<u>A077926</u> <u>001</u>	Sep 19, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077926</u> <u>002</u>	Sep 19, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077926</u> <u>003</u>	Sep 19, 2008
<u>AP</u>	+ TEVA PARENTERAL	<u>50MG/5ML (10MG/ML)</u>	<u>A077139</u> <u>001</u>	Sep 21, 2005
<u>AP</u>	+	<u>150MG/15ML (10MG/ML)</u>	<u>A077139</u> <u>002</u>	Sep 21, 2005
<u>AP</u>	+	<u>450MG/45ML (10MG/ML)</u>	<u>A077139</u> <u>003</u>	Sep 21, 2005
<u>AP</u>	+	<u>600MG/60ML (10MG/ML)</u>	<u>A077139</u> <u>004</u>	Sep 21, 2005
<u>AP</u>	WATSON LABS	<u>50MG/5ML (10MG/ML)</u>	<u>A077861</u> <u>001</u>	Jan 18, 2007
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077861</u> <u>002</u>	Jan 18, 2007
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077861</u> <u>003</u>	Jan 18, 2007
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077861</u> <u>004</u>	Jan 18, 2007
	CARBOPLATIN			
	+ ONCO THERAPIES LTD	<u>1GM/100ML (10MG/ML)</u>	<u>A091478</u> <u>001</u>	Nov 23, 2011

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

HEMABATE

+ PHARMACIA AND UPJOHN EQ 0.25MG BASE/ML

N017989 001

CARGLUMIC ACID

TABLET; ORAL

CARBAGLU

+ ORPHAN EUROPE 200MG

N022562 001 Mar 18, 2010

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

<u>AA</u>	ADVENT PHARMS	<u>350MG</u>	<u>A040576</u> <u>001</u>	Jun 07, 2005
<u>AA</u>	AUROBINDO PHARMA	<u>350MG</u>	<u>A040792</u> <u>001</u>	Aug 06, 2009
<u>AA</u>	COREPHARMA	<u>350MG</u>	<u>A040397</u> <u>001</u>	Sep 21, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 79 (of 424)

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

<u>AA</u>	MIRROR PHARMS	<u>350MG</u>	<u>A040823</u>	<u>001</u>	Oct 22, 2008
<u>AA</u>	MUTUAL PHARM	<u>350MG</u>	<u>A089346</u>	<u>001</u>	Oct 17, 1991
<u>AA</u>	PROSAM LABS	<u>350MG</u>	<u>A040188</u>	<u>001</u>	Mar 07, 1997
<u>AA</u>	SUN PHARM INDs LTD	<u>350MG</u>	<u>A040755</u>	<u>001</u>	Feb 27, 2007
<u>AA</u>	VINTAGE PHARMS	<u>350MG</u>	<u>A040245</u>	<u>001</u>	Sep 08, 1997
<u>AA</u>	WATSON LABS	<u>350MG</u>	<u>A087499</u>	<u>001</u>	Apr 20, 1982
<u>AA</u>	WEST WARD	<u>350MG</u>	<u>A040124</u>	<u>001</u>	Jan 24, 1996
	<u>SOMA</u>				
<u>AA</u>	MEDA PHARMS	<u>350MG</u>	<u>N011792</u>	<u>001</u>	
	SOMA				
+ MEDA PHARMS		250MG	N011792	004	Sep 13, 2007

CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

+ EISAI INC	7.7MG	N020637	001	Sep 23, 1996
-------------	-------	---------	-----	--------------

INJECTABLE; INJECTION

BICNU

+ BRISTOL	100MG/VIAL	N017422	001	
-----------	------------	---------	-----	--

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

<u>AT</u>	ALCON	<u>1%</u>	<u>A075476</u>	<u>001</u>	Jan 03, 2000
<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>A075546</u>	<u>001</u>	Jan 20, 2000
<u>AT</u>	NOVEX	<u>1%</u>	<u>A076097</u>	<u>001</u>	Feb 06, 2002
	<u>OCUPRESS</u>				
<u>AT</u>	+ NOVARTIS	<u>1%</u>	<u>N019972</u>	<u>001</u>	May 23, 1990

CARVEDILOL

TABLET; ORAL

CARVEDILOL

<u>AB</u>	APOTEX INC	<u>3.125MG</u>	<u>A078165</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078165</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078165</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078165</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>3.125MG</u>	<u>A078332</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078332</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078332</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078332</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	BEXIMCO USA	<u>3.125MG</u>	<u>A078384</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078384</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078384</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078384</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	CARACO	<u>3.125MG</u>	<u>A077346</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077346</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077346</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077346</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>3.125MG</u>	<u>A076649</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076649</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076649</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076649</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	GLENMARK GENERICS	<u>3.125MG</u>	<u>A078251</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078251</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078251</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078251</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	HIKMA	<u>3.125MG</u>	<u>A077887</u>	<u>001</u>	Sep 07, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 80 (of 424)

CARVEDILOL

TABLET; ORAL

CARVEDILOL

<u>AB</u>	HIKMA	<u>6.25MG</u>	<u>A077887</u>	<u>002</u>	Sep 07, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077887</u>	<u>003</u>	Sep 07, 2007
<u>AB</u>		<u>25MG</u>	<u>A077887</u>	<u>004</u>	Sep 07, 2007
<u>AB</u>	LUPIN	<u>3.125MG</u>	<u>A078217</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078217</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078217</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078217</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	MYLAN	<u>3.125MG</u>	<u>A077316</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077316</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077316</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077316</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	PLIVA HRVATSKA DOO	<u>3.125MG</u>	<u>A078240</u>	<u>001</u>	Oct 30, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078240</u>	<u>002</u>	Oct 30, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078240</u>	<u>003</u>	Oct 30, 2007
<u>AB</u>		<u>25MG</u>	<u>A078240</u>	<u>004</u>	Oct 30, 2007
<u>AB</u>	RANBAXY	<u>3.125MG</u>	<u>A076989</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076989</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076989</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076989</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	SANDOZ	<u>3.125MG</u>	<u>A078227</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078227</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078227</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078227</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	TARO	<u>3.125MG</u>	<u>A077780</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077780</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077780</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077780</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	TEVA	<u>3.125MG</u>	<u>A076373</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076373</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076373</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076373</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	WATSON LABS	<u>3.125MG</u>	<u>A077474</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077474</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077474</u>	<u>003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077474</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>	WOCKHARDT	<u>3.125MG</u>	<u>A078786</u>	<u>001</u>	Dec 22, 2009
<u>AB</u>		<u>6.25MG</u>	<u>A078786</u>	<u>002</u>	Dec 22, 2009
<u>AB</u>		<u>12.5MG</u>	<u>A078786</u>	<u>003</u>	Dec 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A078786</u>	<u>004</u>	Dec 22, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>3.125MG</u>	<u>A077614</u>	<u>004</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077614</u>	<u>001</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077614</u>	<u>002</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077614</u>	<u>003</u>	Sep 05, 2007
<u> </u>	<u>COREG</u>				
<u>AB</u>	SMITHKLINE BEECHAM	<u>3.125MG</u>	<u>N020297</u>	<u>004</u>	May 29, 1997
<u>AB</u>		<u>6.25MG</u>	<u>N020297</u>	<u>003</u>	Sep 14, 1995
<u>AB</u>	+	<u>12.5MG</u>	<u>N020297</u>	<u>002</u>	Sep 14, 1995
<u>AB</u>		<u>25MG</u>	<u>N020297</u>	<u>001</u>	Sep 14, 1995

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

COREG CR

SB PHARMCO

10MG

N022012 001 Oct 20, 2006

20MG

N022012 002 Oct 20, 2006

+

40MG

N022012 003 Oct 20, 2006

80MG

N022012 004 Oct 20, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 81 (of 424)

CASPOFUNGIN ACETATE

INJECTABLE; IV (INFUSION)

+ MERCK	50MG/VIAL	N021227 001	Jan 26, 2001
+	70MG/VIAL	N021227 002	Jan 26, 2001

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>	RANBAXY	<u>EQ 250MG BASE</u>	<u>A064156 001</u>	Aug 28, 1997
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A064156 002</u>	Aug 28, 1997
<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

CEFACLOR

+ 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

TABLET, CHEWABLE; ORAL

RANICLOR

RANBAXY	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

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>	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	
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 500MG BASE</u>	<u>A065309 001</u>	Sep 18, 2006	
<u>AB</u>	RANBAXY	<u>EQ 500MG BASE</u>	<u>A065015 001</u>	Jun 22, 1999	
<u>AB</u>	SANDOZ	<u>EQ 500MG BASE</u>	<u>A062291 001</u>		
<u>AB</u>	+	<u>TEVA PHARMS</u>	<u>EQ 500MG BASE</u>	<u>A065282 001</u>	Jan 20, 2006

FOR SUSPENSION; ORAL

CEFADROXIL

<u>AB</u>	LUPIN	<u>EQ 250MG BASE/5ML</u>	<u>A065396 001</u>	Feb 21, 2008
<u>AB</u>	+	<u>EQ 500MG BASE/5ML</u>	<u>A065396 002</u>	Feb 21, 2008
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE/5ML</u>	<u>A065307 002</u>	Oct 16, 2006
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A065307 003</u>	Oct 16, 2006
<u>AB</u>	RANBAXY	<u>EQ 250MG BASE/5ML</u>	<u>A065115 002</u>	Mar 26, 2003
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A065115 003</u>	Mar 26, 2003
<u>AB</u>	TEVA PHARMS	<u>EQ 250MG BASE/5ML</u>	<u>A065278 001</u>	Jan 20, 2006
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>A065278 002</u>	Jan 20, 2006

CEFADROXIL

RANBAXY

EQ 125MG BASE/5ML

A065115 001 Mar 26, 2003

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
<u>AB</u>	RANBAXY	<u>EQ 1GM BASE</u>	<u>A065018 001</u>	Apr 23, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 82 (of 424)

CEFADROXIL/CEFADROXIL HEMIHYDRATE

TABLET; ORAL
 CEFADROXIL
 + IVAX SUB 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</u> <u>001</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065303</u> <u>002</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065306</u> <u>001</u>	Oct 22, 2008
<u>AP</u>	+ APP PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A064169</u> <u>001</u>	Aug 14, 1998
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>A064169</u> <u>002</u>	Aug 14, 1998
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>A064170</u> <u>001</u>	Mar 18, 1998
<u>AP</u>	+	<u>EQ 20GM BASE/VIAL</u>	<u>A064170</u> <u>002</u>	Mar 18, 1998
<u>AP</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A065395</u> <u>001</u>	Aug 08, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065395</u> <u>002</u>	Aug 08, 2008
<u>AP</u>	CEPHAZONE PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A065280</u> <u>001</u>	Mar 18, 2009
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065280</u> <u>002</u>	Mar 18, 2009
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065295</u> <u>001</u>	Mar 18, 2009
<u>AP</u>		<u>EQ 20GM BASE/VIAL</u>	<u>A065296</u> <u>001</u>	Mar 18, 2009
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 500MG BASE/VIAL</u>	<u>A065047</u> <u>001</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065047</u> <u>002</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065143</u> <u>001</u>	Oct 18, 2004
<u>AP</u>	HOSPIRA INC	<u>EQ 500MG BASE/VIAL</u>	<u>A065226</u> <u>001</u>	Apr 21, 2005
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065226</u> <u>002</u>	Apr 21, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065244</u> <u>001</u>	Aug 12, 2005
<u>AP</u>		<u>EQ 20GM BASE/VIAL</u>	<u>A065247</u> <u>001</u>	Aug 12, 2005
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A062831</u> <u>001</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062831</u> <u>002</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065345</u> <u>001</u>	May 09, 2007
<u>AP</u>		<u>EQ 20GM BASE/VIAL</u>	<u>A062831</u> <u>003</u>	Sep 25, 1992
<u>AP</u>	STERI PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A063214</u> <u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A063216</u> <u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A063207</u> <u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 20GM BASE/VIAL</u>	<u>A063208</u> <u>001</u>	Dec 27, 1991
<u>AP</u>			<u>A063209</u> <u>001</u>	Dec 27, 1991
<u>AP</u>			<u>A063209</u> <u>002</u>	Apr 30, 1999
	<u>KEFZOL</u>			
<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A061773</u> <u>002</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A061773</u> <u>003</u>	
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A061773</u> <u>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
	CEFAZOLIN SODIUM			
+ SAMSON MEDCL		EQ 100GM BASE/VIAL	A065141 001	Nov 29, 2006
+		EQ 300GM BASE/VIAL	A065141 002	Nov 29, 2006

CEFDINIR

CAPSULE; ORAL

CEFDINIR

<u>AB</u>	AUROBINDO PHARMA	<u>300MG</u>	<u>A065434</u> <u>001</u>	Jan 07, 2008
<u>AB</u>	LUPIN	<u>300MG</u>	<u>A065264</u> <u>001</u>	May 19, 2006
<u>AB</u>	ORCHID HLTHCARE	<u>300MG</u>	<u>A065418</u> <u>001</u>	Jul 18, 2007
<u>AB</u>	+ SANDOZ	<u>300MG</u>	<u>A065330</u> <u>001</u>	Apr 06, 2007
<u>AB</u>	TEVA PHARMS	<u>300MG</u>	<u>A065368</u> <u>001</u>	May 09, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 83 (of 424)

CEFDINIR

FOR SUSPENSION; ORAL

CEFDINIR

<u>AB</u>	AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065473</u>	<u>001</u>	Dec 14, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065473</u>	<u>002</u>	Dec 14, 2007
<u>AB</u>	LUPIN	<u>125MG/5ML</u>	<u>A065259</u>	<u>001</u>	May 31, 2006
<u>AB</u>		<u>250MG/5ML</u>	<u>A065259</u>	<u>002</u>	May 07, 2007
<u>AB</u>	ORCHID HLTHCARE	<u>125MG/5ML</u>	<u>A065429</u>	<u>001</u>	Jul 18, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065429</u>	<u>002</u>	Jul 18, 2007
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065337</u>	<u>001</u>	Apr 06, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065337</u>	<u>002</u>	Apr 06, 2007
<u>AB</u>	+ TEVA PHARMS	<u>125MG/5ML</u>	<u>A065332</u>	<u>001</u>	May 04, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065332</u>	<u>002</u>	May 04, 2007

CEFDITOREN PIVOXIL

TABLET; ORAL

SPECTRACEF

CORNERSTONE THERAP	200MG	N021222	001	Aug 29, 2001
+	400MG	N021222	002	Jul 21, 2008

CEFEPIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

<u>AP</u>	ACS DOBFAR	<u>EQ 1GM BASE/VIAL</u>	<u>A065441</u>	<u>001</u>	Mar 20, 2008
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065441</u>	<u>002</u>	Mar 20, 2008
<u>AP</u>	HOSPIRA INC	<u>EQ 500MG BASE/VIAL</u>	<u>A065369</u>	<u>001</u>	Jun 18, 2007
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065369</u>	<u>002</u>	Jun 18, 2007
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065369</u>	<u>003</u>	Jun 18, 2007
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A090291</u>	<u>001</u>	Dec 21, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090291</u>	<u>002</u>	Dec 21, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090291</u>	<u>003</u>	Dec 21, 2010
	<u>MAXIPIME</u>				
<u>AP</u>	+ HOSPIRA INC	<u>EQ 500MG BASE/VIAL</u>	<u>N050679</u>	<u>001</u>	Jan 18, 1996
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N050679</u>	<u>002</u>	Jan 18, 1996
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>N050679</u>	<u>003</u>	Jan 18, 1996
	CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER				
	B BRAUN	EQ 1GM BASE/VIAL	N050821	001	May 06, 2010
		EQ 2GM BASE/VIAL	N050821	002	May 06, 2010
	CEFEPIME IN PLASTIC CONTAINER				
	+ BAXTER HLTHCARE	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)	N050817	001	Aug 05, 2008
	+	EQ 2GM BASE/100ML (EQ 20MG BASE/ML)	N050817	002	Aug 05, 2008

CEFIXIME

SUSPENSION; ORAL

SUPRAX

LUPIN PHARMS	100MG/5ML	A065129	001	Feb 23, 2004
+	200MG/5ML	A065355	001	Apr 10, 2007

TABLET; ORAL

SUPRAX

+ LUPIN PHARMS	400MG	A065130	001	Feb 12, 2004
----------------	-------	---------	-----	--------------

TABLET, CHEWABLE; ORAL

SUPRAX

LUPIN LTD	100MG	A065380	001	Oct 25, 2010
	150MG	A065380	002	Oct 25, 2010
+	200MG	A065380	003	Oct 25, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 84 (of 424)

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME

<u>AP</u>	APP PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A064200 001</u>	Mar 24, 2000
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A064200 002</u>	Mar 24, 2000
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A064200 003</u>	Mar 24, 2000
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A064201 001</u>	Mar 24, 2000
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>A065072 001</u>	Nov 20, 2002
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065072 002</u>	Nov 20, 2002
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065072 003</u>	Nov 20, 2002
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065071 001</u>	Nov 20, 2002
<u>AP</u>	WOCKHARDT	<u>EQ 1GM BASE/VIAL</u>	<u>A065197 001</u>	Aug 29, 2006
	<u>CEFOTAXIME SODIUM</u>			
<u>AP</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE/VIAL</u>	<u>A065517 001</u>	Nov 06, 2009
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065517 002</u>	Nov 06, 2009
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065517 003</u>	Nov 06, 2009
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065516 001</u>	Nov 06, 2009
<u>AP</u>	CEPHAZONE PHARMA	<u>EQ 10GM BASE/VIAL</u>	<u>A065348 001</u>	Jan 25, 2010
<u>AP</u>	HOSPIRA INC	<u>EQ 500MG BASE/VIAL</u>	<u>A065290 001</u>	Aug 11, 2006
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065290 002</u>	Aug 11, 2006
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065293 001</u>	Aug 10, 2006
<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 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
	<u>CLAFORAN</u>			
<u>AP</u>	+ SANOFI AVENTIS US	<u>EQ 500MG BASE/VIAL</u>	<u>N050547 001</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062659 001</u>	Jan 13, 1987
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N050547 002</u>	
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062659 002</u>	Jan 13, 1987
<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
	<u>CEFOTAXIME</u>			
	+ APP PHARMS	<u>EQ 20GM BASE/VIAL</u>	<u>A064201 002</u>	Mar 24, 2000
	CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER			
	+ SANOFI AVENTIS US	<u>EQ 20MG BASE/ML</u>	<u>N050596 002</u>	May 20, 1985
		<u>EQ 40MG BASE/ML</u>	<u>N050596 004</u>	May 20, 1985

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTETAN

<u>AP</u>	+ APP PHARMS	<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>	WEST-WARD PHARM CORP	<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>		<u>EQ 10GM BASE/VIAL</u>	<u>A091030 001</u>	Oct 26, 2011
	CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER			
	+ B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>A065430 001</u>	Aug 09, 2007
	+	<u>EQ 2GM BASE/VIAL</u>	<u>A065430 002</u>	Aug 09, 2007

CEFOXITIN

INJECTABLE; INJECTION

CEFOXITIN

<u>AP</u>	ANTIBIOTICOS BRASIL	<u>EQ 1GM BASE/VIAL</u>	<u>A065467 001</u>	Aug 31, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065467 002</u>	Aug 31, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 85 (of 424)

CEFOXITIN

INJECTABLE; INJECTION

CEFOXITIN

<u>AP</u>	ANTIBIOTICOS BRASIL	<u>EQ 10GM BASE/VIAL</u>	<u>A065464 001</u>	Aug 31, 2011
-----------	---------------------	--------------------------	--------------------	--------------

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>	+ APP PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A065012 001</u>	Jul 03, 2000
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>A065012 002</u>	Jul 03, 2000
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>A065011 001</u>	Jul 03, 2000
<u>AP</u>	BAXTER HLTHCARE	<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
<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>CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER</u>				
<u>AP</u>	B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>A065214 001</u>	Mar 10, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065214 002</u>	Mar 10, 2006
MEFOXIN IN PLASTIC CONTAINER				
+ BIONICHE PHARMA		<u>EQ 20MG BASE/ML</u>	<u>A063182 001</u>	Jan 25, 1993
+ BIONICHE PHARMA		<u>EQ 40MG BASE/ML</u>	<u>A063182 002</u>	Jan 25, 1993

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETIL

<u>AB</u>	AUROBINDO PHARMA	<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>	RANBAXY	<u>EQ 50MG BASE/5ML</u>	<u>A065082 001</u>	May 31, 2002
<u>AB</u>		<u>EQ 100MG BASE/5ML</u>	<u>A065082 002</u>	May 31, 2002
<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
<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>	RANBAXY	<u>EQ 100MG BASE</u>	<u>A065083 001</u>	Aug 20, 2003
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A065083 002</u>	Aug 20, 2003
<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>	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>	RANBAXY	<u>125MG/5ML</u>	<u>A065202 001</u>	Jun 30, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 86 (of 424)

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

<u>AB</u>	RANBAXY	<u>250MG/5ML</u>	<u>A065202</u>	<u>002</u>	Jun 30, 2006
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065257</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065257</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	TEVA PHARMS	<u>125MG/5ML</u>	<u>A065236</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065236</u>	<u>002</u>	Dec 08, 2005

TABLET; ORAL

CEFPROZIL

<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A065327</u>	<u>001</u>	Mar 26, 2008
<u>AB</u>		<u>500MG</u>	<u>A065327</u>	<u>002</u>	Mar 26, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A065340</u>	<u>001</u>	May 24, 2007
<u>AB</u>		<u>500MG</u>	<u>A065340</u>	<u>002</u>	May 24, 2007
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A065276</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>	+	<u>500MG</u>	<u>A065276</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>250MG</u>	<u>A065267</u>	<u>001</u>	Dec 19, 2005
<u>AB</u>		<u>500MG</u>	<u>A065267</u>	<u>002</u>	Dec 19, 2005
<u>AB</u>	RANBAXY	<u>250MG</u>	<u>A065198</u>	<u>001</u>	Dec 13, 2006
<u>AB</u>		<u>500MG</u>	<u>A065198</u>	<u>002</u>	Dec 13, 2006
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065235</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>		<u>500MG</u>	<u>A065235</u>	<u>002</u>	Nov 14, 2005
<u>AB</u>	TEVA	<u>250MG</u>	<u>A065208</u>	<u>001</u>	Dec 06, 2005
<u>AB</u>		<u>500MG</u>	<u>A065208</u>	<u>002</u>	Dec 06, 2005
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A065428</u>	<u>001</u>	Jun 14, 2007
<u>AB</u>		<u>500MG</u>	<u>A065428</u>	<u>002</u>	Jun 14, 2007

CEFTAROLINE FOSAMIL

POWDER; IV (INFUSION)

CEREXA	400MG/VIAL	N200327	001	Oct 29, 2010
+	600MG/VIAL	N200327	002	Oct 29, 2010

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

<u>AP</u>	ACS DOBFAR	<u>500MG/VIAL</u>	<u>A062640</u>	<u>001</u>	Nov 20, 1985
<u>AP</u>		<u>1GM/VIAL</u>	<u>A062640</u>	<u>002</u>	Nov 20, 1985
<u>AP</u>		<u>2GM/VIAL</u>	<u>A062640</u>	<u>003</u>	Nov 20, 1985
<u>AP</u>		<u>6GM/VIAL</u>	<u>A062640</u>	<u>004</u>	Feb 03, 1992
<u>AP</u>	AUROBINDO PHARMA	<u>500MG/VIAL</u>	<u>A065481</u>	<u>001</u>	May 28, 2010
<u>AP</u>		<u>1GM/VIAL</u>	<u>A065481</u>	<u>002</u>	May 28, 2010
<u>AP</u>		<u>2GM/VIAL</u>	<u>A065481</u>	<u>003</u>	May 28, 2010
<u>AP</u>		<u>6GM/VIAL</u>	<u>A065482</u>	<u>001</u>	May 28, 2010
<u>AP</u>	WOCKHARDT	<u>1GM/VIAL</u>	<u>A065196</u>	<u>001</u>	Oct 15, 2008
	<u>FORTAZ</u>				
<u>AP</u>	+ GLAXOSMITHKLINE	<u>500MG/VIAL</u>	<u>N050578</u>	<u>001</u>	Jul 19, 1985
<u>AP</u>	+	<u>1GM/VIAL</u>	<u>N050578</u>	<u>002</u>	Jul 19, 1985
<u>AP</u>	+	<u>2GM/VIAL</u>	<u>N050578</u>	<u>003</u>	Jul 19, 1985
<u>AP</u>	+	<u>6GM/VIAL</u>	<u>N050578</u>	<u>004</u>	Jul 19, 1985
	<u>TAZICEF</u>				
<u>AP</u>	HOSPIRA	<u>500MG/VIAL</u>	<u>A062662</u>	<u>001</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>A062662</u>	<u>002</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>A064032</u>	<u>001</u>	Oct 31, 1993
<u>AP</u>		<u>2GM/VIAL</u>	<u>A062662</u>	<u>003</u>	Mar 06, 1986
<u>AP</u>		<u>2GM/VIAL</u>	<u>A064032</u>	<u>002</u>	Oct 31, 1993
<u>AP</u>		<u>6GM/VIAL</u>	<u>A062662</u>	<u>004</u>	Mar 06, 1986
	CEFTAZIDIME IN DEXTROSE CONTAINER				
	B BRAUN	EQ 1GM BASE	N050823	001	Jun 13, 2011
	+	EQ 2GM BASE	N050823	002	Jun 13, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 87 (of 424)

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION			
FORTAZ IN PLASTIC CONTAINER			
+ GLAXOSMITHKLINE	EQ 20MG BASE/ML	N050634 002	Apr 28, 1989
+	EQ 40MG BASE/ML	N050634 003	Apr 28, 1989

CEFTIBUTEN DIHYDRATE

CAPSULE; 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

CEFTRIAXONE SODIUM

INJECTABLE; IM-IV

CEFTRIAXONE

AP APP PHARMS	<u>EQ 250MG BASE/VIAL</u>	<u>A065245 001</u>	Feb 15, 2006
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065245 002</u>	Feb 15, 2006
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065245 003</u>	Feb 15, 2006
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065245 004</u>	Feb 15, 2006
AP AUROBINDO PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A065505 001</u>	Jul 31, 2008
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065505 002</u>	Jul 31, 2008
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065505 003</u>	Jul 31, 2008
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065505 004</u>	Jul 31, 2008
AP BEDFORD	<u>EQ 250MG BASE/VIAL</u>	<u>A065465 001</u>	Aug 18, 2008
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065465 002</u>	Aug 18, 2008
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065465 003</u>	Aug 18, 2008
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065465 004</u>	Aug 18, 2008
AP CEPHAZONE PHARMA	<u>EQ 250MG BASE/VIAL</u>	<u>A065294 001</u>	Mar 26, 2007
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065294 002</u>	Mar 26, 2007
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065294 003</u>	Mar 26, 2007
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065294 004</u>	Mar 26, 2007
AP HIKMA FARMACEUTICA	<u>EQ 250MG BASE/VIAL</u>	<u>A065342 001</u>	Jan 10, 2008
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065342 002</u>	Jan 10, 2008
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065342 003</u>	Jan 10, 2008
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065342 004</u>	Jan 10, 2008
AP HOSPIRA INC	<u>EQ 250MG BASE/VIAL</u>	<u>A065230 001</u>	Aug 02, 2005
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065230 002</u>	Aug 02, 2005
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065230 003</u>	Aug 02, 2005
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065230 004</u>	Aug 02, 2005
AP LUITPOLD	<u>EQ 250MG BASE/VIAL</u>	<u>A065305 001</u>	Jan 11, 2008
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065305 002</u>	Jan 11, 2008
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065305 003</u>	Jan 11, 2008
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065305 004</u>	Jan 11, 2008
AP LUPIN	<u>EQ 250MG BASE/VIAL</u>	<u>A065125 001</u>	Sep 30, 2003
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065125 002</u>	Sep 30, 2003
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065125 003</u>	Sep 30, 2003
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065125 004</u>	Sep 30, 2003
AP + SANDOZ	<u>EQ 250MG BASE/VIAL</u>	<u>A065169 001</u>	May 09, 2005
AP +	<u>EQ 500MG BASE/VIAL</u>	<u>A065169 002</u>	May 09, 2005
AP +	<u>EQ 1GM BASE/VIAL</u>	<u>A065169 003</u>	May 09, 2005
AP +	<u>EQ 2GM BASE/VIAL</u>	<u>A065169 004</u>	May 09, 2005
AP STERI PHARMA	<u>EQ 1GM BASE/VIAL</u>	<u>A065268 001</u>	Feb 28, 2007
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065268 002</u>	Feb 28, 2007
AP TEVA	<u>EQ 1GM BASE/VIAL</u>	<u>A065262 001</u>	Jun 29, 2006
AP	<u>EQ 2GM BASE/VIAL</u>	<u>A065262 002</u>	Jun 29, 2006
AP TEVA PARENTERAL	<u>EQ 250MG BASE/VIAL</u>	<u>A065227 001</u>	Mar 15, 2007
AP	<u>EQ 500MG BASE/VIAL</u>	<u>A065227 002</u>	Mar 15, 2007
AP	<u>EQ 1GM BASE/VIAL</u>	<u>A065227 003</u>	Mar 15, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 88 (of 424)

CEFTRIAXONE SODIUM

INJECTABLE; IM-IV

CEFTRIAXONE

<u>AP</u>	TEVA PARENTERAL	<u>EQ 2GM BASE/VIAL</u>	<u>A065227</u>	<u>004</u>	Mar 15, 2007
<u>AP</u>	WOCKHARDT	<u>EQ 250MG BASE/VIAL</u>	<u>A065391</u>	<u>001</u>	Apr 12, 2007
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065391</u>	<u>002</u>	Apr 12, 2007
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065391</u>	<u>003</u>	Apr 12, 2007

INJECTABLE; INJECTION

CEFTRIAXONE

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065329</u>	<u>001</u>	Jul 24, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065329</u>	<u>002</u>	Jul 24, 2008
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065329</u>	<u>003</u>	Jul 24, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065328</u>	<u>001</u>	Jul 24, 2008
<u>AP</u>	APP PHARMS	<u>EQ 10GM BASE/VIAL</u>	<u>A065252</u>	<u>001</u>	Feb 15, 2006
<u>AP</u>	AUROBINDO PHARMA	<u>EQ 10GM BASE/VIAL</u>	<u>A065504</u>	<u>001</u>	Jul 31, 2008
<u>AP</u>	BEDFORD	<u>EQ 10GM BASE/VIAL</u>	<u>A065475</u>	<u>001</u>	Aug 18, 2008
<u>AP</u>	HOSPIRA INC	<u>EQ 1GM BASE/VIAL</u>	<u>A065231</u>	<u>001</u>	Aug 02, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065231</u>	<u>002</u>	Aug 02, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065232</u>	<u>001</u>	Aug 02, 2005
<u>AP</u>	LUPIN	<u>EQ 10GM BASE/VIAL</u>	<u>A065263</u>	<u>001</u>	Sep 12, 2006
<u>AP</u>	SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A065204</u>	<u>001</u>	May 03, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065204</u>	<u>002</u>	May 03, 2005
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>A065168</u>	<u>001</u>	May 17, 2005
<u>AP</u>	STERI PHARMA	<u>EQ 10GM BASE/VIAL</u>	<u>A065269</u>	<u>001</u>	Feb 28, 2007
<u>AP</u>	TEVA	<u>EQ 10GM BASE/VIAL</u>	<u>A065274</u>	<u>001</u>	May 01, 2006
<u>AP</u>	WOCKHARDT	<u>EQ 1GM BASE/VIAL</u>	<u>A065180</u>	<u>001</u>	May 12, 2006
		<u>CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER</u>			
<u>AP</u>	+ B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N050796</u>	<u>001</u>	Apr 20, 2005
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>N050796</u>	<u>002</u>	Apr 20, 2005
		<u>ROCEPHIN</u>			
<u>AP</u>	+ HOFFMANN LA ROCHE	<u>EQ 500MG BASE/VIAL</u>	<u>A063239</u>	<u>002</u>	Aug 13, 1993
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>A063239</u>	<u>003</u>	Aug 13, 1993
		<u>CEFTRIAXONE IN PLASTIC CONTAINER</u>			
	BAXTER HLTHCARE	<u>EQ 20MG BASE/ML</u>	<u>A065224</u>	<u>001</u>	Aug 23, 2005
		<u>EQ 40MG BASE/ML</u>	<u>A065224</u>	<u>002</u>	Aug 23, 2005

CEFUROXIME AXETIL

FOR SUSPENSION; ORAL

CEFTIN

<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 125MG BASE/5ML</u>	<u>N050672</u>	<u>001</u>	Jun 30, 1994
<u>AB</u>	+	<u>EQ 250MG BASE/5ML</u>	<u>N050672</u>	<u>002</u>	Apr 29, 1997

CEFUROXIME AXETIL

<u>AB</u>	RANBAXY	<u>EQ 125MG BASE/5ML</u>	<u>A065323</u>	<u>001</u>	Feb 05, 2008
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065323</u>	<u>002</u>	Feb 05, 2008

TABLET; ORAL

CEFTIN

<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 125MG BASE</u>	<u>N050605</u>	<u>001</u>	Dec 28, 1987
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N050605</u>	<u>002</u>	Dec 28, 1987
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N050605</u>	<u>003</u>	Dec 28, 1987

CEFUROXIME AXETIL

<u>AB</u>	ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A065496</u>	<u>001</u>	Jun 07, 2010
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065496</u>	<u>002</u>	Jun 07, 2010

APOTEX

<u>AB</u>	APOTEX	<u>EQ 250MG BASE</u>	<u>A065069</u>	<u>001</u>	Oct 02, 2002
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065069</u>	<u>002</u>	Oct 02, 2002

AUROBINDO PHARMA LTD

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 125MG BASE</u>	<u>A065308</u>	<u>001</u>	Mar 29, 2006
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065308</u>	<u>002</u>	Mar 29, 2006

LUPIN

<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>A065135</u>	<u>001</u>	Jul 25, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065135</u>	<u>002</u>	Jul 25, 2003

ORCHID HLTHCARE

<u>AB</u>	ORCHID HLTHCARE	<u>EQ 125MG BASE</u>	<u>A065359</u>	<u>001</u>	Feb 15, 2008
-----------	-----------------	----------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 89 (of 424)

CEFUROXIME AXETIL

TABLET; ORAL

CEFUROXIME AXETIL

<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE</u>	<u>A065359 002</u>	Feb 15, 2008
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065359 003</u>	Feb 15, 2008
<u>AB</u>	RANBAXY	<u>EQ 125MG BASE</u>	<u>A065043 003</u>	Feb 15, 2002
<u>AB</u>		<u>EQ 125MG BASE</u>	<u>A065118 001</u>	Apr 25, 2003
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065043 002</u>	Feb 15, 2002
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065118 002</u>	Apr 25, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065043 001</u>	Feb 15, 2002
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065118 003</u>	Apr 25, 2003
<u>AB</u>	TEVA	<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>	WOCKHARDT	<u>EQ 125MG BASE</u>	<u>A065166 001</u>	Jul 29, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065166 002</u>	Jul 29, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065166 003</u>	Jul 29, 2005

CEFUROXIME SODIUM

INJECTABLE; IM-IV

CEFUROXIME SODIUM

<u>AB</u>	APP PHARMS	<u>EQ 750MG BASE/VIAL</u>	<u>A065001 001</u>	May 30, 2001
<u>AB</u>	HIKMA FARMACEUTICA	<u>EQ 750MG BASE/VIAL</u>	<u>A065048 001</u>	Jan 09, 2004
<u>AB</u>	STERI PHARMA	<u>EQ 750MG BASE/VIAL</u>	<u>A064125 001</u>	May 30, 1997
<u>AB</u>	TEVA	<u>EQ 750MG BASE/VIAL</u>	<u>A064192 002</u>	Apr 16, 1998
	ZINACEF			
<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 750MG BASE/VIAL</u>	<u>N050558 002</u>	Oct 19, 1983
	CEFUROXIME SODIUM			
<u>AP</u>	HOSPIRA INC	<u>EQ 750MG BASE/VIAL</u>	<u>A065483 001</u>	Oct 15, 2008

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>	APP PHARMS	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065001 002</u>	May 30, 2001
<u>AP</u>		<u>EQ 7.5GM BASE/VIAL</u>	<u>A065002 001</u>	Sep 28, 1998

<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065048 002</u>	Jan 09, 2004
<u>AP</u>	HOSPIRA INC	<u>EQ 7.5GM BASE/VIAL</u>	<u>A065046 001</u>	Jan 09, 2004
<u>AP</u>		<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
<u>AP</u>	STERI PHARMA	<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>	TEVA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A064192 001</u>	Apr 16, 1998
<u>AP</u>		<u>EQ 7.5GM BASE/VIAL</u>	<u>A064191 001</u>	Apr 16, 1998

ZINACEF

<u>AP</u>	+ GLAXOSMITHKLINE	<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
	CEFUROXIME SODIUM IN PLASTIC CONTAINER			
+ SAMSON MEDCL		<u>EQ 75GM BASE/VIAL</u>	<u>A065251 001</u>	Dec 30, 2009
+		<u>EQ 225GM BASE/VIAL</u>	<u>A065251 002</u>	Dec 30, 2009
	ZINACEF IN PLASTIC CONTAINER			
+ GLAXOSMITHKLINE		<u>EQ 15MG BASE/ML</u>	<u>N050643 001</u>	Apr 28, 1989
+		<u>EQ 30MG BASE/ML</u>	<u>N050643 002</u>	Apr 28, 1989

CELECOXIB

CAPSULE; ORAL

CELEBREX

GD SEARLE	50MG	<u>N020998 004</u>	Dec 15, 2006
	100MG	<u>N020998 001</u>	Dec 31, 1998
	200MG	<u>N020998 002</u>	Dec 31, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 90 (of 424)

CELECOXIB

CAPSULE; ORAL
 CELEBREX
 + GD SEARLE 400MG N020998 003 Aug 29, 2002

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM	<u>EQ 250MG BASE</u>	<u>A090836</u> <u>001</u>	Dec 20, 2010
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A090836</u> <u>002</u>	Dec 20, 2010
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A065253</u> <u>001</u>	Nov 16, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065253</u> <u>002</u>	Nov 16, 2005
<u>AB</u>	BELCHER PHARMS	<u>EQ 250MG BASE</u>	<u>A062713</u> <u>001</u>	Jul 15, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062713</u> <u>002</u>	Jul 15, 1988
<u>AB</u>	FACTA FARMA	<u>EQ 250MG BASE</u>	<u>A062118</u> <u>001</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062118</u> <u>002</u>	
<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>A065215</u> <u>001</u>	Jan 24, 2006
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065215</u> <u>002</u>	Jan 24, 2006
<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>A065229</u> <u>001</u>	Nov 25, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065229</u> <u>002</u>	Nov 25, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE</u>	<u>A065248</u> <u>001</u>	Jun 28, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065248</u> <u>002</u>	Jun 28, 2005
<u>AB</u>	RANBAXY	<u>EQ 250MG BASE</u>	<u>A065007</u> <u>001</u>	Sep 16, 1999
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065007</u> <u>002</u>	Sep 16, 1999
<u>AB</u>	STEVENS J	<u>EQ 250MG BASE</u>	<u>A062870</u> <u>001</u>	Mar 17, 1988
<u>AB</u>	SUN PHARM INDNS (IN)	<u>EQ 250MG BASE</u>	<u>A062791</u> <u>001</u>	Jun 11, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062791</u> <u>002</u>	Jun 11, 1987
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A062702</u> <u>001</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062702</u> <u>002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 250MG BASE</u>	<u>A065152</u> <u>001</u>	Feb 24, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065152</u> <u>002</u>	Feb 24, 2005

KEFLEX

<u>AB</u>	SHIONOGI INC	<u>EQ 250MG BASE</u>	<u>N050405</u> <u>002</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N050405</u> <u>003</u>	

KEFLEX

+ SHIONOGI INC EQ 750MG BASE N050405 005 May 12, 2006

FOR SUSPENSION; ORAL

CEPHALEXIN

<u>AB</u>	FACTA FARMA	<u>EQ 125MG BASE/5ML</u>	<u>A062117</u> <u>002</u>	
<u>AB</u>	+	<u>EQ 250MG BASE/5ML</u>	<u>A062117</u> <u>003</u>	
<u>AB</u>	HIKMA PHARMS	<u>EQ 125MG BASE/5ML</u>	<u>A065444</u> <u>001</u>	Aug 28, 2009
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065444</u> <u>002</u>	Aug 28, 2009
<u>AB</u>	LUPIN	<u>EQ 125MG BASE/5ML</u>	<u>A065234</u> <u>001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065234</u> <u>002</u>	Aug 17, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 125MG BASE/5ML</u>	<u>A065326</u> <u>001</u>	Jul 10, 2006
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065326</u> <u>002</u>	Jul 10, 2006
<u>AB</u>	RANBAXY	<u>EQ 125MG BASE/5ML</u>	<u>A065081</u> <u>001</u>	Jul 27, 2001
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065081</u> <u>002</u>	Jul 27, 2001
<u>AB</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>A062703</u> <u>001</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A062703</u> <u>002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 125MG BASE/5ML</u>	<u>A065336</u> <u>001</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065336</u> <u>002</u>	Jul 25, 2007

TABLET; ORAL

CEPHALEXIN

<u>AB</u>	TEVA	EQ 250MG BASE	A063023 001	Jan 12, 1989
<u>AB</u>	+	EQ 500MG BASE	A063024 001	Jan 12, 1989

PRESCRIPTION DRUG PRODUCT LIST

3 - 91 (of 424)

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>5MG/5ML</u>	<u>A090766</u>	<u>001</u>	Oct 07, 2009
<u>AA</u>	APOTEX INC	<u>5MG/5ML</u>	<u>A078412</u>	<u>001</u>	Jun 18, 2008
<u>AA</u>	AUROBINDO PHARMA	<u>5MG/5ML</u>	<u>A090751</u>	<u>001</u>	Dec 16, 2009
<u>AA</u>	CYPRESS PHARM	<u>5MG/5ML</u>	<u>A078488</u>	<u>001</u>	Oct 06, 2008
<u>AA</u>	DR REDDYS LABS LTD	<u>5MG/5ML</u>	<u>A078870</u>	<u>001</u>	Apr 27, 2009
<u>AA</u>	PERRIGO R AND D	<u>5MG/5ML</u>	<u>A078398</u>	<u>001</u>	Jun 17, 2008
<u>AA</u>	RANBAXY	<u>5MG/5ML</u>	<u>A077472</u>	<u>001</u>	Jun 18, 2008
<u>AA</u>	SUN PHARM INDs INC	<u>5MG/5ML</u>	<u>A090191</u>	<u>001</u>	Nov 12, 2009
<u>AA</u>	TARO	<u>5MG/5ML</u>	<u>A076601</u>	<u>001</u>	Jun 20, 2008
<u>AA</u>	TEVA PHARMS	<u>5MG/5ML</u>	<u>A077279</u>	<u>001</u>	May 27, 2008
<u>AA</u>	VINTAGE	<u>5MG/5ML</u>	<u>A078496</u>	<u>001</u>	Sep 25, 2009
<u>AA</u>	WOCKHARDT	<u>5MG/5ML</u>	<u>A078757</u>	<u>001</u>	Aug 28, 2009
	<u>ZYRTEC</u>				
<u>AA</u>	+ MCNEIL CONSUMER	<u>5MG/5ML</u>	<u>N020346</u>	<u>001</u>	Sep 27, 1996

CETRORELIX

INJECTABLE; INJECTION

CETROTIDE

+ EMD SERONO	EQ 0.25MG BASE/ML	N021197	001	Aug 11, 2000
+	EQ 3MG BASE/ML	N021197	002	Aug 11, 2000

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL

CEVIMELINE

<u>AB</u>	APOTEX INC	<u>30MG</u>	<u>A091260</u>	<u>001</u>	Aug 25, 2011
<u>AB</u>	+ DAIICHI SANKYO CO	<u>30MG</u>	<u>N020989</u>	<u>002</u>	Jan 11, 2000

CHENODIOL

TABLET; ORAL

CHENODIOL

+ NEXGEN PHARMA	250MG	A091019	001	Oct 22, 2009
-----------------	-------	---------	-----	--------------

CHLORAMBUCIL

TABLET; ORAL

LEUKERAN

+ PBS	2MG	N010669	002	
-------	-----	---------	-----	--

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

+ APP PHARMS	EQ 1GM BASE/VIAL	A062365	001	Aug 25, 1982
--------------	------------------	---------	-----	--------------

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

<u>AB</u>	BARR	<u>5MG</u>	<u>A084768</u>	<u>001</u>
<u>AB</u>		<u>10MG</u>	<u>A083116</u>	<u>001</u>
<u>AB</u>		<u>25MG</u>	<u>A084769</u>	<u>001</u>
<u>AB</u>	USL PHARMA	<u>10MG</u>	<u>A084623</u>	<u>001</u>
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>A086383</u>	<u>001</u>
<u>AB</u>		<u>10MG</u>	<u>A086294</u>	<u>001</u>
<u>AB</u>		<u>25MG</u>	<u>A086382</u>	<u>001</u>
	<u>LIBRIUM</u>			
<u>AB</u>	VALEANT PHARM INTL	<u>5MG</u>	<u>A085461</u>	<u>001</u>
<u>AB</u>		<u>10MG</u>	<u>A085472</u>	<u>001</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 93 (of 424)

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

<u>AB</u>	MYLAN	<u>500MG</u>	<u>A084217 001</u>	
<u>AB</u>	WEST WARD	<u>250MG</u>	<u>A086028 001</u>	Jul 14, 1982
<u>AB</u>		<u>500MG</u>	<u>A087736 001</u>	Jul 14, 1982

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

<u>AP</u>	APP PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A090896 001</u>	Oct 16, 2009
<u>AP</u>	SUN PHARMA GLOBAL	<u>EQ 500MG BASE/VIAL</u>	<u>A091546 001</u>	Jul 26, 2011
	<u>DIURIL</u>			
<u>AP</u>	+ LUNDBECK INC	<u>EQ 500MG BASE/VIAL</u>	<u>N011145 005</u>	

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

ZUTRIPRO

+ CYPRESS PHARM	4MG/5ML;5MG/5ML;60MG/5ML	N022439 001	Jun 08, 2011
-----------------	--------------------------	-------------	--------------

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

CAPSULE, EXTENDED RELEASE; ORAL

TUSSICAPS

HI-TECH PHARMA CO	<u>EQ 4MG MALEATE;EQ 5MG BITARTRATE</u>	A077273 002	Sep 24, 2007
+	<u>EQ 8MG MALEATE;EQ 10MG BITARTRATE</u>	A077273 001	Sep 24, 2007

SUSPENSION, EXTENDED RELEASE; ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

<u>AB</u>	TRIS PHARMA INC	<u>EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML</u>	<u>A091632 001</u>	Oct 01, 2010
<u>AB</u>	+ UCB INC	<u>EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML</u>	<u>N019111 001</u>	Dec 31, 1987

CHLORPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

+ BAXTER HLTHCARE	25MG/ML	A083329 001
-------------------	---------	-------------

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

BP	SANDOZ	10MG	A080439 001
BP		25MG	A080439 002
BP		50MG	A080439 003
BP	+	100MG	A080439 004
BP		200MG	A080439 005
BP	USL PHARMA	10MG	A083386 001
BP		25MG	A084112 001
BP		50MG	A084113 001
BP		100MG	A084114 001
BP		200MG	A084115 001

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

<u>AB</u>	MYLAN	<u>100MG</u>	<u>A088549 002</u>	Jun 01, 1984
<u>AB</u>		<u>250MG</u>	<u>A088549 001</u>	Jun 01, 1984
<u>AB</u>	PLIVA	<u>100MG</u>	<u>A088921 001</u>	Apr 12, 1985
<u>AB</u>		<u>250MG</u>	<u>A088922 001</u>	Apr 12, 1985
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>A088852 001</u>	Sep 26, 1984
<u>AB</u>		<u>250MG</u>	<u>A088826 001</u>	Sep 26, 1984

PRESCRIPTION DRUG PRODUCT LIST

3 - 94 (of 424)

CHLORPROPAMIDE

TABLET; ORAL			
<u>DIABINESE</u>			
<u>AB</u>	PFIZER	<u>100MG</u>	<u>N011641 003</u>
<u>AB</u> +		<u>250MG</u>	<u>N011641 006</u>

CHLORTHALIDONE

TABLET; ORAL			
<u>CHLORTHALIDONE</u>			
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A086831 002</u>
<u>AB</u> +		<u>50MG</u>	<u>A086831 001</u>
<u>AB</u>	PLIVA	<u>25MG</u>	<u>A088902 001</u>
<u>AB</u>		<u>50MG</u>	<u>A088903 001</u>
THALITONE			Sep 19, 1985
+ MONARCH PHARMS		15MG	<u>N019574 001</u>
			Dec 20, 1988

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL			
CLORPRES			
	MYLAN	15MG; 0.1MG	<u>A071325 003</u>
		15MG; 0.2MG	<u>A071325 002</u>
+		15MG; 0.3MG	<u>A071325 001</u>

CHLORZOXAZONE

TABLET; ORAL			
<u>CHLORZOXAZONE</u>			
<u>AA</u>	BARR	<u>500MG</u>	<u>A089895 001</u>
<u>AA</u>	WATSON LABS	<u>500MG</u>	<u>A040137 001</u>
<u>AA</u>		<u>500MG</u>	<u>A081040 001</u>
<u>AA</u>		<u>500MG</u>	<u>A089859 001</u>
<u>PARAFON FORTE DSC</u>			
<u>AA</u> +	JANSSEN R AND D	<u>500MG</u>	<u>N011529 002</u>
CHLORZOXAZONE			
	MIKART	375MG	<u>A040861 001</u>
		750MG	<u>A040861 002</u>
			Jun 01, 2010
			Jun 01, 2010

CHOLESTYRAMINE

POWDER; ORAL			
<u>CHOLESTYRAMINE</u>			
<u>AB</u>	PAR PHARM	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077204 002</u>
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>A077204 001</u>
<u>AB</u> +	SANDOZ	<u>EQ 4GM RESIN/PACKET</u>	<u>A074557 001</u>
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074557 002</u>
<u>CHOLESTYRAMINE LIGHT</u>			
<u>AB</u>	PAR PHARM	<u>EQ 4GM RESIN/PACKET</u>	<u>A077203 001</u>
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A077203 002</u>
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074558 002</u>
<u>AB</u> +		<u>EQ 4GM RESIN/PACKET</u>	<u>A074558 001</u>
<u>LOCHOLEST</u>			
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074561 002</u>
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>A074561 001</u>
<u>LOCHOLEST LIGHT</u>			
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/PACKET</u>	<u>A074562 001</u>
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A074562 002</u>
<u>PREVALITE</u>			
<u>AB</u>	UPSHER SMITH	<u>EQ 4GM RESIN/PACKET</u>	<u>A073263 001</u>
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>A073263 002</u>
			Feb 22, 1996
			Oct 30, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 95 (of 424)

CHOLINE FENOFLIBRATE

CAPSULE, DELAYED RELEASE; ORAL TRILIPIX		
ABBOTT LABS	EQ 45MG FENOFLIBRIC ACID	N022224 001 Dec 15, 2008
+	EQ 135MG FENOFLIBRIC ACID	N022224 002 Dec 15, 2008

CHORIOGONADOTROPIN ALFA

INJECTABLE; SUBCUTANEOUS OVIDREL		
+	EMD SERONO EQ 0.25MG / 0.5ML	N021149 002 Oct 06, 2003

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		
NYCOMED US	0.08MG/INH	N021658 002 Jan 10, 2008
+	0.16MG/INH	N021658 003 Jan 10, 2008
SPRAY, METERED; NASAL OMNARIS		
+	NYCOMED US 0.05MG/INH	N022004 001 Oct 20, 2006

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

<u>AB</u> ALTANA	<u>0.77%</u>	<u>A076435</u> <u>001</u> Dec 29, 2004
<u>AB</u> G AND W LABS INC	<u>0.77%</u>	<u>A078463</u> <u>001</u> Dec 20, 2010
<u>AB</u> GLENMARK PHARMS	<u>0.77%</u>	<u>A090273</u> <u>001</u> Nov 10, 2009
<u>AB</u> PERRIGO	<u>0.77%</u>	<u>A077364</u> <u>001</u> Mar 03, 2006
<u>AB</u> TARO	<u>0.77%</u>	<u>A076790</u> <u>001</u> Apr 12, 2005
<u>LOPROX</u>		
<u>AB</u> + MEDICIS	<u>0.77%</u>	<u>N018748</u> <u>001</u> Dec 30, 1982

GEL; TOPICAL

CICLOPIROX

<u>AB</u> NYCOMED US	<u>0.77%</u>	<u>A077896</u> <u>001</u> Jun 10, 2008
<u>AB</u> PADDICK LLC	<u>0.77%</u>	<u>A078266</u> <u>001</u> Jan 07, 2009

LOPROX

<u>AB</u> + MEDICIS	<u>0.77%</u>	<u>N020519</u> <u>001</u> Jul 21, 1997
---------------------	--------------	--

SHAMPOO; TOPICAL

CICLOPIROX

<u>AT</u> NYCOMED US	<u>1%</u>	<u>A090146</u> <u>001</u> May 25, 2010
<u>AT</u> PADDICK LLC	<u>1%</u>	<u>A090490</u> <u>001</u> Nov 24, 2009
<u>AT</u> PERRIGO	<u>1%</u>	<u>A078594</u> <u>001</u> Feb 16, 2010
<u>AT</u> TARO	<u>1%</u>	<u>A090269</u> <u>001</u> Feb 23, 2011
<u>LOPROX</u>		
<u>AT</u> + MEDICIS	<u>1%</u>	<u>N021159</u> <u>001</u> Feb 28, 2003

SOLUTION; TOPICAL

CICLOPIROX

<u>AT</u> ACTAVIS MID ATLANTIC	<u>8%</u>	<u>A078046</u> <u>001</u> Sep 18, 2007
<u>AT</u> APOTEX CORP	<u>8%</u>	<u>A078172</u> <u>001</u> Sep 18, 2007
<u>AT</u> G AND W LABS	<u>8%</u>	<u>A078233</u> <u>001</u> Sep 18, 2007
<u>AT</u> HI TECH PHARMA	<u>8%</u>	<u>A078270</u> <u>001</u> Sep 18, 2007
<u>AT</u> PERRIGO NEW YORK	<u>8%</u>	<u>A077623</u> <u>001</u> Sep 18, 2007
<u>AT</u> SYNERX PHARMA	<u>8%</u>	<u>A078567</u> <u>001</u> Sep 18, 2007
<u>AT</u> TARO PHARM IND	<u>8%</u>	<u>A078144</u> <u>001</u> Sep 18, 2007
<u>AT</u> TEVA PHARMS	<u>8%</u>	<u>A078079</u> <u>001</u> Sep 18, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 96 (of 424)

CICLOPIROX

SOLUTION; TOPICAL

CICLOPIROX

<u>AT</u>	TOLMAR	<u>8%</u>	<u>A077687</u>	<u>001</u>	Sep 18, 2007
<u>AT</u>	VERSAPHARM	<u>8%</u>	<u>A078975</u>	<u>001</u>	Feb 17, 2010
<u>AT</u>	WATSON LABS	<u>8%</u>	<u>A078124</u>	<u>001</u>	Sep 18, 2007
	<u>PENLAC</u>				
<u>AT</u>	+ SANOFI AVENTIS US	<u>8%</u>	<u>N021022</u>	<u>001</u>	Dec 17, 1999
	SUSPENSION; TOPICAL				
	<u>CICLOPIROX</u>				
<u>AB</u>	ALTANA	<u>0.77%</u>	<u>A076422</u>	<u>001</u>	Aug 06, 2004
<u>AB</u>	PERRIGO NEW YORK	<u>0.77%</u>	<u>A077676</u>	<u>001</u>	Dec 15, 2006
<u>AB</u>	TARO	<u>0.77%</u>	<u>A077092</u>	<u>001</u>	Aug 10, 2005
	<u>LOPROX</u>				
<u>AB</u>	+ MEDICIS	<u>0.77%</u>	<u>N019824</u>	<u>001</u>	Dec 30, 1988

CIDOFOVIR

INJECTABLE; INJECTION

VISTIDE

+ GILEAD

EQ 75MG BASE/ML

N020638 001 Jun 26, 1996

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

IMIPENEM AND CILASTATIN

<u>AP</u>	ACS DOBFAR	<u>EQ 250MG BASE/VIAL;250MG/VIAL</u>	<u>A090577</u>	<u>001</u>	Dec 21, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL;500MG/VIAL</u>	<u>A090577</u>	<u>002</u>	Dec 21, 2011
<u>AP</u>	HOSPIRA INC	<u>EQ 250MG BASE/VIAL;250MG/VIAL</u>	<u>A090825</u>	<u>001</u>	Nov 16, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL;500MG/VIAL</u>	<u>A090825</u>	<u>002</u>	Nov 16, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL;500MG/VIAL</u>	<u>A091007</u>	<u>001</u>	Nov 16, 2011
	<u>PRIMAXIN</u>				
<u>AP</u>	MERCK	<u>EQ 250MG BASE/VIAL;250MG/VIAL</u>	<u>A062756</u>	<u>001</u>	Jan 08, 1987
<u>AP</u>	+	<u>EQ 250MG BASE/VIAL;250MG/VIAL</u>	<u>N050587</u>	<u>001</u>	Nov 26, 1985
<u>AP</u>		<u>EQ 500MG BASE/VIAL;500MG/VIAL</u>	<u>A062756</u>	<u>002</u>	Jan 08, 1987
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL;500MG/VIAL</u>	<u>N050587</u>	<u>002</u>	Nov 26, 1985
<u>AP</u>		<u>EQ 500MG BASE/VIAL;500MG/VIAL</u>	<u>N050630</u>	<u>001</u>	Dec 14, 1990

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

<u>AB</u>	ALPHAPHARM	<u>50MG</u>	<u>A077019</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>		<u>100MG</u>	<u>A077019</u>	<u>002</u>	Nov 23, 2004
<u>AB</u>	APOTEX INC	<u>50MG</u>	<u>A077030</u>	<u>001</u>	Dec 10, 2004
<u>AB</u>		<u>100MG</u>	<u>A077030</u>	<u>002</u>	Dec 10, 2004
<u>AB</u>	BRECKENRIDGE PHARM	<u>50MG</u>	<u>A077708</u>	<u>001</u>	Sep 28, 2009
<u>AB</u>		<u>100MG</u>	<u>A077708</u>	<u>002</u>	Sep 28, 2009
<u>AB</u>	COREPHARMA	<u>50MG</u>	<u>A077150</u>	<u>001</u>	Mar 11, 2005
<u>AB</u>		<u>100MG</u>	<u>A077022</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A077323</u>	<u>002</u>	Apr 20, 2006
<u>AB</u>		<u>100MG</u>	<u>A077323</u>	<u>001</u>	Apr 20, 2006
<u>AB</u>	PLIVA HRVATSKA DOO	<u>50MG</u>	<u>A077898</u>	<u>001</u>	Oct 29, 2007
<u>AB</u>		<u>100MG</u>	<u>A077898</u>	<u>002</u>	Oct 29, 2007
<u>AB</u>	ROXANE	<u>50MG</u>	<u>A077024</u>	<u>001</u>	May 17, 2005
<u>AB</u>		<u>100MG</u>	<u>A077024</u>	<u>002</u>	May 17, 2005
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>A077310</u>	<u>001</u>	Nov 08, 2005
<u>AB</u>		<u>100MG</u>	<u>A077021</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>	TEVA	<u>50MG</u>	<u>A077027</u>	<u>001</u>	Nov 24, 2004
<u>AB</u>		<u>100MG</u>	<u>A077027</u>	<u>002</u>	Nov 24, 2004
	<u>PLETAL</u>				
<u>AB</u>	+ OTSUKA	<u>50MG</u>	<u>N020863</u>	<u>001</u>	Jan 15, 1999
<u>AB</u>	+	<u>100MG</u>	<u>N020863</u>	<u>002</u>	Jan 15, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 97 (of 424)

CIMETIDINE

TABLET; ORAL

CIMETIDINE

<u>AB</u>	APOTEX	<u>200MG</u>	<u>A074890</u>	<u>001</u>	Dec 18, 1998
<u>AB</u>		<u>300MG</u>	<u>A074890</u>	<u>002</u>	Dec 18, 1998
<u>AB</u>		<u>400MG</u>	<u>A074890</u>	<u>003</u>	Dec 18, 1998
<u>AB</u>		<u>800MG</u>	<u>A074890</u>	<u>004</u>	Dec 18, 1998
<u>AB</u>	DAVA PHARMS INC	<u>300MG</u>	<u>A074340</u>	<u>001</u>	Jun 23, 1995
<u>AB</u>		<u>400MG</u>	<u>A074340</u>	<u>002</u>	Jun 23, 1995
<u>AB</u>		<u>800MG</u>	<u>A074339</u>	<u>001</u>	Jun 23, 1995
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>200MG</u>	<u>A074424</u>	<u>001</u>	Jul 28, 1995
<u>AB</u>		<u>300MG</u>	<u>A074424</u>	<u>002</u>	Jul 28, 1995
<u>AB</u>		<u>400MG</u>	<u>A074424</u>	<u>003</u>	Jul 28, 1995
<u>AB</u>		<u>800MG</u>	<u>A074424</u>	<u>004</u>	Jul 28, 1995
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A074246</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074246</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074246</u>	<u>003</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>A074246</u>	<u>004</u>	May 17, 1994
<u>AB</u>	PLIVA	<u>200MG</u>	<u>A074568</u>	<u>001</u>	Feb 27, 1997
<u>AB</u>		<u>300MG</u>	<u>A074568</u>	<u>002</u>	Feb 27, 1997
<u>AB</u>		<u>400MG</u>	<u>A074568</u>	<u>003</u>	Feb 27, 1997
<u>AB</u>		<u>800MG</u>	<u>A074566</u>	<u>001</u>	Feb 27, 1997
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A074506</u>	<u>001</u>	Jan 24, 1996
<u>AB</u>		<u>300MG</u>	<u>A074506</u>	<u>002</u>	Jan 24, 1996
<u>AB</u>		<u>400MG</u>	<u>A074506</u>	<u>003</u>	Jan 24, 1996
<u>AB</u>		<u>800MG</u>	<u>A074506</u>	<u>004</u>	Jan 24, 1996
<u>AB</u>	TEVA	<u>200MG</u>	<u>A074151</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>A074151</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>A074151</u>	<u>003</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>A074463</u>	<u>001</u>	May 17, 1994
<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>A074349</u>	<u>001</u>	Aug 30, 1996
<u>AB</u>		<u>300MG</u>	<u>A074349</u>	<u>002</u>	Aug 30, 1996
<u>AB</u>		<u>400MG</u>	<u>A074349</u>	<u>003</u>	Aug 30, 1996
<u>AB</u>		<u>800MG</u>	<u>A074316</u>	<u>001</u>	Feb 28, 1996
	<u>TAGAMET</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>200MG</u>	<u>N017920</u>	<u>002</u>	
<u>AB</u>		<u>300MG</u>	<u>N017920</u>	<u>003</u>	
<u>AB</u>		<u>400MG</u>	<u>N017920</u>	<u>004</u>	Dec 14, 1983
<u>AB</u> +		<u>800MG</u>	<u>N017920</u>	<u>005</u>	Apr 30, 1986

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

<u>AP</u>	DAVA PHARMS INC	<u>EQ 300MG BASE/2ML</u>	<u>A074428</u>	<u>001</u>	Apr 25, 1996
<u>AP</u>	HOSPIRA	<u>EQ 300MG BASE/2ML</u>	<u>A074344</u>	<u>001</u>	Jan 31, 1995
<u>AP</u>		<u>EQ 300MG BASE/2ML</u>	<u>A074345</u>	<u>001</u>	Jan 31, 1995

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

<u>AA</u> +	HI TECH PHARMA	<u>EQ 300MG BASE/5ML</u>	<u>A074664</u>	<u>001</u>	Oct 28, 1997
<u>AA</u>	NOVEX	<u>EQ 300MG BASE/5ML</u>	<u>A075560</u>	<u>001</u>	Mar 15, 2000
<u>AA</u>	PHARM ASSOC	<u>EQ 300MG BASE/5ML</u>	<u>A074553</u>	<u>001</u>	Jan 27, 1997
<u>AA</u>	TEVA	<u>EQ 300MG BASE/5ML</u>	<u>A074610</u>	<u>001</u>	Sep 26, 1996
<u>AA</u>	WOCKHARDT	<u>EQ 300MG BASE/5ML</u>	<u>A074757</u>	<u>001</u>	Oct 17, 1997

CINACALCET HYDROCHLORIDE

TABLET; ORAL

SENSIPAR

	AMGEN	<u>EQ 30MG BASE</u>	<u>N021688</u>	<u>001</u>	Mar 08, 2004
+		<u>EQ 60MG BASE</u>	<u>N021688</u>	<u>002</u>	Mar 08, 2004
+		<u>EQ 90MG BASE</u>	<u>N021688</u>	<u>003</u>	Mar 08, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 98 (of 424)

CIPROFLOXACIN

FOR SUSPENSION; ORAL

CIPRO

BAYER HLTHCARE	250MG/5ML	N020780 001	Sep 26, 1997
+	500MG/5ML	N020780 002	Sep 26, 1997

INJECTABLE; INJECTION

CIPRO

AP + BAYER HLTHCARE	<u>400MG/40ML (10MG/ML)</u>	<u>N019847 001</u>	Dec 26, 1990
AP +	<u>200MG/20ML (10MG/ML)</u>	<u>N019847 002</u>	Dec 26, 1990
	<u>CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP + BAYER HLTHCARE	<u>200MG/100ML</u>	<u>N019857 001</u>	Dec 26, 1990
AP +	<u>400MG/200ML</u>	<u>N019857 002</u>	Dec 26, 1990
	<u>CIPROFLOXACIN</u>		
AP CLARIS LIFESCIENCES	<u>200MG/20ML (10MG/ML)</u>	<u>A078062 001</u>	Apr 29, 2008
AP	<u>400MG/40ML (10MG/ML)</u>	<u>A078062 002</u>	Apr 29, 2008
AP HIKMA FARMACEUTICA	<u>200MG/20ML (10MG/ML)</u>	<u>A076717 001</u>	Dec 22, 2009
AP	<u>400MG/40ML (10MG/ML)</u>	<u>A076717 002</u>	Dec 22, 2009
AP HOSPIRA	<u>400MG/40ML (10MG/ML)</u>	<u>A077245 002</u>	Aug 28, 2006
AP	<u>200MG/20ML (10MG/ML)</u>	<u>A077245 001</u>	Aug 28, 2006
AP TEVA PARENTERAL	<u>400MG/40ML (10MG/ML)</u>	<u>A077782 002</u>	Aug 28, 2006
AP	<u>200MG/20ML (10MG/ML)</u>	<u>A077782 001</u>	Aug 28, 2006
	<u>CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP ACS DOBFAR INFO SA	<u>200MG/100ML</u>	<u>A078252 001</u>	Mar 18, 2008
AP	<u>400MG/200ML</u>	<u>A078252 002</u>	Mar 18, 2008
AP BAXTER HLTHCARE	<u>200MG/100ML</u>	<u>A077888 001</u>	Mar 18, 2008
AP	<u>400MG/200ML</u>	<u>A077888 002</u>	Mar 18, 2008
AP CLARIS LIFESCIENCES	<u>200MG/100ML</u>	<u>A078024 001</u>	Mar 18, 2008
AP	<u>400MG/200ML</u>	<u>A078024 002</u>	Mar 18, 2008
AP HIKMA FARMACEUTICA	<u>400MG/200ML</u>	<u>A078431 001</u>	Nov 18, 2009
AP HOSPIRA	<u>200MG/100ML</u>	<u>A077753 001</u>	Mar 18, 2008
AP	<u>400MG/200ML</u>	<u>A077753 002</u>	Mar 18, 2008

CIPROFLOXACIN HYDROCHLORIDE

OINTMENT; OPHTHALMIC

CILOXAN

+ ALCON PHARMS LTD	EQ 0.3% BASE	N020369 001	Mar 30, 1998
--------------------	--------------	-------------	--------------

SOLUTION/DROPS; OPHTHALMIC

CILOXAN

AT + ALCON PHARMS LTD	<u>EQ 0.3% BASE</u>	<u>N019992 001</u>	Dec 31, 1990
	<u>CIPROFLOXACIN HYDROCHLORIDE</u>		
AT AKORN INC	<u>EQ 0.3% BASE</u>	<u>A076555 001</u>	Dec 11, 2008
AT APOTEX	<u>EQ 0.3% BASE</u>	<u>A075928 001</u>	Jun 09, 2004
AT BAUSCH AND LOMB	<u>EQ 0.3% BASE</u>	<u>A076754 001</u>	Jun 09, 2004
AT FDC LTD	<u>EQ 0.3% BASE</u>	<u>A077568 001</u>	Jun 30, 2008
AT HITECH PHARMA	<u>EQ 0.3% BASE</u>	<u>A076673 001</u>	Jan 21, 2005
AT NEXUS PHARMS	<u>EQ 0.3% BASE</u>	<u>A077689 001</u>	Dec 13, 2006
AT PHARMAFORCE	<u>EQ 0.3% BASE</u>	<u>A078598 001</u>	Jan 16, 2008

SOLUTION/DROPS; OTIC

+ WRASER PHARMS	EQ 0.2% BASE	N021918 001	May 01, 2009
-----------------	--------------	-------------	--------------

TABLET; ORAL

CIPRO

AB BAYER HLTHCARE	<u>EQ 100MG BASE</u>	<u>N019537 001</u>	Apr 08, 1996
AB	<u>EQ 250MG BASE</u>	<u>N019537 002</u>	Oct 22, 1987
AB +	<u>EQ 500MG BASE</u>	<u>N019537 003</u>	Oct 22, 1987
AB	<u>EQ 750MG BASE</u>	<u>N019537 004</u>	Oct 22, 1987
	<u>CIPROFLOXACIN HYDROCHLORIDE</u>		
AB APOTEX	<u>EQ 250MG BASE</u>	<u>A076896 001</u>	Nov 04, 2004
AB	<u>EQ 500MG BASE</u>	<u>A076896 002</u>	Nov 04, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 99 (of 424)

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

AB	APOTEX	<u>EQ 750MG BASE</u>	<u>A076896</u> <u>003</u>	Nov 04, 2004
AB	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A077859</u> <u>001</u>	Apr 26, 2007
AB		<u>EQ 500MG BASE</u>	<u>A077859</u> <u>002</u>	Apr 26, 2007
AB		<u>EQ 750MG BASE</u>	<u>A077859</u> <u>003</u>	Apr 26, 2007
AB	CARLSBAD	<u>EQ 250MG BASE</u>	<u>A076126</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A076126</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A076126</u> <u>004</u>	Jun 09, 2004
AB	DR REDDYS LABS LTD	<u>EQ 100MG BASE</u>	<u>A075593</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 250MG BASE</u>	<u>A075593</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A075593</u> <u>004</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A075593</u> <u>001</u>	Jun 09, 2004
AB	HIKMA	<u>EQ 250MG BASE</u>	<u>A076558</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A076558</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A076558</u> <u>004</u>	Jun 09, 2004
AB	IVAX SUB TEVA PHARMS	<u>EQ 250MG BASE</u>	<u>A076089</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A076089</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A076089</u> <u>004</u>	Jun 09, 2004
AB	MYLAN	<u>EQ 100MG BASE</u>	<u>A075817</u> <u>001</u>	Jun 25, 2007
AB		<u>EQ 250MG BASE</u>	<u>A075685</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A075817</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A075685</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A075817</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A075685</u> <u>001</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A075817</u> <u>004</u>	Jun 09, 2004
AB	RANBAXY	<u>EQ 250MG BASE</u>	<u>A075747</u> <u>001</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A075747</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A075747</u> <u>003</u>	Jun 09, 2004
AB	TARO	<u>EQ 100MG BASE</u>	<u>A076912</u> <u>001</u>	Feb 18, 2005
AB		<u>EQ 250MG BASE</u>	<u>A076912</u> <u>002</u>	Oct 06, 2004
AB		<u>EQ 500MG BASE</u>	<u>A076912</u> <u>003</u>	Oct 06, 2004
AB		<u>EQ 750MG BASE</u>	<u>A076912</u> <u>004</u>	Oct 06, 2004
AB	UNIQUE PHARM LABS	<u>EQ 250MG BASE</u>	<u>A076639</u> <u>001</u>	Sep 10, 2004
AB		<u>EQ 500MG BASE</u>	<u>A076639</u> <u>002</u>	Sep 10, 2004
AB		<u>EQ 750MG BASE</u>	<u>A076639</u> <u>003</u>	Sep 10, 2004
AB	WATSON LABS	<u>EQ 100MG BASE</u>	<u>A076794</u> <u>001</u>	Feb 10, 2005
AB		<u>EQ 250MG BASE</u>	<u>A076794</u> <u>002</u>	Jun 09, 2004
AB		<u>EQ 500MG BASE</u>	<u>A076794</u> <u>003</u>	Jun 09, 2004
AB		<u>EQ 750MG BASE</u>	<u>A076794</u> <u>004</u>	Jun 09, 2004
<u>CIPROFLOXACIN HYDROCHLORIDE</u>				
BX	PLIVA	<u>EQ 100MG BASE</u>	<u>A076426</u> <u>001</u>	Jun 15, 2005
BX		<u>EQ 250MG BASE</u>	<u>A076426</u> <u>002</u>	Jun 15, 2005
BX		<u>EQ 500MG BASE</u>	<u>A076426</u> <u>003</u>	Jun 15, 2005
BX		<u>EQ 750MG BASE</u>	<u>A076426</u> <u>004</u>	Jun 15, 2005

CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS; OTIC

+ ALCON PHARMS LTD EQ 0.2% BASE;1%

N020805 001 Feb 10, 1998

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPRO XR

AB	+ BAYER HLTHCARE	<u>212.6MG;EQ 287.5MG BASE</u>	<u>N021473</u> <u>001</u>	Dec 13, 2002
AB	+ ANCHEN PHARMS	<u>425.2MG;EQ 574.9MG BASE</u>	<u>N021473</u> <u>002</u>	Aug 28, 2003
<u>CIPROFLOXACIN EXTENDED RELEASE</u>				
AB	ANCHEN PHARMS	<u>212.6MG;EQ 287.5MG BASE</u>	<u>A078166</u> <u>002</u>	Nov 27, 2007
AB		<u>425.2MG;EQ 574.9MG BASE</u>	<u>A078166</u> <u>001</u>	Nov 27, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 100 (of 424)

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPROFLOXACIN EXTENDED RELEASE

<u>AB</u>	DR REDDYS LABS LTD	<u>212.6MG;EQ 287.5MG BASE</u>	<u>A077902 001</u>	Oct 31, 2007
<u>AB</u>		<u>425.2MG;EQ 574.9MG BASE</u>	<u>A077701 001</u>	Mar 26, 2007
<u>AB</u>	MYLAN	<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
<u>AB</u>	WATSON LABS FLORIDA	<u>212.6MG;EQ 287.5MG BASE</u>	<u>A077417 001</u>	Nov 30, 2010
<u>AB</u>		<u>425.2MG;EQ 574.9MG BASE</u>	<u>A077809 001</u>	Nov 30, 2010

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS; OTIC

+ ALCON PHARMS LTD	0.3%;0.1%	N021537 001	Jul 18, 2003
--------------------	-----------	-------------	--------------

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

NIMBEX

+ ABBOTT	EQ 2MG BASE/ML	N020551 001	Dec 15, 1995
NIMBEX PRESERVATIVE FREE			
+ ABBOTT	EQ 2MG BASE/ML	N020551 003	Dec 15, 1995
+	EQ 10MG BASE/ML	N020551 002	Dec 15, 1995

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

<u>AP</u> + APP PHARMS	<u>1MG/ML</u>	<u>A074735 001</u>	Jul 16, 1999
<u>AP</u> BEDFORD	<u>1MG/ML</u>	<u>A075036 001</u>	Nov 07, 2000
<u>AP</u> PHARMACHEMIE	<u>1MG/ML</u>	<u>A074656 001</u>	May 16, 2000
<u>AP</u> TEVA PARENTERAL	<u>1MG/ML</u>	<u>A074814 001</u>	May 16, 2000

CITALOPRAM HYDROBROMIDE

CAPSULE; ORAL

CITALOPRAM HYDROBROMIDE

ALPHAPHARM	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

<u>AA</u> + FOREST LABS	<u>EQ 10MG BASE/5ML</u>	<u>N021046 001</u>	Dec 22, 1999
<u>CITALOPRAM HYDROBROMIDE</u>			
<u>AA</u> AUROBINDO PHARMA LTD	<u>EQ 10MG BASE/5ML</u>	<u>A077812 001</u>	Aug 28, 2006
<u>AA</u> ROXANE	<u>EQ 10MG BASE/5ML</u>	<u>A077043 001</u>	Dec 13, 2004
<u>AA</u> SILARX	<u>EQ 10MG BASE/5ML</u>	<u>A077629 001</u>	Jun 15, 2006

TABLET; ORAL

CELEXA

<u>AB</u> FOREST LABS	<u>EQ 10MG BASE</u>	<u>N020822 001</u>	Apr 27, 2000
	<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
<u>CITALOPRAM HYDROBROMIDE</u>			
<u>AB</u> ALPHAPHARM	<u>EQ 10MG BASE</u>	<u>A077037 001</u>	Nov 05, 2004
	<u>EQ 20MG BASE</u>	<u>A077037 002</u>	Nov 05, 2004
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A077037 003</u>	Nov 05, 2004
<u>AB</u> AMNEAL PHARMS NY	<u>EQ 10MG BASE</u>	<u>A077289 001</u>	Nov 30, 2006
	<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>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

PRESCRIPTION DRUG PRODUCT LIST

3 - 101 (of 424)

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

AB	AUROBINDO	EQ 10MG BASE	A077031 001	Oct 28, 2004
AB		EQ 20MG BASE	A077031 002	Oct 28, 2004
AB		EQ 40MG BASE	A077031 003	Oct 28, 2004
AB	CARACO	EQ 10MG BASE	A077032 001	Nov 12, 2004
AB		EQ 20MG BASE	A077032 002	Nov 12, 2004
AB		EQ 40MG BASE	A077032 003	Nov 12, 2004
AB	COREPHARMA	EQ 10MG BASE	A077036 001	Oct 28, 2004
AB		EQ 20MG BASE	A077036 002	Oct 28, 2004
AB		EQ 40MG BASE	A077036 003	Oct 28, 2004
AB	DR REDDYS LABS LTD	EQ 10MG BASE	A077038 001	Oct 28, 2004
AB		EQ 20MG BASE	A077038 002	Oct 28, 2004
AB		EQ 40MG BASE	A077038 003	Oct 28, 2004
AB	EPIC PHARMA	EQ 10MG BASE	A077045 003	Apr 29, 2005
AB		EQ 20MG BASE	A077045 002	Apr 29, 2005
AB		EQ 40MG BASE	A077045 001	Apr 29, 2005
AB	GLENMARK GENERICS	EQ 10MG BASE	A077654 001	Feb 27, 2009
AB		EQ 20MG BASE	A077654 002	Feb 27, 2009
AB		EQ 40MG BASE	A077654 003	Feb 27, 2009
AB	INVAGEN PHARMS	EQ 10MG BASE	A077534 001	Oct 03, 2006
AB		EQ 20MG BASE	A077534 002	Oct 03, 2006
AB		EQ 40MG BASE	A077534 003	Oct 03, 2006
AB	IVAX SUB TEVA PHARMS	EQ 10MG BASE	A077048 001	Nov 16, 2004
AB		EQ 20MG BASE	A077048 002	Nov 16, 2004
AB		EQ 40MG BASE	A077048 003	Nov 16, 2004
AB	MYLAN	EQ 10MG BASE	A077039 001	Feb 03, 2005
AB		EQ 10MG BASE	A077042 001	Nov 05, 2004
AB		EQ 20MG BASE	A077039 002	Feb 03, 2005
AB		EQ 20MG BASE	A077042 002	Nov 05, 2004
AB		EQ 40MG BASE	A077039 003	Feb 03, 2005
AB		EQ 40MG BASE	A077042 003	Nov 05, 2004
AB	NATCO PHARMA LTD	EQ 20MG BASE	A077141 002	Apr 10, 2008
AB		EQ 40MG BASE	A077141 001	Apr 10, 2008
AB	PLIVA	EQ 10MG BASE	A077232 001	Oct 31, 2005
AB		EQ 20MG BASE	A077232 002	Oct 31, 2005
AB		EQ 40MG BASE	A077232 003	Oct 31, 2005
AB	SANDOZ	EQ 10MG BASE	A077035 001	Oct 28, 2004
AB		EQ 20MG BASE	A077035 002	Oct 28, 2004
AB		EQ 40MG BASE	A077035 003	Oct 28, 2004
AB	TORRENT PHARMS	EQ 10MG BASE	A078216 001	Mar 27, 2007
AB		EQ 20MG BASE	A078216 002	Mar 27, 2007
AB		EQ 40MG BASE	A078216 003	Mar 27, 2007
AB	WATSON LABS	EQ 10MG BASE	A077034 001	Jun 30, 2005
AB		EQ 10MG BASE	A077044 001	Nov 05, 2004
AB		EQ 20MG BASE	A077034 002	Jun 30, 2005
AB		EQ 20MG BASE	A077044 002	Nov 05, 2004
AB		EQ 40MG BASE	A077034 003	Jun 30, 2005
AB		EQ 40MG BASE	A077044 003	Nov 05, 2004

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; UREA C-13

FOR SOLUTION, TABLET, FOR SOLUTION; ORAL

IDKIT:HP

+ EXALENZ BIOSCIENCE N/A,4GM;75MG,N/A N021314 001 Dec 17, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 102 (of 424)

CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE

<u>AP</u>	APP PHARMS	<u>1MG/ML</u>	<u>A076571</u>	<u>001</u>	Apr 22, 2004
<u>AP</u>	BEDFORD	<u>1MG/ML</u>	<u>A075405</u>	<u>001</u>	Feb 28, 2000
<u>AP</u>	ONCO THERAPIES LTD	<u>1MG/ML</u>	<u>A200510</u>	<u>001</u>	Oct 06, 2011
	<u>LEUSTATIN</u>				
<u>AP</u>	+ JANSSEN PHARMS	<u>1MG/ML</u>	<u>N020229</u>	<u>001</u>	Feb 26, 1993

CLARITHROMYCIN

FOR SUSPENSION; ORAL

BIAXIN

<u>AB</u>	ABBOTT	<u>125MG/5ML</u>	<u>N050698</u>	<u>001</u>	Dec 23, 1993
<u>AB</u>	+	<u>250MG/5ML</u>	<u>N050698</u>	<u>002</u>	Dec 23, 1993

CLARITHROMYCIN

<u>AB</u>	RANBAXY	<u>125MG/5ML</u>	<u>A065382</u>	<u>001</u>	Aug 30, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065382</u>	<u>002</u>	Aug 30, 2007
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065283</u>	<u>002</u>	Sep 04, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065283</u>	<u>003</u>	Sep 04, 2007

TABLET; ORAL

BIAXIN

<u>AB</u>	+ ABBOTT	<u>250MG</u>	<u>N050662</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>	+	<u>500MG</u>	<u>N050662</u>	<u>002</u>	Oct 31, 1991

CLARITHROMYCIN

<u>AB</u>	APOTEX CORP	<u>250MG</u>	<u>A065384</u>	<u>001</u>	Aug 20, 2007
<u>AB</u>		<u>500MG</u>	<u>A065384</u>	<u>002</u>	Aug 20, 2007
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A065195</u>	<u>001</u>	Mar 11, 2005
<u>AB</u>		<u>500MG</u>	<u>A065195</u>	<u>002</u>	Mar 11, 2005
<u>AB</u>	RANBAXY	<u>250MG</u>	<u>A065174</u>	<u>001</u>	Sep 24, 2004
<u>AB</u>		<u>500MG</u>	<u>A065174</u>	<u>002</u>	Sep 24, 2004
<u>AB</u>	ROXANE	<u>250MG</u>	<u>A065178</u>	<u>002</u>	May 25, 2004
<u>AB</u>		<u>500MG</u>	<u>A065178</u>	<u>001</u>	May 25, 2004
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065144</u>	<u>001</u>	Oct 18, 2005
<u>AB</u>		<u>500MG</u>	<u>A065136</u>	<u>001</u>	Aug 25, 2005
<u>AB</u>	TEVA	<u>250MG</u>	<u>A065155</u>	<u>001</u>	May 31, 2005
<u>AB</u>		<u>500MG</u>	<u>A065155</u>	<u>002</u>	May 31, 2005
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A065266</u>	<u>001</u>	May 31, 2006
<u>AB</u>		<u>500MG</u>	<u>A065266</u>	<u>002</u>	May 31, 2006

TABLET, EXTENDED RELEASE; ORAL

BIAXIN XL

<u>AB</u>	+ ABBOTT	<u>500MG</u>	<u>N050775</u>	<u>001</u>	Mar 03, 2000
	<u>CLARITHROMYCIN</u>				
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065250</u>	<u>001</u>	Aug 25, 2005

CLARITHROMYCIN

<u>AB</u>	TEVA	<u>500MG</u>	<u>A065154</u>	<u>001</u>	May 18, 2005
<u>AB</u>	WATSON LABS FLORIDA	<u>500MG</u>	<u>A065145</u>	<u>001</u>	Jun 24, 2004

CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TIMENTIN

+ GLAXOSMITHKLINE	EQ 1GM BASE/VIAL;EQ 30GM BASE/VIAL EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL		N050590	003	Aug 18, 1987
	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
	TIMENTIN IN PLASTIC CONTAINER				
+ GLAXOSMITHKLINE	EQ 100MG BASE/100ML;EQ 3GM BASE/100ML		N050658	001	Dec 15, 1989

PRESCRIPTION DRUG PRODUCT LIST

3 - 103 (of 424)

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

<u>AA</u>	NOVEX	<u>EQ 0.5MG BASE/5ML</u>	<u>A075703</u>	<u>001</u>	Nov 27, 2000
<u>AA</u>	SILARX	<u>EQ 0.5MG BASE/5ML</u>	<u>A074884</u>	<u>001</u>	Dec 17, 1997
<u>AA</u> +	TEVA	<u>EQ 0.5MG BASE/5ML</u>	<u>A073399</u>	<u>001</u>	Jun 30, 1994
<u>AA</u>	WOCKHARDT	<u>EQ 0.5MG BASE/5ML</u>	<u>A074863</u>	<u>001</u>	Mar 13, 1998

TABLET; ORAL

CLEMASTINE FUMARATE

<u>AB</u>	SANDOZ	<u>2.68MG</u>	<u>A073459</u>	<u>001</u>	Oct 31, 1993
<u>AB</u> +	TEVA	<u>2.68MG</u>	<u>A073283</u>	<u>001</u>	Jan 31, 1992

CLEVIDIPINE BUTYRATE

EMULSION; INTRAVENOUS

CLEVIPREX

+ MEDICINES CO	25MG/50ML (0.5MG/ML)	N022156	001	Aug 01, 2008
+	50MG/100ML (0.5MG/ML)	N022156	002	Aug 01, 2008

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN HYDROCHLORIDE

<u>AB</u>	PHARMACIA AND UPJOHN	<u>EQ 75MG BASE</u>	<u>N050162</u>	<u>001</u>	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N050162</u>	<u>002</u>	
<u>AB</u> +		<u>EQ 300MG BASE</u>	<u>N050162</u>	<u>003</u>	Apr 14, 1988

CLINDAMYCIN HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 150MG BASE</u>	<u>A065442</u>	<u>001</u>	Aug 26, 2009
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065442</u>	<u>002</u>	Aug 26, 2009
<u>AB</u>	COREPHARMA	<u>EQ 150MG BASE</u>	<u>A065194</u>	<u>001</u>	Mar 22, 2004
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065194</u>	<u>002</u>	Mar 22, 2004
<u>AB</u>	LANNETT	<u>EQ 75MG BASE</u>	<u>A065242</u>	<u>001</u>	Aug 12, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065242</u>	<u>002</u>	Aug 12, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065243</u>	<u>001</u>	Aug 12, 2005
<u>AB</u>	MATRIX LABS LTD	<u>EQ 75MG BASE</u>	<u>A091225</u>	<u>001</u>	May 31, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091225</u>	<u>002</u>	May 31, 2011
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A091225</u>	<u>003</u>	May 31, 2011
<u>AB</u>	RANBAXY	<u>EQ 150MG BASE</u>	<u>A065061</u>	<u>001</u>	Feb 02, 2001
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065061</u>	<u>002</u>	Feb 02, 2001
<u>AB</u>	TEVA	<u>EQ 150MG BASE</u>	<u>A063029</u>	<u>001</u>	Sep 20, 1989
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A063029</u>	<u>002</u>	Aug 05, 2005
<u>AB</u>	WATSON LABS	<u>EQ 150MG BASE</u>	<u>A063083</u>	<u>001</u>	Jul 31, 1991
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A063083</u>	<u>002</u>	Mar 18, 2003
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 75MG BASE</u>	<u>A065217</u>	<u>001</u>	Jan 31, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065217</u>	<u>002</u>	Jan 31, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A065217</u>	<u>003</u>	Jan 31, 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>AA</u>	CLINDAMYCIN PALMITATE HYDROCHLORIDE				
<u>AA</u>	PADDOCK LABS	<u>EQ 75MG BASE/5ML</u>	<u>A090902</u>	<u>001</u>	Jul 07, 2010

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AT</u>	PERRIGO UK FINCO	<u>1%</u>	<u>A090785</u>	<u>001</u>	Mar 31, 2010
<u>AT</u>	EVOCLIN				
<u>AT</u> +	STIEFEL LABS INC	<u>1%</u>	<u>N050801</u>	<u>001</u>	Oct 22, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 104 (of 424)

CLINDAMYCIN PHOSPHATE

CREAM; VAGINAL			
<u>CLEOCIN</u>			
AB + PHARMACIA AND UPJOHN	<u>EQ 2% BASE</u>	<u>N050680 002</u>	Mar 02, 1998
<u>CLINDAMYCIN PHOSPHATE</u>			
AB NYCOMED US	<u>EQ 2% BASE</u>	<u>A065139 001</u>	Dec 27, 2004
CLINDESSE			
+ KV PHARM	<u>EQ 2% BASE</u>	<u>N050793 001</u>	Nov 30, 2004
GEL; TOPICAL			
<u>CLEOCIN T</u>			
AB + PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050615 001</u>	Jan 07, 1987
<u>CLINDAMYCIN PHOSPHATE</u>			
AB ALTANA	<u>EQ 1% BASE</u>	<u>A064160 001</u>	Jan 28, 2000
CLINDAGEL			
BT + GALDERMA LABS LP	<u>EQ 1% BASE</u>	<u>N050782 001</u>	Nov 27, 2000
INJECTABLE; INJECTION			
<u>CLEOCIN PHOSPHATE</u>			
AP PHARMACIA AND UPJOHN	<u>EQ 150MG BASE/ML</u>	<u>A062803 001</u>	Oct 16, 1987
AP +	<u>EQ 150MG BASE/ML</u>	<u>N050441 001</u>	
<u>CLINDAMYCIN PHOSPHATE</u>			
AP APP PHARMS	<u>EQ 150MG BASE/ML</u>	<u>A065346 001</u>	Mar 29, 2007
AP	<u>EQ 150MG BASE/ML</u>	<u>A065347 001</u>	May 09, 2007
AP BAXTER HLTHCARE	<u>EQ 150MG BASE/ML</u>	<u>A062889 001</u>	Apr 25, 1988
AP BEDFORD	<u>EQ 150MG BASE/ML</u>	<u>A065206 001</u>	Sep 24, 2004
AP HOSPIRA	<u>EQ 150MG BASE/ML</u>	<u>A062800 001</u>	Jul 24, 1987
AP	<u>EQ 150MG BASE/ML</u>	<u>A062801 001</u>	Jul 24, 1987
AP	<u>EQ 150MG BASE/ML</u>	<u>A062943 001</u>	Sep 29, 1988
AP SAGENT STRIDES	<u>EQ 150MG BASE/ML</u>	<u>A090108 001</u>	Sep 30, 2011
AP	<u>EQ 150MG BASE/ML</u>	<u>A090109 001</u>	Sep 30, 2011
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER			
+ PHARMACIA AND UPJOHN	<u>EQ 6MG BASE/ML</u>	<u>N050639 001</u>	Aug 30, 1989
	<u>EQ 12MG BASE/ML</u>	<u>N050639 002</u>	Aug 30, 1989
	<u>EQ 18MG BASE/ML</u>	<u>N050639 003</u>	Apr 10, 1991
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%			
+ ABRAXIS PHARM	<u>EQ 900MG BASE/100ML</u>	<u>N050635 001</u>	Dec 22, 1989
LOTION; TOPICAL			
<u>CLEOCIN T</u>			
AB + PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050600 001</u>	May 31, 1989
<u>CLINDAMYCIN PHOSPHATE</u>			
AB ALTANA	<u>EQ 1% BASE</u>	<u>A065067 001</u>	Jan 31, 2002
SOLUTION; TOPICAL			
<u>CLEOCIN T</u>			
AT + PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050537 001</u>	
<u>CLINDA-DERM</u>			
AT PADDOCK LLC	<u>EQ 1% BASE</u>	<u>A063329 001</u>	Sep 30, 1992
<u>CLINDAMYCIN PHOSPHATE</u>			
AT ALTANA	<u>EQ 1% BASE</u>	<u>A065254 001</u>	Feb 14, 2006
AT FOUGERA	<u>EQ 1% BASE</u>	<u>A064159 001</u>	Jun 05, 1997
AT PERRIGO NEW YORK	<u>EQ 1% BASE</u>	<u>A064050 001</u>	Nov 30, 1995
AT TARO PHARM INDs	<u>EQ 1% BASE</u>	<u>A065184 001</u>	Mar 31, 2004
AT WOCKHARDT	<u>EQ 1% BASE</u>	<u>A063304 001</u>	Jul 15, 1997
SUPPOSITORY; VAGINAL			
CLEOCIN			
+ PHARMACIA AND UPJOHN	100MG	<u>N050767 001</u>	Aug 13, 1999
SWAB; TOPICAL			
<u>CLEOCIN</u>			
AT PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050537 002</u>	Feb 22, 1994
<u>CLINDAMYCIN PHOSPHATE</u>			
AT PERRIGO NEW YORK	<u>EQ 1% BASE</u>	<u>A065049 001</u>	May 25, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 105 (of 424)

CLINDAMYCIN PHOSPHATE

SWAB; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AT</u>	VERSAPHARM	<u>EQ 1% BASE</u>	<u>A065513 001</u>	Jun 17, 2010
<u>AT</u>	<u>CLINDETS</u>			

PERRIGOEQ 1% BASEA064136 001

Sep 30, 1996

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL

VELTIN

BX	STIEFEL GSK	1.2%;0.025%	N050803 001	Jul 16, 2010
BX	ZIANA			

+ MEDICIS1.2%;0.025%N050802 001

Nov 07, 2006

CLOBAZAM

TABLET; ORAL

ONFI

LUNDBECK INC

5MG

N202067 001

Oct 21, 2011

10MG

N202067 002

Oct 21, 2011

+

20MG

N202067 003

Oct 21, 2011

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	PERRIGO ISRAEL	<u>0.05%</u>	<u>A077763 001</u>	Mar 10, 2008
-----------	----------------	--------------	--------------------	--------------

OLUX

<u>AB</u>	+ STIEFEL LABS INC	<u>0.05%</u>	<u>N021142 001</u>	May 26, 2000
	OLUX E			

+ STIEFEL LABS INC0.05%N022013 001

Jan 12, 2007

CREAM; TOPICAL

CLOBETASOL PROPIONATE

<u>AB1</u>	FOUGERA	<u>0.05%</u>	<u>A074392 001</u>	Sep 30, 1996
------------	---------	--------------	--------------------	--------------

<u>AB1</u>	TARO	<u>0.05%</u>	<u>A074249 001</u>	Jul 08, 1996
------------	------	--------------	--------------------	--------------

<u>AB1</u>	TEVA PHARMS	<u>0.05%</u>	<u>A074087 001</u>	Feb 16, 1994
------------	-------------	--------------	--------------------	--------------

CORMAX

<u>AB1</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A074220 001</u>	May 16, 1997
------------	----------------	--------------	--------------------	--------------

TEMOVATE

<u>AB1</u>	+ NYCOMED US	<u>0.05%</u>	<u>N019322 001</u>	Dec 27, 1985
------------	--------------	--------------	--------------------	--------------

CLOBETASOL PROPIONATE (EMOLlient)

<u>AB2</u>	ALTANA	<u>0.05%</u>	<u>A075430 001</u>	May 26, 1999
------------	--------	--------------	--------------------	--------------

<u>AB2</u>	TARO	<u>0.05%</u>	<u>A075633 001</u>	May 17, 2000
------------	------	--------------	--------------------	--------------

EMBELINE E

<u>AB2</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A075325 001</u>	Dec 24, 1998
------------	----------------	--------------	--------------------	--------------

TEMOVATE E

<u>AB2</u>	+ FOUGERA PHARMS	<u>0.05%</u>	<u>N020340 001</u>	Jun 17, 1994
------------	------------------	--------------	--------------------	--------------

GEL; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A075368 001</u>	Feb 15, 2000
-----------	--------	--------------	--------------------	--------------

<u>AB</u>	PERRIGO	<u>0.05%</u>	<u>A075027 001</u>	Oct 31, 1997
-----------	---------	--------------	--------------------	--------------

TARO

<u>AB</u>	EMBELINE	<u>0.05%</u>	<u>A075279 001</u>	May 28, 1999
-----------	----------	--------------	--------------------	--------------

HI TECH PHARMA

<u>AB</u>	TEMOVATE	<u>0.05%</u>	<u>A076141 001</u>	Apr 12, 2002
-----------	----------	--------------	--------------------	--------------

NYCOMED US

<u>AB</u>	+ NYCOMED US	<u>0.05%</u>	<u>N020337 001</u>	Apr 29, 1994
-----------	--------------	--------------	--------------------	--------------

LOTION; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A078223 001</u>	Dec 04, 2008
-----------	----------------------	--------------	--------------------	--------------

CLOBEX

<u>AB</u>	+ GALDERMA LABS LP	<u>0.05%</u>	<u>N021535 001</u>	Jul 24, 2003
-----------	--------------------	--------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 106 (of 424)

CLOBETASOL PROPIONATE

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	NYCOMED US	<u>0.05%</u>	<u>A074407</u>	<u>001</u>	Feb 23, 1996
<u>AB</u>	TARO	<u>0.05%</u>	<u>A074248</u>	<u>001</u>	Jul 12, 1996
<u>AB</u>	TEVA PHARMS	<u>0.05%</u>	<u>A074089</u>	<u>001</u>	Feb 16, 1994
<u>EMBELINE</u>					
<u>AB</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A074221</u>	<u>001</u>	Mar 31, 1995
<u>TEMOVATE</u>					
<u>AB</u>	+ NYCOMED US	<u>0.05%</u>	<u>N019323</u>	<u>001</u>	Dec 27, 1985

SHAMPOO; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A078854</u>	<u>001</u>	Jun 07, 2011
<u>CLOBEX</u>					
<u>AB</u>	+ GALDERMA LABS	<u>0.05%</u>	<u>N021644</u>	<u>001</u>	Feb 05, 2004

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

<u>AT</u>	NYCOMED US	<u>0.05%</u>	<u>A075391</u>	<u>001</u>	Feb 08, 1999
<u>AT</u>	TARO	<u>0.05%</u>	<u>A075224</u>	<u>001</u>	Nov 16, 1998
<u>AT</u>		<u>0.05%</u>	<u>A075363</u>	<u>001</u>	Dec 29, 2000
<u>AT</u>	TOLMAR	<u>0.05%</u>	<u>A076977</u>	<u>001</u>	Aug 05, 2005
<u>AT</u>	WOCKHARDT	<u>0.05%</u>	<u>A075205</u>	<u>001</u>	Nov 13, 1998

EMBELINE

<u>AT</u>	HI TECH PHARMA	<u>0.05%</u>	<u>A074222</u>	<u>001</u>	Dec 06, 1995
<u>TEMOVATE</u>					
<u>AT</u>	+ NYCOMED US	<u>0.05%</u>	<u>N019966</u>	<u>001</u>	Feb 22, 1990

SPRAY; TOPICAL

CLOBETASOL PROPIONATE

<u>AT</u>	PADDOCK LLC	<u>0.05%</u>	<u>A090898</u>	<u>001</u>	Jun 16, 2011
<u>CLOBEX</u>					
<u>AT</u>	+ GALDERMA LABS LP	<u>0.05%</u>	<u>N021835</u>	<u>001</u>	Oct 27, 2005

CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLODERM

+ PROMIUS PHARMA LLC	0.1%	N017765	001
----------------------	------	---------	-----

CLOFARABINE

INJECTABLE; IV (INFUSION)

CLOLAR

+ GENZYME	20MG/20ML (1MG/ML)	N021673	001	Dec 28, 2004
-----------	--------------------	---------	-----	--------------

CLOFAZIMINE

CAPSULE; ORAL

LAMPRENE

+ NOVARTIS	50MG	N019500	002	Dec 15, 1986
------------	------	---------	-----	--------------

CLOMIPHENE CITRATE

TABLET; ORAL

CLOMID

<u>AB</u>	+ SANOFI AVENTIS US	<u>50MG</u>	<u>N016131</u>	<u>002</u>
<u>CLOMIPHENE CITRATE</u>				

<u>AB</u>	PAR PHARM	<u>50MG</u>	<u>A075528</u>	<u>001</u>
<u>SEROPHEN</u>				

<u>AB</u>	EMD SERONO	<u>50MG</u>	<u>N018361</u>	<u>001</u>
<u>50MG</u>				

Mar 22, 1982

PRESCRIPTION DRUG PRODUCT LIST

3 - 107 (of 424)

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

ANAFRANIL

<u>AB</u>	MALLINCKRODT LLC	<u>25MG</u>	<u>N019906</u>	<u>001</u>	Dec 29, 1989
<u>AB</u>	<u>+</u>	<u>50MG</u>	<u>N019906</u>	<u>002</u>	Dec 29, 1989
<u>AB</u>		<u>75MG</u>	<u>N019906</u>	<u>003</u>	Dec 29, 1989
<u>CLOMIPRAMINE HYDROCHLORIDE</u>					
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A074947</u>	<u>001</u>	Apr 30, 1998
<u>AB</u>		<u>50MG</u>	<u>A074947</u>	<u>002</u>	Apr 30, 1998
<u>AB</u>		<u>75MG</u>	<u>A074947</u>	<u>003</u>	Apr 30, 1998
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A074364</u>	<u>001</u>	Mar 29, 1996
<u>AB</u>		<u>50MG</u>	<u>A074364</u>	<u>002</u>	Mar 29, 1996
<u>AB</u>		<u>75MG</u>	<u>A074364</u>	<u>003</u>	Mar 29, 1996
<u>AB</u>	TARO	<u>25MG</u>	<u>A074694</u>	<u>001</u>	Dec 31, 1996
<u>AB</u>		<u>50MG</u>	<u>A074694</u>	<u>002</u>	Dec 31, 1996
<u>AB</u>		<u>75MG</u>	<u>A074694</u>	<u>003</u>	Dec 31, 1996
<u>AB</u>	TEVA	<u>25MG</u>	<u>A074958</u>	<u>001</u>	Aug 26, 1997
<u>AB</u>		<u>50MG</u>	<u>A074958</u>	<u>002</u>	Aug 26, 1997
<u>AB</u>		<u>75MG</u>	<u>A074958</u>	<u>003</u>	Aug 26, 1997

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

<u>AB</u>	ACCORD HLTHCARE INC	<u>0.5MG</u>	<u>A077147</u>	<u>001</u>	May 02, 2005
<u>AB</u>		<u>1MG</u>	<u>A077147</u>	<u>002</u>	May 02, 2005
<u>AB</u>		<u>2MG</u>	<u>A077147</u>	<u>003</u>	May 02, 2005
<u>CLONAZEPAM</u>					
<u>AB</u>	ACTAVIS ELIZABETH	<u>0.5MG</u>	<u>A074869</u>	<u>001</u>	Oct 31, 1996
<u>AB</u>		<u>1MG</u>	<u>A074869</u>	<u>002</u>	Oct 31, 1996
<u>AB</u>		<u>2MG</u>	<u>A074869</u>	<u>003</u>	Oct 31, 1996
<u>AB</u>	ALPHAPHARM	<u>0.5MG</u>	<u>A074940</u>	<u>001</u>	Oct 30, 1997
<u>AB</u>		<u>1MG</u>	<u>A074940</u>	<u>002</u>	Oct 30, 1997
<u>AB</u>		<u>2MG</u>	<u>A074940</u>	<u>003</u>	Oct 30, 1997
<u>AB</u>	APOTEX	<u>0.5MG</u>	<u>A075468</u>	<u>001</u>	Oct 06, 2000
<u>AB</u>		<u>1MG</u>	<u>A075468</u>	<u>002</u>	Oct 06, 2000
<u>AB</u>		<u>2MG</u>	<u>A075468</u>	<u>003</u>	Oct 06, 2000
<u>AB</u>	CARACO	<u>0.5MG</u>	<u>A075423</u>	<u>001</u>	Apr 27, 2001
<u>AB</u>		<u>1MG</u>	<u>A075423</u>	<u>002</u>	Apr 27, 2001
<u>AB</u>		<u>2MG</u>	<u>A075423</u>	<u>003</u>	Apr 27, 2001
<u>AB</u>	MYLAN	<u>0.5MG</u>	<u>A075150</u>	<u>001</u>	Oct 05, 1998
<u>AB</u>		<u>1MG</u>	<u>A075150</u>	<u>002</u>	Oct 05, 1998
<u>AB</u>		<u>2MG</u>	<u>A075150</u>	<u>003</u>	Oct 05, 1998
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A074979</u>	<u>001</u>	Aug 29, 1997
<u>AB</u>		<u>1MG</u>	<u>A074979</u>	<u>002</u>	Aug 29, 1997
<u>AB</u>		<u>2MG</u>	<u>A074979</u>	<u>003</u>	Aug 29, 1997
<u>AB</u>	TEVA	<u>0.5MG</u>	<u>A074569</u>	<u>001</u>	Sep 10, 1996
<u>AB</u>		<u>1MG</u>	<u>A074569</u>	<u>002</u>	Sep 10, 1996
<u>AB</u>		<u>2MG</u>	<u>A074569</u>	<u>003</u>	Sep 10, 1996
<u>AB</u>	VINTAGE PHARMS	<u>0.5MG</u>	<u>A077856</u>	<u>001</u>	Jun 28, 2006
<u>AB</u>		<u>1MG</u>	<u>A077856</u>	<u>002</u>	Jun 28, 2006
<u>AB</u>		<u>2MG</u>	<u>A077856</u>	<u>003</u>	Jun 28, 2006
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>A074964</u>	<u>001</u>	Dec 30, 1997
<u>AB</u>		<u>1MG</u>	<u>A074964</u>	<u>002</u>	Dec 30, 1997
<u>AB</u>		<u>2MG</u>	<u>A074964</u>	<u>003</u>	Dec 30, 1997
<u>KLONOPIN</u>					
<u>AB</u>	ROCHE	<u>0.5MG</u>	<u>N017533</u>	<u>001</u>	
<u>AB</u>	<u>+</u>	<u>1MG</u>	<u>N017533</u>	<u>002</u>	
<u>AB</u>		<u>2MG</u>	<u>N017533</u>	<u>003</u>	

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	BARR	<u>0.125MG</u>	<u>A077194</u>	<u>001</u>	Aug 10, 2005
-----------	------	----------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 108 (of 424)

CLONAZEPAM

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	BARR	<u>0.25MG</u>	<u>A077194</u>	<u>002</u>	Aug 10, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077194</u>	<u>003</u>	Aug 10, 2005
<u>AB</u>		<u>1MG</u>	<u>A077194</u>	<u>004</u>	Aug 10, 2005
<u>AB</u>		<u>2MG</u>	<u>A077194</u>	<u>005</u>	Aug 10, 2005
<u>AB</u>	PAR PHARM	<u>0.125MG</u>	<u>A077171</u>	<u>001</u>	Aug 03, 2005
<u>AB</u>		<u>0.25MG</u>	<u>A077171</u>	<u>002</u>	Aug 03, 2005
<u>AB</u>		<u>0.5MG</u>	<u>A077171</u>	<u>003</u>	Aug 03, 2005
<u>AB</u>	+	<u>1MG</u>	<u>A077171</u>	<u>004</u>	Aug 03, 2005
<u>AB</u>		<u>2MG</u>	<u>A077171</u>	<u>005</u>	Aug 03, 2005

CLONIDINE

FILM, EXTENDED RELEASE; TRANSDERMAL

CATAPRES-TTS-1

<u>AB</u>	BOEHRINGER INGELHEIM	<u>0.1MG/24HR</u>	<u>N018891</u>	<u>001</u>	Oct 10, 1984
-----------	----------------------	-------------------	----------------	------------	--------------

CATAPRES-TTS-2

<u>AB</u>	BOEHRINGER INGELHEIM	<u>0.2MG/24HR</u>	<u>N018891</u>	<u>002</u>	Oct 10, 1984
-----------	----------------------	-------------------	----------------	------------	--------------

CATAPRES-TTS-3

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.3MG/24HR</u>	<u>N018891</u>	<u>003</u>	Oct 10, 1984
-----------	---	----------------------	-------------------	----------------	------------	--------------

CLONIDINE

<u>AB</u>	AVEVA	<u>0.1MG/24HR</u>	<u>A076157</u>	<u>001</u>	Aug 18, 2009
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076157</u>	<u>002</u>	Aug 18, 2009
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076157</u>	<u>003</u>	Aug 18, 2009
<u>AB</u>	BARR	<u>0.1MG/24HR</u>	<u>A079090</u>	<u>001</u>	Aug 20, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A079090</u>	<u>002</u>	Aug 20, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A079090</u>	<u>003</u>	Aug 20, 2010
<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.1MG/24HR</u>	<u>A076166</u>	<u>001</u>	Jul 16, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076166</u>	<u>002</u>	Jul 16, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076166</u>	<u>003</u>	Jul 16, 2010

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

<u>AP</u>	APP PHARMS	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200673</u>	<u>001</u>	Jul 08, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200673</u>	<u>002</u>	Jul 08, 2011
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200300</u>	<u>001</u>	Jan 26, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200300</u>	<u>002</u>	Jan 26, 2011
<u>AP</u>	LUITPOLD	<u>1 MG/10 ML (0.1 MG/ML)</u>	<u>A091104</u>	<u>001</u>	Oct 08, 2009
<u>AP</u>		<u>5 MG/10 ML (0.5 MG/ML)</u>	<u>A091104</u>	<u>002</u>	Oct 08, 2009
<u>AP</u>	DURACLON				
<u>AP</u>	BIONICHE PHARMA USA	<u>1 MG/10 ML (0.1 MG/ML)</u>	<u>N020615</u>	<u>001</u>	Oct 02, 1996
<u>AP</u>	+	<u>5 MG/10 ML (0.5 MG/ML)</u>	<u>N020615</u>	<u>002</u>	Apr 27, 1999

TABLET; ORAL

CATAPRES

<u>AB</u>	BOEHRINGER INGELHEIM	<u>0.1MG</u>	<u>N017407</u>	<u>001</u>	
<u>AB</u>		<u>0.2MG</u>	<u>N017407</u>	<u>002</u>	
<u>AB</u>	+	<u>0.3MG</u>	<u>N017407</u>	<u>003</u>	

CLONIDINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A070974</u>	<u>001</u>	Dec 16, 1986
<u>AB</u>		<u>0.2MG</u>	<u>A070975</u>	<u>001</u>	Dec 16, 1986
<u>AB</u>		<u>0.3MG</u>	<u>A070976</u>	<u>001</u>	Dec 16, 1986
<u>AB</u>	ALEMBIC PHARMS LTD	<u>0.1MG</u>	<u>A091368</u>	<u>001</u>	Dec 06, 2011
<u>AB</u>		<u>0.2MG</u>	<u>A091368</u>	<u>002</u>	Dec 06, 2011
<u>AB</u>		<u>0.3MG</u>	<u>A091368</u>	<u>003</u>	Dec 06, 2011
<u>AB</u>	DAVA PHARMS INC	<u>0.1MG</u>	<u>A071783</u>	<u>001</u>	Apr 05, 1988
<u>AB</u>		<u>0.2MG</u>	<u>A071784</u>	<u>001</u>	Apr 05, 1988
<u>AB</u>		<u>0.3MG</u>	<u>A071785</u>	<u>001</u>	Apr 05, 1988
<u>AB</u>	IMPAX LABS	<u>0.1MG</u>	<u>A078099</u>	<u>001</u>	Aug 27, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 109 (of 424)

CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE

<u>AB</u>	IMPAK LABS	<u>0.2MG</u>	<u>A078099</u> <u>002</u>	Aug 27, 2009
<u>AB</u>		<u>0.3MG</u>	<u>A078099</u> <u>003</u>	Aug 27, 2009
<u>AB</u>	MUTUAL PHARM	<u>0.1MG</u>	<u>A070925</u> <u>001</u>	Sep 04, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070924</u> <u>001</u>	Sep 04, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070923</u> <u>001</u>	Sep 04, 1987
<u>AB</u>	MYLAN	<u>0.1MG</u>	<u>A070317</u> <u>002</u>	Jul 09, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070317</u> <u>003</u>	Jun 09, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070317</u> <u>001</u>	Jun 09, 1987
<u>AB</u>	UNICHEM	<u>0.1MG</u>	<u>A078895</u> <u>001</u>	Aug 26, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078895</u> <u>002</u>	Aug 26, 2009
<u>AB</u>		<u>0.3MG</u>	<u>A078895</u> <u>003</u>	Aug 26, 2009
<u>AB</u>	VINTAGE	<u>0.1MG</u>	<u>A077901</u> <u>001</u>	Mar 09, 2007
<u>AB</u>		<u>0.2MG</u>	<u>A077901</u> <u>002</u>	Mar 09, 2007
<u>AB</u>		<u>0.3MG</u>	<u>A077901</u> <u>003</u>	Mar 09, 2007
<u>AB</u>	WATSON LABS	<u>0.3MG</u>	<u>A070963</u> <u>001</u>	Jul 08, 1986
TABLET, EXTENDED RELEASE; ORAL				
KAPVAY				
+ SHIONOGI INC		0.1MG	N022331 003	Sep 28, 2010

CLOPIDOGREL BISULFATE

TABLET; ORAL

PLAVIX

SANOFI AVENTIS US	EQ 75MG BASE	N020839 001	Nov 17, 1997
+	EQ 300MG BASE	N020839 002	Sep 20, 2007

CLORAZEPATE DIPOTASSIUM

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

<u>AB</u>	MYLAN	<u>3.75MG</u>	<u>A071858</u> <u>002</u>	Jul 17, 1987
<u>AB</u>		<u>7.5MG</u>	<u>A071858</u> <u>003</u>	Jul 17, 1987
<u>AB</u>		<u>15MG</u>	<u>A071858</u> <u>001</u>	Jul 17, 1987
<u>AB</u>	RANBAXY	<u>3.75MG</u>	<u>A076911</u> <u>001</u>	Sep 29, 2004
<u>AB</u>		<u>7.5MG</u>	<u>A076911</u> <u>002</u>	Sep 29, 2004
<u>AB</u>		<u>15MG</u>	<u>A076911</u> <u>003</u>	Sep 29, 2004
<u>AB</u>	TARO	<u>3.75MG</u>	<u>A075731</u> <u>003</u>	Apr 27, 2000
<u>AB</u>		<u>7.5MG</u>	<u>A075731</u> <u>002</u>	Apr 27, 2000
<u>AB</u>		<u>15MG</u>	<u>A075731</u> <u>001</u>	Apr 27, 2000
<u>AB</u>	WATSON LABS	<u>3.75MG</u>	<u>A071852</u> <u>001</u>	Feb 09, 1988
<u>AB</u>		<u>7.5MG</u>	<u>A071853</u> <u>001</u>	Feb 09, 1988
<u>AB</u>		<u>15MG</u>	<u>A071854</u> <u>001</u>	Feb 09, 1988
GEN-XENE				
<u>AB</u>	ALRA	<u>3.75MG</u>	<u>A071787</u> <u>001</u>	Apr 26, 1988
<u>AB</u>		<u>7.5MG</u>	<u>A071788</u> <u>001</u>	Apr 26, 1988
<u>AB</u>		<u>15MG</u>	<u>A071789</u> <u>001</u>	Apr 26, 1988
TRANXENE				
<u>AB</u>	LUNDBECK INC	<u>3.75MG</u>	<u>N017105</u> <u>006</u>	
<u>AB</u>		<u>7.5MG</u>	<u>N017105</u> <u>007</u>	
<u>AB</u>	+	<u>15MG</u>	<u>N017105</u> <u>008</u>	

CLOTTRIMAZOLE

CREAM; TOPICAL

CLOTTRIMAZOLE

<u>AB</u>	GLENMARK PHARMS	<u>1%</u>	<u>A090219</u> <u>001</u>	Aug 03, 2010
<u>AB</u>	NYCOMED US	<u>1%</u>	<u>A078338</u> <u>001</u>	Sep 02, 2008
<u>AB</u>	+	<u>1%</u>	<u>A072640</u> <u>001</u>	Aug 31, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 110 (of 424)

CLOTRIMAZOLE

SOLUTION; TOPICAL

CLOTRIMAZOLE

<u>AT</u>	TARO	<u>1%</u>	<u>A074580</u>	<u>001</u>	Jul 29, 1996
<u>AT</u>	TEVA	<u>1%</u>	<u>A073306</u>	<u>001</u>	Feb 28, 1995

MYCELEX

<u>AT</u>	+ BAYER HLTHCARE	<u>1%</u>	<u>N018181</u>	<u>001</u>
-----------	------------------	-----------	----------------	------------

TROCHE / LOZENGE; ORAL

CLOTRIMAZOLE

<u>AB</u>	PADDICK LLC	<u>10MG</u>	<u>A076763</u>	<u>001</u>	Oct 28, 2005
<u>AB</u>	ROXANE	<u>10MG</u>	<u>A076387</u>	<u>001</u>	Jul 29, 2004

MYCELEX

<u>AB</u>	+ BAYER HLTHCARE	<u>10MG</u>	<u>N018713</u>	<u>001</u>	Jun 17, 1983
-----------	------------------	-------------	----------------	------------	--------------

CLOZAPINE

TABLET; ORAL

CLOZAPINE

<u>AB</u>	CARACO	<u>25MG</u>	<u>A075713</u>	<u>001</u>	Nov 15, 2002
<u>AB</u>		<u>50MG</u>	<u>A075713</u>	<u>003</u>	Aug 19, 2005
<u>AB</u>		<u>100MG</u>	<u>A075713</u>	<u>002</u>	Nov 15, 2002
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A074949</u>	<u>001</u>	Nov 26, 1997
<u>AB</u>		<u>50MG</u>	<u>A074949</u>	<u>004</u>	Apr 25, 2005
<u>AB</u>		<u>50MG</u>	<u>A076809</u>	<u>003</u>	Dec 16, 2005
<u>AB</u>		<u>100MG</u>	<u>A074949</u>	<u>002</u>	Nov 26, 1997
<u>AB</u>		<u>100MG</u>	<u>A076809</u>	<u>002</u>	Dec 16, 2005
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A075417</u>	<u>001</u>	May 27, 1999
<u>AB</u>		<u>100MG</u>	<u>A075417</u>	<u>002</u>	May 27, 1999

CLOZARIL

<u>AB</u>	NOVARTIS	<u>25MG</u>	<u>N019758</u>	<u>001</u>	Sep 26, 1989
<u>AB</u>	+ CLOZAPINE	<u>100MG</u>	<u>N019758</u>	<u>002</u>	Sep 26, 1989

CLOZAPINE

IVAX SUB TEVA PHARMS	12.5MG	<u>A074949</u>	<u>003</u>	Jul 31, 2003
	200MG	<u>A076809</u>	<u>001</u>	Dec 16, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

FAZACLO ODT

AZUR PHARMA INTL	12.5MG	<u>N021590</u>	<u>004</u>	May 30, 2007
	25MG	<u>N021590</u>	<u>001</u>	Feb 10, 2004
+	100MG	<u>N021590</u>	<u>002</u>	Feb 10, 2004
	150MG	<u>N021590</u>	<u>005</u>	Jul 09, 2010
	200MG	<u>N021590</u>	<u>006</u>	Jul 09, 2010

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</u>	<u>001</u>	Dec 07, 2006

PROMETH VC W/ CODEINE

<u>AA</u>	+ ACTAVIS MID ATLANTIC	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>A088764</u>	<u>001</u>	Oct 31, 1984
-----------	------------------------	--------------------------------------	----------------	------------	--------------

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</u>	<u>001</u>	Oct 31, 1984
<u>AA</u>	HI TECH PHARMA	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A040151</u>	<u>001</u>	Aug 26, 1997
<u>AA</u>	PHARM ASSOC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A089647</u>	<u>001</u>	Dec 22, 1988
<u>AA</u>	SUN PHARM INDs INC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A090180</u>	<u>001</u>	Mar 17, 2010
<u>AA</u>	WOCKHARDT	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A088875</u>	<u>001</u>	Dec 17, 1984

PROMETHAZINE WITH CODEINE

<u>AA</u>	VINTAGE	<u>10MG/5ML; 6.25MG/5ML</u>	<u>A040650</u>	<u>001</u>	Jan 31, 2006
-----------	---------	-----------------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 111 (of 424)

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL
 TRIACIN-C
 + STI PHARMA LLC 10MG/5ML;30MG/5ML;1.25MG/5ML A088704 001 Mar 22, 1985

CODEINE SULFATE

SOLUTION; ORAL
 CODEINE SULFATE
 + ROXANE 30MG/5ML N202245 001 Jun 30, 2011

TABLET; ORAL
 CODEINE SULFATE
 ROXANE 15MG N022402 001 Jul 16, 2009
 30MG N022402 002 Jul 16, 2009
 + 60MG N022402 003 Jul 16, 2009

COLCHICINE

TABLET; ORAL
 COLCRYS
 + AR HOLDING CO INC 0.6MG N022352 001 Jul 29, 2009

COLCHICINE; PROBENECID

TABLET; ORAL
COL-PROBENECID
AB + WATSON LABS 0.5MG;500MG A084279 001
PROBENECID AND COLCHICINE
AB MIRROR PHARMS 0.5MG;500MG A040618 001 May 13, 2008

COLESEVELAM HYDROCHLORIDE

FOR SUSPENSION; ORAL
 WELCHOL
 DAIICHI SANKYO 1.875GM/PACKET N022362 001 Oct 02, 2009
 + 3.75GM/PACKET N022362 002 Oct 02, 2009

TABLET; ORAL
 WELCHOL
 + DAIICHI SANKYO 625MG N021176 001 May 26, 2000

COlestipol HYDROCHLORIDE

GRANULE; ORAL
COlestid
AB PHARMACIA AND UPJOHN 5GM/SCOOPFUL N017563 003 Sep 22, 1995
AB + 5GM/PACKET N017563 004 Sep 22, 1995

COlestipol HYDROCHLORIDE
AB IMPAX LABS 5GM/PACKET A077277 002 May 02, 2006
AB 5GM/SCOOPFUL A077277 001 May 02, 2006

FLAVORED COlestid
 PHARMACIA AND UPJOHN 5GM/PACKET N017563 001
 5GM/SCOOPFUL N017563 002

TABLET; ORAL
COlestid
AB + PHARMACIA AND 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 APP PHARMS EQ 150MG BASE/VIAL A065364 001 Apr 17, 2008
AP PADDOCK LLC EQ 150MG BASE/VIAL A065177 001 Mar 19, 2004
AP X GEN PHARMS EQ 150MG BASE/VIAL A064216 001 Feb 26, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 112 (of 424)

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

COLY-MYCIN MAP + JHP PHARMSEQ 150MG BASE/VIALN050108 002COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS; OTIC

COLY-MYCIN S

+ JHP PHARMS

EQ 3MG BASE/ML; 10MG/ML; EQ 3.3MG
BASE/ML; 0.5MG/ML

N050356 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; IV (INFUSION)

+ ASTELLAS

20MG/100ML (0.2MG/ML)

N021697 002 Oct 08, 2008

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

PARAGARD T 380A

+ DURAMED RES

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

+ QUESTCOR PHARMS

80 UNITS/ML

N008372 008

CORTISONE ACETATE

TABLET; ORAL

CORTISONE ACETATE

+ WEST WARD

25MG

A080776 002

COSYNTROPIN

INJECTABLE; INJECTION

CORTROSYNAP + AMPHASTAR PHARMS INC0.25MG/VIALN016750 001COSYNTROPINAP BIONICHE PHARMA0.25MG/VIALA090574 001 Dec 17, 2009

SOLUTION; INTRAVENOUS

COSYNTROPIN

+ SANDOZ

0.25MG/ML (0.25MG/ML)

N022028 001 Feb 21, 2008

CRIZOTINIB

CAPSULE; ORAL

XALKORI

+ PFIZER

200MG

N202570 001 Aug 26, 2011

+

250MG

N202570 002 Aug 26, 2011

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

+ KING PHARMS

0.8MG/INH

N018887 001 Dec 05, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 113 (of 424)

CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUMAA PACK PHARMS LLC 100MG/5ML A202583 001 Oct 27, 2011GASTROCROMAA + AZUR PHARMA 100MG/5ML N020479 001 Feb 29, 1996

SOLUTION; INHALATION

CROMOLYN SODIUMAN BAUSCH AND LOMB 10MG/ML A075585 001 Dec 21, 2000AN DEY 10MG/ML A074209 001 Apr 26, 1994AN NOVEX 10MG/ML A075333 001 Apr 30, 2002AN + TEVA PARENTERAL 10MG/ML A075271 001 Jan 18, 2000AN WATSON LABS 10MG/ML A076469 001 Jun 17, 2005AN WOCKHARDT 10MG/ML A075346 001 Oct 25, 1999

SOLUTION/DROPS; OPHTHALMIC

CROLOMAT BAUSCH AND LOMB 4% A074443 001 Jan 30, 1995CROMOLYN SODIUMAT AKORN 4% A074706 001 Apr 29, 1998AT ALCON 4% A075282 001 Jun 16, 1999AT NOVEX 4% A075615 001 Jan 26, 2001OPTICROMAT + ALLERGAN 4% N018155 001 Oct 03, 1984CROTAMITON

CREAM; TOPICAL

EURAX

+ RANBAXY 10% N006927 001

LOTION; TOPICAL

CROTANAT SUMMERS 10% A087204 001EURAXAT + RANBAXY 10% N009112 003CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE IN PLASTIC CONTAINER

+ HOSPIRA EQ 0.4MG COPPER/ML N018960 001 Jun 26, 1986CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMINAP + LUITPOLD 1MG/ML A080737 001VIBISONEAP + APP PHARMS 1MG/ML A080557 003

SPRAY, METERED; NASAL

NASCOBAL

+ PAR PHARM 0.5MG/SPRAY N021642 001 Jan 31, 2005CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

AMRIXAB ANESTA AG 15MG N021777 001 Feb 01, 2007AB + 30MG N021777 002 Feb 01, 2007CYCLOBENZAPRINE HYDROCHLORIDEAB MYLAN 15MG A090738 001 Apr 18, 2011AB 30MG A090738 002 Apr 18, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 114 (of 424)

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078643</u>	<u>001</u>	Sep 26, 2008
<u>AB</u>		<u>10MG</u>	<u>A078643</u>	<u>002</u>	Sep 26, 2008
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A090478</u>	<u>001</u>	Jul 23, 2010
<u>AB</u>		<u>10MG</u>	<u>A090478</u>	<u>002</u>	Jul 23, 2010
<u>AB</u>	JUBILANT CADISTA	<u>5MG</u>	<u>A077563</u>	<u>001</u>	Apr 19, 2006
<u>AB</u>		<u>10MG</u>	<u>A077563</u>	<u>002</u>	Apr 19, 2006
<u>AB</u>	KVK TECH	<u>5MG</u>	<u>A078048</u>	<u>001</u>	Feb 28, 2011
<u>AB</u>		<u>10MG</u>	<u>A078048</u>	<u>002</u>	Feb 28, 2011
<u>AB</u>	MUTUAL PHARM	<u>5MG</u>	<u>A073541</u>	<u>002</u>	Apr 06, 2006
<u>AB</u>		<u>10MG</u>	<u>A073541</u>	<u>001</u>	May 23, 1995
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A073144</u>	<u>002</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A073144</u>	<u>001</u>	May 30, 1991
<u>AB</u>	ORIT LABS LLC	<u>10MG</u>	<u>A078218</u>	<u>001</u>	Apr 18, 2008
<u>AB</u>	PLIVA	<u>10MG</u>	<u>A074421</u>	<u>001</u>	Sep 29, 1995
<u>AB</u>	PROSAM LABS	<u>5MG</u>	<u>A077291</u>	<u>001</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A077209</u>	<u>001</u>	Oct 04, 2005
<u>AB</u>	RANBAXY	<u>5MG</u>	<u>A078722</u>	<u>001</u>	May 12, 2008
<u>AB</u>		<u>7.5MG</u>	<u>A078722</u>	<u>002</u>	May 12, 2008
<u>AB</u>		<u>10MG</u>	<u>A078722</u>	<u>003</u>	May 12, 2008
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A072854</u>	<u>002</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A072854</u>	<u>001</u>	Nov 19, 1991
<u>AB</u>	VINTAGE PHARMS	<u>5MG</u>	<u>A077797</u>	<u>001</u>	Feb 28, 2007
<u>AB</u>		<u>10MG</u>	<u>A077797</u>	<u>002</u>	Feb 28, 2007
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>A071611</u>	<u>002</u>	Feb 03, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A071611</u>	<u>003</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A071611</u>	<u>001</u>	May 03, 1989
<u>FLEXERIL</u>					
<u>AB</u>	JANSSEN R AND D	<u>5MG</u>	<u>N017821</u>	<u>001</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N017821</u>	<u>002</u>	

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPENTOLATE

<u>AT</u>	AKORN	<u>1%</u>	<u>A040164</u>	<u>001</u>	Jan 13, 1997
<u>AT</u>		<u>2%</u>	<u>A040165</u>	<u>001</u>	Jan 13, 1997
<u>CYCLOGYL</u>					
<u>AT</u>	+ ALCON	<u>1%</u>	<u>A084110</u>	<u>001</u>	
<u>AT</u>	+	<u>2%</u>	<u>A084108</u>	<u>001</u>	
<u>PENTOLAIR</u>					
<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>A040075</u>	<u>001</u>	Apr 29, 1994
CYCLOGYL					
	+ ALCON	0.5%		<u>A084109</u>	001

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOMYDRIL

+ ALCON	0.2%;1%	<u>A084300</u>	<u>001</u>
---------	---------	----------------	------------

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

<u>AP</u>	+ BAXTER HLTHCARE	<u>500MG/VIAL</u>	<u>A040745</u>	<u>001</u>	May 21, 2008
<u>AP</u>	+	<u>1GM/VIAL</u>	<u>A040745</u>	<u>002</u>	May 21, 2008
<u>AP</u>	+	<u>2GM/VIAL</u>	<u>A040745</u>	<u>003</u>	May 21, 2008
<u>CYTOXAN</u>					
<u>AP</u>	BAXTER HLTHCARE	<u>500MG/VIAL</u>	<u>N012142</u>	<u>003</u>	
<u>AP</u>		<u>1GM/VIAL</u>	<u>N012142</u>	<u>004</u>	Aug 30, 1982

PRESCRIPTION DRUG PRODUCT LIST

3 - 115 (of 424)

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYTOXAN

<u>AP</u>	BAXTER HLTHCARE	<u>2GM/VIAL</u>	<u>N012142 005</u>	Aug 30, 1982
	TABLET; ORAL			
	CYCLOPHOSPHAMIDE			
	ROXANE	25MG	A040032 001	Aug 17, 1999

+

50MG

A040032 002 Aug 17, 1999

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>	PLIVA	<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>	ABBOTT	<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

SANDIMMUNE

BX	NOVARTIS	50MG	N050625 003	Nov 23, 1992
----	----------	------	-------------	--------------

EMULSION; OPHTHALMIC

RESTASIS

+

ALLERGAN

0.05%

N050790 001 Dec 23, 2002

INJECTABLE; INJECTION

CYCLOSPORINE

<u>AP</u>	BEDFORD	<u>50MG/ML</u>	<u>A065004 001</u>	Oct 29, 1999
<u>AP</u>	LUITPOLD	<u>50MG/ML</u>	<u>A065151 001</u>	Oct 07, 2003

SANDIMMUNE

<u>AP</u>	+	NOVARTIS	<u>50MG/ML</u>	<u>N050573 001</u>	Nov 14, 1983
-----------	---	----------	----------------	--------------------	--------------

SOLUTION; ORAL

CYCLOSPORINE

<u>AB1</u>	ABBOTT	<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

NOVEX

<u>AB1</u>	WATSON LABS	<u>100MG/ML</u>	<u>A065054 001</u>	Dec 18, 2001
<u>AB1</u>				

NEORAL

<u>AB1</u>	+	NOVARTIS	<u>100MG/ML</u>	<u>N050716 001</u>	Jul 14, 1995
------------	---	----------	-----------------	--------------------	--------------

CYCLOSPORINE

<u>AB2</u>	WOCKHARDT	<u>100MG/ML</u>	<u>A065133 001</u>	Sep 17, 2004
------------	-----------	-----------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 116 (of 424)

CYCLOSPORINE

SOLUTION; ORAL

SANDIMMUNEAB2 + NOVARTIS 100MG/ML N050574 001 Nov 14, 1983CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDEAA + LYNE 2MG/5ML A040668 001 Jun 28, 2006

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	COREPHARMA	<u>4MG</u>	<u>A040537 001</u>	Sep 30, 2003
<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>	STASON PHARMS	<u>4MG</u>	<u>A040644 001</u>	May 30, 2006

CYSTEAMINE BITARTRATE

CAPSULE; ORAL

CYSTAGON

MYLAN EQ 50MG BASE
+ EQ 150MG BASEN020392 001 Aug 15, 1994
N020392 002 Aug 15, 1994CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

<u>AP</u> + APP PHARMS	<u>100MG/ML</u>	<u>A076512 001</u>	Jan 15, 2004
<u>AP</u> BEDFORD	<u>100MG/VIAL</u>	<u>A071471 001</u>	Aug 02, 1989
<u>AP</u>	<u>500MG/VIAL</u>	<u>A071472 001</u>	Aug 02, 1989
<u>AP</u>	<u>1GM/VIAL</u>	<u>A074245 001</u>	Aug 31, 1994
<u>AP</u>	<u>2GM/VIAL</u>	<u>A074245 002</u>	Aug 31, 1994
<u>AP</u> + HOSPIRA	<u>20MG/ML</u>	<u>A071868 001</u>	Jun 04, 1990
<u>AP</u> +	<u>20MG/ML</u>	<u>A072168 001</u>	Aug 31, 1990
<u>AP</u> +	<u>20MG/ML</u>	<u>A072945 001</u>	Feb 28, 1994
<u>AP</u>	<u>100MG/ML</u>	<u>A075383 001</u>	Nov 22, 1999
<u>AP</u> ONCO THERAPIES LTD	<u>20MG/ML</u>	<u>A200914 001</u>	Dec 13, 2011
<u>AP</u>	<u>20MG/ML</u>	<u>A200915 001</u>	Dec 13, 2011
<u>AP</u>	<u>20MG/ML</u>	<u>A200916 001</u>	Dec 13, 2011
CYTOSAR-U			
<u>AP</u> TEVA PARENTERAL	<u>100MG/VIAL</u>	<u>A075206 001</u>	Dec 30, 1998
<u>AP</u> +	<u>500MG/VIAL</u>	<u>A075206 002</u>	Dec 30, 1998
<u>AP</u> +	<u>1GM/VIAL</u>	<u>A075206 004</u>	Dec 30, 1998
<u>AP</u> +	<u>2GM/VIAL</u>	<u>A075206 003</u>	Dec 30, 1998

INJECTABLE, LIPOSOMAL; INJECTION

DEPOCYT

+ PACIRA PHARMS INC 10MG/ML

N021041 001 Apr 01, 1999

DABIGATRAN ETEXILATE MESYLATE

CAPSULE; ORAL

PRADAXA

BOEHRINGER INGELHEIM 75MG

+ 150MG

N022512 001 Oct 18, 2010

N022512 002 Oct 18, 2010

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

<u>AP</u> APP PHARMS	<u>100MG/VIAL</u>	<u>A075371 001</u>	Aug 27, 1999
<u>AP</u>	<u>200MG/VIAL</u>	<u>A075371 002</u>	Aug 27, 1999
<u>AP</u> BEDFORD	<u>200MG/VIAL</u>	<u>A075812 001</u>	Jun 15, 2001
<u>AP</u>	<u>500MG/VIAL</u>	<u>A075812 002</u>	Oct 31, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 117 (of 424)

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

<u>AP</u>	HOSPIRA	<u>200MG/VIAL</u>	<u>A075940</u>	<u>001</u>	Oct 18, 2001
<u>AP</u>	TEVA PARENTERAL	<u>200MG/VIAL</u>	<u>A075259</u>	<u>002</u>	Aug 27, 1998
<u>AP</u>	<u>+</u>	<u>500MG/VIAL</u>	<u>A075259</u>	<u>001</u>	Sep 22, 2000
	<u>DTIC-DOME</u>				
<u>AP</u>	<u>+</u> BAYER HLTHCARE	<u>100MG/VIAL</u>	<u>N017575</u>	<u>001</u>	
<u>AP</u>	<u>+</u>	<u>200MG/VIAL</u>	<u>N017575</u>	<u>002</u>	

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

<u>AP</u>	<u>+</u> LUNDBECK INC	<u>0.5MG/VIAL</u>	<u>N050682</u>	<u>001</u>	
<u>AP</u>	<u>DACTINOMYCIN</u>				
	BEDFORD	<u>0.5MG/VIAL</u>	<u>A090304</u>	<u>001</u>	Mar 16, 2010

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

AMPYRA

+ ACORDA	10MG	N022250	001	Jan 22, 2010
----------	------	---------	-----	--------------

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; IV (INFUSION)

SYNERCID

+ KING PHARMS	350MG/VIAL;150MG/VIAL	N050748	001	Sep 21, 1999
---------------	-----------------------	---------	-----	--------------

DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

EISAI 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
	10,000IU/0.4ML (25,000IU/ML)	N020287	002	May 01, 2007
	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
	95,000IU/9.5ML (10,000IU/ML)	N020287	007	Apr 04, 2002

DANAZOL

CAPSULE; ORAL

DANAZOL

<u>AB</u>	BARR	<u>50MG</u>	<u>A074582</u>	<u>003</u>	May 29, 1998
<u>AB</u>		<u>100MG</u>	<u>A074582</u>	<u>002</u>	May 29, 1998
<u>AB</u>	<u>+</u>	<u>200MG</u>	<u>A074582</u>	<u>001</u>	Aug 09, 1996
<u>AB</u>	LANNETT	<u>50MG</u>	<u>A078214</u>	<u>001</u>	Apr 19, 2007
<u>AB</u>		<u>100MG</u>	<u>A078214</u>	<u>002</u>	Apr 19, 2007
<u>AB</u>		<u>200MG</u>	<u>A077246</u>	<u>001</u>	Sep 28, 2005

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUM

<u>AB</u>	JHP PHARMS	<u>25MG</u>	<u>N017443</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>N017443</u>	<u>003</u>	
<u>AB</u>	<u>+</u>	<u>100MG</u>	<u>N017443</u>	<u>002</u>	
	<u>DANTROLENE SODIUM</u>				
<u>AB</u>	IMPAK LABS	<u>25MG</u>	<u>A076856</u>	<u>001</u>	Mar 01, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 118 (of 424)

DANTROLENE SODIUM

CAPSULE; ORAL

DANTROLENE SODIUM

<u>AB</u>	IMPAK LABS	<u>50MG</u>	<u>A076856 002</u>	Mar 01, 2005
<u>AB</u>		<u>100MG</u>	<u>A076856 003</u>	Mar 01, 2005
INJECTABLE; INJECTION				
<u>DANTRIUM</u>				
<u>AP</u>	+ JHP PHARMS	<u>20MG/VIAL</u>	<u>N018264 001</u>	
<u>DANTROLENE SODIUM</u>				
<u>AP</u>	US WORLDMEDS	<u>20MG/VIAL</u>	<u>A078378 001</u>	Jul 24, 2007

DAPSONE

GEL; TOPICAL

ACZONE

+ ALLERGAN	5%	N021794 001	Jul 07, 2005
------------	----	-------------	--------------

TABLET; ORAL

DAPSONE

JACOBUS

+ 100MG

A086841 001

A086842 001

DAPTOMYCIN

INJECTABLE; IV (INFUSION)

+ CUBIST	500MG/VIAL	N021572 002	Sep 12, 2003
----------	------------	-------------	--------------

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

ENABLEX

WARNER CHILCOTT LLC	EQ 7.5MG BASE	N021513 001	Dec 22, 2004
+	EQ 15MG BASE	N021513 002	Dec 22, 2004

DARUNAVIR ETHANOLATE

SUSPENSION; ORAL

PREZISTA

+ TIBOTEC	EQ 100MG BASE/ML	N202895 001	Dec 16, 2011
TABLET; ORAL			
PREZISTA			
TIBOTEC	EQ 75MG BASE	N021976 004	Dec 18, 2008
	EQ 150MG BASE	N021976 005	Dec 18, 2008
	EQ 400MG BASE	N021976 003	Oct 21, 2008
+	EQ 600MG BASE	N021976 002	Feb 25, 2008

DASATINIB

TABLET; ORAL

SPRYCEL

BRISTOL MYERS SQUIBB	20MG	N021986 001	Jun 28, 2006
	50MG	N021986 002	Jun 28, 2006
	70MG	N021986 003	Jun 28, 2006
	80MG	N021986 005	Oct 28, 2010
	100MG	N021986 004	May 30, 2008
+	140MG	N021986 006	Oct 28, 2010

DAUNORUBICIN CITRATE

INJECTABLE, LIPOSOMAL; INJECTION

DAUNOXOME

+ GALEN (UK)	EQ 2MG BASE/ML	N050704 002	Apr 08, 1996
--------------	----------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 119 (of 424)

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

<u>AP</u>	+ BEDFORD	<u>EQ 20MG BASE/VIAL</u>	<u>A064103 001</u>	Feb 03, 1995
	<u>DAUNORUBICIN HYDROCHLORIDE</u>			
<u>AP</u>	APP PHARMS	<u>EQ 20MG BASE/VIAL</u>	<u>A065000 001</u>	May 25, 1999
<u>AP</u>	+ BEDFORD	<u>EQ 5MG BASE/ML</u>	<u>N050731 001</u>	Jan 30, 1998
<u>AP</u>	TEVA PARENTERAL	<u>EQ 5MG BASE/ML</u>	<u>A065035 001</u>	Jan 24, 2000
	<u>DAUNORUBICIN HYDROCHLORIDE</u>			
	APP PHARMS	<u>EQ 5MG BASE/VIAL</u>	<u>A065034 001</u>	Nov 20, 2001

DECITABINE

INJECTABLE; INTRAVENOUS

DACOGEN

+ EISAI INC	50MG/VIAL	<u>N021790 001</u>	May 02, 2006
-------------	-----------	--------------------	--------------

DEFERASIROX

TABLET, FOR SUSPENSION; ORAL

EXJADE

NOVARTIS	125MG	<u>N021882 001</u>	Nov 02, 2005
	250MG	<u>N021882 002</u>	Nov 02, 2005
+	500MG	<u>N021882 003</u>	Nov 02, 2005

DEFERIPRONE

TABLET; ORAL

FERRIPROX

+ APOPHARMA INC	500MG	<u>N021825 001</u>	Oct 14, 2011
-----------------	-------	--------------------	--------------

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

<u>AP</u>	APP PHARMS	<u>500MG/VIAL</u>	<u>A078718 001</u>	Sep 15, 2009
<u>AP</u>		<u>2GM/VIAL</u>	<u>A078718 002</u>	Sep 15, 2009
<u>AP</u>	BEDFORD	<u>500MG/VIAL</u>	<u>A078086 001</u>	May 30, 2007
<u>AP</u>		<u>2GM/VIAL</u>	<u>A078086 002</u>	May 30, 2007
<u>AP</u>	HOSPIRA	<u>500MG/VIAL</u>	<u>A076019 001</u>	Mar 17, 2004
<u>AP</u>		<u>2GM/VIAL</u>	<u>A076019 002</u>	Mar 17, 2004
<u>AP</u>	WATSON LABS	<u>500MG/VIAL</u>	<u>A076806 001</u>	Mar 31, 2006
<u>AP</u>		<u>2GM/VIAL</u>	<u>A076806 002</u>	Mar 31, 2006
	<u>DESFERAL</u>			
<u>AP</u>	+ NOVARTIS	<u>500MG/VIAL</u>	<u>N016267 001</u>	
<u>AP</u>	+	<u>2GM/VIAL</u>	<u>N016267 002</u>	May 25, 2000

DEGARELIX ACETATE

POWDER; SUBCUTANEOUS

FIRMAGON

FERRING	EQ 80MG BASE/VIAL	<u>N022201 001</u>	Dec 24, 2008
+	EQ 120MG BASE/VIAL	<u>N022201 002</u>	Dec 24, 2008

DELAVIRDINE MESYLATE

TABLET; ORAL

RESCRIPTOR

VIVI HLTHCARE	100MG	<u>N020705 001</u>	Apr 04, 1997
+	200MG	<u>N020705 002</u>	Jul 14, 1999

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DECLOMYCIN

<u>AB</u>	COREPHARMA	<u>150MG</u>	<u>N050261 002</u>
-----------	------------	--------------	--------------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 120 (of 424)

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DECLOMYCIN

<u>AB</u>	+ COREPHARMA	<u>300MG</u>	<u>N050261</u> <u>003</u>	
<u>DEMECLOCYCLINE HYDROCHLORIDE</u>				
<u>AB</u>	AMNEAL PHARM	<u>150MG</u>	<u>A065425</u> <u>001</u>	Feb 27, 2008
<u>AB</u>		<u>300MG</u>	<u>A065425</u> <u>002</u>	Feb 27, 2008
<u>AB</u>	BARR	<u>150MG</u>	<u>A065171</u> <u>001</u>	Dec 13, 2004
<u>AB</u>		<u>300MG</u>	<u>A065171</u> <u>002</u>	Dec 13, 2004
<u>AB</u>	IMPAK LABS	<u>150MG</u>	<u>A065094</u> <u>001</u>	Mar 22, 2004
<u>AB</u>		<u>300MG</u>	<u>A065094</u> <u>002</u>	Mar 22, 2004
<u>AB</u>	VERSAPHARM	<u>150MG</u>	<u>A065389</u> <u>001</u>	Dec 01, 2008
<u>AB</u>		<u>300MG</u>	<u>A065389</u> <u>002</u>	Dec 01, 2008

DESFLURANE

LIQUID; INHALATION

SUPRANE

+ BAXTER HLTHCARE CORP 99.9%

N020118 001 Sep 18, 1992

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS TOTOWA	<u>10MG</u>	<u>A074430</u> <u>001</u>	Feb 09, 1996
<u>AB</u>		<u>25MG</u>	<u>A071601</u> <u>001</u>	Jun 05, 1987
<u>AB</u>		<u>50MG</u>	<u>A071588</u> <u>001</u>	Jun 05, 1987
<u>AB</u>		<u>75MG</u>	<u>A071602</u> <u>001</u>	Oct 05, 1987
<u>AB</u>		<u>100MG</u>	<u>A071766</u> <u>001</u>	Oct 05, 1987
<u>AB</u>		<u>150MG</u>	<u>A074430</u> <u>002</u>	Feb 09, 1996
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A072099</u> <u>001</u>	May 24, 1988
<u>AB</u>		<u>25MG</u>	<u>A072100</u> <u>001</u>	May 24, 1988
<u>AB</u>		<u>50MG</u>	<u>A072101</u> <u>001</u>	May 24, 1988
<u>AB</u>		<u>75MG</u>	<u>A072102</u> <u>001</u>	Jun 20, 1988
<u>AB</u>		<u>100MG</u>	<u>A072103</u> <u>001</u>	Jun 20, 1988
<u>AB</u>		<u>150MG</u>	<u>A072104</u> <u>001</u>	Jun 20, 1988
<u>NORPRAMIN</u>				
<u>AB</u>	SANOFI AVENTIS US	<u>10MG</u>	<u>N014399</u> <u>007</u>	Feb 11, 1982
<u>AB</u>		<u>25MG</u>	<u>N014399</u> <u>001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>N014399</u> <u>003</u>	
<u>AB</u>		<u>75MG</u>	<u>N014399</u> <u>004</u>	
<u>AB</u>	+	<u>100MG</u>	<u>N014399</u> <u>005</u>	
<u>AB</u>		<u>150MG</u>	<u>N014399</u> <u>006</u>	

DESIRUDIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS

IPRIVASK

+ CANYON 15MG/VIAL

N021271 001 Apr 04, 2003

DESLORATADINE

SYRUP; ORAL

CLARINEX

+ SCHERING 0.5MG/ML

N021300 001 Sep 01, 2004

TABLET; ORAL

CLARINEX

<u>AB</u>	+ SCHERING PLOUGH	<u>5MG</u>	<u>N021165</u> <u>001</u>	Dec 21, 2001
<u>DESLORATADINE</u>				
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A078365</u> <u>001</u>	Mar 08, 2011
<u>AB</u>	LUPIN PHARMS	<u>5MG</u>	<u>A078352</u> <u>001</u>	Oct 25, 2010
<u>AB</u>	ORCHID HLTHCARE	<u>5MG</u>	<u>A078357</u> <u>001</u>	Feb 19, 2010
<u>AB</u>	PERRIGO R AND D	<u>5MG</u>	<u>A078361</u> <u>001</u>	Dec 22, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 121 (of 424)

DESLORATADINE

TABLET; ORAL

DESLORATADINE

<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A078364</u>	<u>001</u>	Dec 03, 2010
<u>AB</u>	SUN PHARM IND'S	<u>5MG</u>	<u>A078359</u>	<u>001</u>	Nov 16, 2010
TABLET, ORALLY DISINTEGRATING; ORAL					
	<u>CLARINEX</u>				
<u>AB</u>	SCHERING	<u>2.5MG</u>	<u>N021312</u>	<u>002</u>	Jul 14, 2005
<u>AB</u>	+	<u>5MG</u>	<u>N021312</u>	<u>001</u>	Jun 26, 2002
	<u>DESLORATADINE</u>				
<u>AB</u>	REDDYS	<u>2.5MG</u>	<u>A078367</u>	<u>001</u>	Jul 12, 2010
<u>AB</u>		<u>5MG</u>	<u>A078367</u>	<u>002</u>	Jul 12, 2010

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX D 24 HOUR

<u>AB</u>	+	SCHERING	<u>5MG; 240MG</u>	<u>N021605</u>	<u>001</u>	Mar 03, 2005
<u>DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR</u>						
<u>AB</u>	DR REDDYS LABS LTD		<u>5MG; 240MG</u>	<u>A078366</u>	<u>001</u>	Apr 26, 2011
	CLARINEX-D 12 HOUR					
	+	SCHERING	2.5MG; 120MG	N021313	001	Feb 01, 2006

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

<u>AP</u>	+	SANOFI AVENTIS US	<u>0.004MG/ML</u>	<u>N018938</u>	<u>001</u>	Mar 30, 1984
<u>DESMOPRESSIN ACETATE</u>						
<u>AP</u>	HOSPIRA		<u>0.004MG/ML</u>	<u>A075220</u>	<u>001</u>	Aug 28, 2000
<u>AP</u>	TEVA PARENTERAL		<u>0.004MG/ML</u>	<u>A074888</u>	<u>001</u>	Oct 15, 1997

SOLUTION; NASAL

DDAVP

+	SANOFI AVENTIS US	0.01%	N017922	001
---	-------------------	-------	---------	-----

SPRAY, METERED; NASAL

DDAVP (NEEDS NO REFRIGERATION)

<u>AB</u>	+	SANOFI AVENTIS US	<u>0.01MG/SPRAY</u>	<u>N017922</u>	<u>003</u>	Aug 07, 1996
<u>DESMOPRESSIN ACETATE</u>						
<u>AB</u>	+	BAUSCH AND LOMB	<u>0.01MG/SPRAY</u>	<u>A074830</u>	<u>001</u>	Jan 25, 1999
<u>DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)</u>						
<u>AB</u>	APOTEX INC		<u>0.01MG/SPRAY</u>	<u>A076703</u>	<u>001</u>	Jan 27, 2005
<u>MINIRIN</u>						
<u>AB</u>	+	FERRING	<u>0.01MG/SPRAY</u>	<u>N021333</u>	<u>001</u>	Sep 16, 2002
STIMATE (NEEDS NO REFRIGERATION)						
<u>AB</u>	+	CSL BEHRING	1.5MG/SPRAY	N020355	002	Oct 24, 2007

TABLET; ORAL

DDAVP

<u>AB</u>	SANOFI AVENTIS US	<u>0.1MG</u>	<u>N019955</u>	<u>001</u>	Sep 06, 1995
<u>AB</u>	+	<u>0.2MG</u>	<u>N019955</u>	<u>002</u>	Sep 06, 1995
<u>DESMOPRESSIN ACETATE</u>					
<u>AB</u>	APOTEX INC	<u>0.1MG</u>	<u>A077414</u>	<u>001</u>	Mar 07, 2006
<u>AB</u>		<u>0.2MG</u>	<u>A077414</u>	<u>002</u>	Mar 07, 2006
<u>AB</u>	TEVA PHARMS	<u>0.1MG</u>	<u>A077122</u>	<u>001</u>	Jan 25, 2006
<u>AB</u>		<u>0.2MG</u>	<u>A077122</u>	<u>002</u>	Jan 25, 2006
<u>AB</u>	WATSON LABS	<u>0.1MG</u>	<u>A076470</u>	<u>001</u>	Jul 01, 2005
<u>AB</u>		<u>0.2MG</u>	<u>A076470</u>	<u>002</u>	Jul 01, 2005
DESMOPRESSIN ACETATE					
	FERRING	0.1MG	N021795	001	May 08, 2008
	+	0.2MG	N021795	002	May 08, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 122 (of 424)

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-28

CYCLESSA

<u>AB</u>	+ ORGANON USA INC	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.</u> <u>025MG</u>	<u>N021090</u> <u>001</u>	Dec 20, 2000
<u>AB</u>	<u>DESOGEN</u>	<u>0.15MG;0.03MG</u>	<u>N020071</u> <u>002</u>	Dec 10, 1992
<u>AB</u>	<u>DESOGESTREL AND ETHINYL ESTRADIOL</u>	<u>0.15MG;0.03MG</u>	<u>A075256</u> <u>002</u>	Aug 12, 1999
<u>AB</u>	DURAMED PHARMS BARR	<u>0.15MG;0.03MG</u>	<u>A076916</u> <u>001</u>	Dec 29, 2008
<u>AB</u>	WATSON LABS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A076915</u> <u>001</u>	Jul 29, 2005
<u>AB</u>	<u>EMOQUETTE</u>	<u>0.15MG;0.03MG</u>	<u>A076675</u> <u>001</u>	Feb 25, 2011
<u>AB</u>	VINTAGE	<u>0.15MG;0.03MG</u>	<u>A075863</u> <u>001</u>	Apr 05, 2002
<u>AB</u>	<u>KARIVA</u>	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>N020713</u> <u>001</u>	Apr 22, 1998
<u>AB</u>	<u>MIRCETTE</u>	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>N020301</u> <u>002</u>	Dec 14, 1992
<u>AB</u>	+ TEVA WOMENS	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>A076455</u> <u>001</u>	Feb 24, 2004
<u>AB</u>	<u>ORTHO-CEPT</u>	<u>0.15MG;0.03MG</u>		
<u>AB</u>	+ JANSSEN PHARMS	<u>0.15MG;0.03MG</u>		
<u>AB</u>	<u>VELVET</u>	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.</u> <u>025MG</u>		

DESONIDEAEROSOL, FOAM; TOPICAL
VERDESO

	+ STIEFEL LABS INC	0.05%	N021978	001	Sep 19, 2006
--	--------------------	-------	---------	-----	--------------

CREAM; TOPICAL

DESONIDE

<u>AB</u>	+ PERRIGO NEW YORK	<u>0.05%</u>	<u>N017010</u>	<u>001</u>	
<u>AB</u>	TARO	<u>0.05%</u>	<u>A073548</u>	<u>001</u>	Jun 30, 1992

DESOWEN

<u>AB</u>	GALDERMA LABS LP	<u>0.05%</u>	<u>N019048</u>	<u>001</u>	Dec 14, 1984
-----------	------------------	--------------	----------------	------------	--------------

GEL; TOPICAL

DESONATE

	+ INTENDIS	0.05%	N021844	001	Oct 20, 2006
--	------------	-------	---------	-----	--------------

LOTION; TOPICAL

DESONIDE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A075860</u>	<u>001</u>	Mar 19, 2002
-----------	--------	--------------	----------------	------------	--------------

DESOWEN

<u>AB</u>	+ GALDERMA LABS LP	<u>0.05%</u>	<u>A072354</u>	<u>001</u>	Jan 24, 1992
-----------	--------------------	--------------	----------------	------------	--------------

OINTMENT; TOPICAL

DESONIDE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A075751</u>	<u>001</u>	Mar 12, 2001
-----------	--------	--------------	----------------	------------	--------------

<u>AB</u>	+ PERRIGO NEW YORK	<u>0.05%</u>	<u>N017426</u>	<u>001</u>	
-----------	--------------------	--------------	----------------	------------	--

TARO

<u>AB</u>	TARO	<u>0.05%</u>	<u>A074254</u>	<u>001</u>	Aug 03, 1994
-----------	------	--------------	----------------	------------	--------------

DESOWEN

<u>AB</u>	GALDERMA LABS LP	<u>0.05%</u>	<u>A071425</u>	<u>001</u>	Jun 15, 1988
-----------	------------------	--------------	----------------	------------	--------------

DESOXIMETASONE

CREAM; TOPICAL

DESOXIMETASONE

<u>AB</u>	NYCOMED US	<u>0.25%</u>	<u>A078369</u>	<u>001</u>	Jun 29, 2010
-----------	------------	--------------	----------------	------------	--------------

<u>AB</u>	PERRIGO NEW YORK	<u>0.25%</u>	<u>A076510</u>	<u>001</u>	Jul 01, 2003
-----------	------------------	--------------	----------------	------------	--------------

<u>AB</u>	+ TARO	<u>0.05%</u>	<u>A073210</u>	<u>001</u>	Nov 30, 1990
-----------	--------	--------------	----------------	------------	--------------

TOPICORT

<u>AB</u>	+ TARO PHARMS NORTH	<u>0.25%</u>	<u>N017856</u>	<u>001</u>	
-----------	---------------------	--------------	----------------	------------	--

TOPICORT LP

<u>AB</u>	TARO PHARMS NORTH	<u>0.05%</u>	<u>N018309</u>	<u>001</u>	
-----------	-------------------	--------------	----------------	------------	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 123 (of 424)

DESOXIMETASONE

GEL; TOPICAL

DESOXIMETASONE

<u>AB</u>	PERRIGO NEW YORK	<u>0.05%</u>	<u>A077552</u> <u>001</u>	Jan 09, 2006
<u>AB</u>	TARO	<u>0.05%</u>	<u>A074904</u> <u>001</u>	Jul 14, 1998
<u>AB</u>	VERSAPHARM	<u>0.05%</u>	<u>A090727</u> <u>001</u>	Mar 10, 2011
		<u>TOPICORT</u>		
<u>AB</u>	+ TARO PHARMS NORTH	<u>0.05%</u>	<u>N018586</u> <u>001</u>	Mar 29, 1982
		OINTMENT; TOPICAL		
		<u>DESOXIMETASONE</u>		
<u>AB</u>	TARO	<u>0.25%</u>	<u>A074286</u> <u>001</u>	Jun 07, 1996
		<u>TOPICORT</u>		
<u>AB</u>	+ TARO PHARMS NORTH	<u>0.25%</u>	<u>N018763</u> <u>001</u>	Sep 30, 1983
		TOPICORT		
	+ TARO PHARMS NORTH	0.05%	N018594 001	Jan 17, 1985

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

PRISTIQ

+ WYETH PHARMS INC	EQ 50MG BASE	N021992 001	Feb 29, 2008
+	EQ 100MG BASE	N021992 002	Feb 29, 2008

DEXAMETHASONE

CONCENTRATE; ORAL

DEXAMETHASONE INTENSOL

+ ROXANE	1MG/ML	A088252 001	Sep 01, 1983
----------	--------	-------------	--------------

ELIXIR; ORAL

DEXAMETHASONE

<u>AA</u>	LYNE	<u>0.5MG/5ML</u>	<u>A090891</u> <u>001</u>	Jul 12, 2011
<u>AA</u>	+ STI PHARMA LLC	<u>0.5MG/5ML</u>	<u>A084754</u> <u>001</u>	
<u>AA</u>	VINTAGE PHARMS	<u>0.5MG/5ML</u>	<u>A091188</u> <u>001</u>	May 11, 2011
<u>AA</u>	+ WOCKHARDT	<u>0.5MG/5ML</u>	<u>A088254</u> <u>001</u>	Jul 27, 1983

IMPLANT; INTRAVITREAL

OZURDEX

+ ALLERGAN	0.7MG	N022315 001	Jun 17, 2009
------------	-------	-------------	--------------

SOLUTION; ORAL

DEXAMETHASONE

+ ROXANE	0.5MG/5ML	A088248 001	Sep 01, 1983
----------	-----------	-------------	--------------

SUSPENSION/DROPS; OPHTHALMIC

+ ALCON	0.1%	N013422 001
---------	------	-------------

TABLET; ORAL

DEXAMETHASONE

<u>AB</u>	ECR	<u>1.5MG</u>	<u>A040700</u> <u>001</u>	Aug 15, 2008
<u>AB</u>	ROXANE	<u>1.5MG</u>	<u>A084610</u> <u>001</u>	
		DEXAMETHASONE		
BP	PAR PHARM	0.5MG	A088148 001	Apr 28, 1983
BP		0.75MG	A088160 001	Apr 28, 1983
BP		1.5MG	A088237 001	Apr 28, 1983
BP		4MG	A088238 001	Apr 28, 1983
BP	+	6MG	A088481 001	Nov 28, 1983
BP	ROXANE	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 124 (of 424)

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

<u>AP</u>	APP PHARMS	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A084916 001</u>	
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040491 001</u>	Apr 11, 2003
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040572 001</u>	Apr 22, 2005
<u>AP</u>	+ BAXTER HLTHCARE	<u>EQ 10MG PHOSPHATE/ML</u>	<u>A087702 001</u>	Sep 07, 1982
<u>AP</u>	+ LUITPOLD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A087440 001</u>	Jul 21, 1982
<u>AP</u>	PFIZER	<u>EQ 4MG PHOSPHATE/ML</u>	<u>A040803 001</u>	Aug 29, 2008
<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>A040802 001</u>	Aug 29, 2008

SOLUTION/DROPS; OPHTHALMIC, OTIC

DEXAMETHASONE SODIUM PHOSPHATE

<u>AT</u>	+ ALCON UNIVERSAL	<u>EQ 0.1% PHOSPHATE</u>	<u>A088771 001</u>	Jan 16, 1985
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.1% PHOSPHATE</u>	<u>A040069 001</u>	Jul 26, 1996

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

MAXITROL

<u>AT</u>	+ FALCON PHARMS	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N050065 002</u>	
		<u>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</u>		
<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064063 001</u>	Jul 25, 1994

ATFERA PHARMS 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A062938 001 Jul 31, 1989

SUSPENSION/DROPS; OPHTHALMIC

DEXASPORIN

<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064135 001</u>	Sep 13, 1995
		<u>MAXITROL</u>		
<u>AT</u>	ALCON	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062341 001</u>	May 22, 1984
<u>AT</u>	+ FALCON PHARMS	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N050023 002</u>	
		<u>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</u>		
<u>AT</u>	ALCON UNIVERSAL	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062721 001</u>	Nov 17, 1986

DEXAMETHASONE; TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBRADEX

+ ALCON 0.1%;0.3% N050616 001 Sep 28, 1988

SUSPENSION/DROPS; OPHTHALMIC

TOBRADEX

<u>AB</u>	+ ALCON	<u>0.1%;0.3%</u>	<u>N050592 001</u>	Aug 18, 1988
		<u>TOBRAMYCIN AND DEXAMETHASONE</u>		
<u>AB</u>	BAUSCH AND LOMB	<u>0.1%;0.3%</u>	<u>A064134 001</u>	Oct 27, 1999

TOBRADEX ST

+ ALCON PHARMS LTD 0.05%;0.3% N050818 001 Feb 13, 2009

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL

DEXCHLORPHENIRAMINE MALEATE

+ WOCKHARDT 2MG/5ML A088251 001 Mar 23, 1984

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE; ORAL

DEXTLANT

TAKEDA PHARMS 30MG N022287 001 Jan 30, 2009

+ 60MG N022287 002 Jan 30, 2009

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRECEDEX

+ HOSPIRA EQ 100MCG BASE/ML (EQ100MCG BASE/ML) N021038 001 Dec 17, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 125 (of 424)

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

FOCALIN XR

NOVARTIS	5MG	N021802 001	May 26, 2005
	10MG	N021802 002	May 26, 2005
	15MG	N021802 004	Aug 01, 2006
	20MG	N021802 003	May 26, 2005
	25MG	N021802 008	Apr 21, 2011
	30MG	N021802 005	Oct 23, 2009
	35MG	N021802 007	Apr 21, 2011
+	40MG	N021802 006	Aug 11, 2010

TABLET; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

AB	TEVA PHARMS	<u>2.5MG</u>	A077107 003	Jan 29, 2007
AB		<u>5MG</u>	A077107 001	Jan 29, 2007
AB		<u>10MG</u>	A077107 002	Jan 29, 2007
	<u>FOCALIN</u>			
AB	NOVARTIS	<u>2.5MG</u>	N021278 001	Nov 13, 2001
AB		<u>5MG</u>	N021278 002	Nov 13, 2001
AB	+	<u>10MG</u>	N021278 003	Nov 13, 2001

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXRAZOXANE HYDROCHLORIDE

AP	BEDFORD	<u>EQ 250MG BASE/VIAL</u>	A076068 001	Sep 28, 2004
AP		<u>EQ 500MG BASE/VIAL</u>	A076068 002	Sep 28, 2004
AP	MYLAN INSTITUTIONAL	<u>EQ 250MG BASE/VIAL</u>	A200752 001	Oct 19, 2011
AP		<u>EQ 500MG BASE/VIAL</u>	A200752 002	Oct 19, 2011
	<u>ZINECARD</u>			
AP	+ PHARMACIA AND UPJOHN	<u>EQ 250MG BASE/VIAL</u>	N020212 001	May 26, 1995
AP		<u>EQ 500MG BASE/VIAL</u>	N020212 002	May 26, 1995
	TOTECT			
+	TOPOTARGET	EQ 500MG BASE/VIAL	N022025 001	Sep 06, 2007

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE

AB	COREPHARMA	<u>5MG</u>	N017078 001	
AB		<u>10MG</u>	N017078 002	
AB	+	<u>15MG</u>	N017078 003	
	<u>DEXTROAMPHETAMINE SULFATE</u>			
AB	BARR	<u>5MG</u>	A076137 001	Jan 18, 2002
AB		<u>10MG</u>	A076137 002	Jan 18, 2002
AB		<u>15MG</u>	A076137 003	Jan 18, 2002
AB	MALLINCKRODT	<u>5MG</u>	A076353 001	May 06, 2003
AB		<u>10MG</u>	A076353 002	May 06, 2003
AB		<u>15MG</u>	A076353 003	May 06, 2003

SOLUTION; ORAL

DEXTROAMPHETAMINE SULFATE

+ OUTLOOK PHARMS 5MG/5ML

A040776 001 Jan 29, 2008

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

AA	BARR	<u>5MG</u>	A040361 001	Jan 31, 2001
AA	+	<u>10MG</u>	A040361 002	Jan 31, 2001
AA	MALLINCKRODT	<u>5MG</u>	A040436 001	Jan 29, 2002
AA		<u>10MG</u>	A040436 002	Jan 29, 2002
AA	MIKART	<u>5MG</u>	A090533 002	Oct 25, 2011
AA		<u>10MG</u>	A090533 004	Oct 25, 2011
AA	NESHER PHARMS	<u>5MG</u>	A040365 001	Oct 31, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 126 (of 424)

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	NESHER PHARMS	<u>10MG</u>	<u>A040367 001</u>	Oct 31, 2002
	DEXTROAMPHETAMINE SULFATE			
	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

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH W/ DEXTROMETHORPHAN

<u>AA</u>	+ ACTAVIS MID ATLANTIC	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A088762 001</u>	Oct 31, 1984
<u>AA</u>	<u>PROMETHAZINE DM</u>	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A040649 001</u>	Feb 14, 2006
<u>AA</u>	VINTAGE	<u>15MG/5ML; 6.25MG/5ML</u>		
<u>AA</u>	<u>PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE</u>			
<u>AA</u>	AMNEAL PHARMS	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A090575 001</u>	Feb 08, 2011
<u>AA</u>	HI TECH PHARMA	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A040027 001</u>	Jul 31, 1996
<u>AA</u>	<u>PROMETHAZINE W/ DEXTROMETHORPHAN</u>			
<u>AA</u>	WOCKHARDT	<u>15MG/5ML; 6.25MG/5ML</u>	<u>A088864 001</u>	Jan 04, 1985

DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE; ORAL

NUEDEXTA

+ AVANIR PHARMS 20MG;10MG

N021879 001 Oct 29, 2010

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

<u>AP</u>	+ B BRAUN	<u>10GM/100ML</u>	<u>N019626 004</u>	Feb 02, 1988
<u>AP</u>	+ BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N016694 001</u>	
<u>AP</u>	+ HOSPIRA	<u>10GM/100ML</u>	<u>N018080 001</u>	
	<u>DEXTROSE 20% IN PLASTIC CONTAINER</u>			
<u>AP</u>	+ BAXTER HLTHCARE	<u>20GM/100ML</u>	<u>N017521 004</u>	

<u>AP</u>	+ HOSPIRA	<u>20GM/100ML</u>	<u>N018564 001</u>	Mar 23, 1982
	<u>DEXTROSE 30% IN PLASTIC CONTAINER</u>			
<u>AP</u>	+ BAXTER HLTHCARE	<u>30GM/100ML</u>	<u>N017521 003</u>	
<u>AP</u>	+ HOSPIRA	<u>30GM/100ML</u>	<u>N019345 001</u>	Jan 26, 1985
	<u>DEXTROSE 40% IN PLASTIC CONTAINER</u>			

<u>AP</u>	+ BAXTER HLTHCARE	<u>40GM/100ML</u>	<u>N017521 002</u>	
<u>AP</u>	+ HOSPIRA	<u>40GM/100ML</u>	<u>N018562 001</u>	Mar 23, 1982
	<u>DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	+ B BRAUN	<u>50MG/ML</u>	<u>N016730 002</u>	

<u>AP</u>	+ B BRAUN	<u>5GM/100ML</u>	<u>N016730 001</u>	
<u>AP</u>	+ BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N019626 002</u>	Feb 02, 1988
<u>AP</u>	+ BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N016673 003</u>	Oct 30, 1985
<u>AP</u>	+ BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N020179 002</u>	Dec 07, 1992
<u>AP</u>	+ BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N016673 001</u>	

<u>AP</u>	+ BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N020179 001</u>	Dec 07, 1992
<u>AP</u>	+ HOSPIRA	<u>50MG/ML</u>	<u>N016367 002</u>	
<u>AP</u>	+ HOSPIRA	<u>50MG/ML</u>	<u>N019222 001</u>	Jul 13, 1984
<u>AP</u>	+ HOSPIRA	<u>50MG/ML</u>	<u>N019466 001</u>	Jul 15, 1985
<u>AP</u>	+ HOSPIRA	<u>5GM/100ML</u>	<u>N019479 001</u>	Sep 17, 1985

<u>AP</u>	+ BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N017521 001</u>	
<u>AP</u>	+ BAXTER HLTHCARE	<u>50GM/100ML</u>	<u>N020047 001</u>	Jul 02, 1991
<u>AP</u>	HOSPIRA	<u>500MG/ML</u>	<u>N019445 001</u>	Jun 03, 1986
<u>AP</u>	+ BAXTER HLTHCARE	<u>50GM/100ML</u>	<u>N018563 001</u>	Mar 23, 1982
	<u>DEXTROSE 50% IN PLASTIC CONTAINER</u>			

PRESCRIPTION DRUG PRODUCT LIST

3 - 127 (of 424)

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	<u>+</u>	HOSPIRA	<u>50GM/100ML</u>	<u>N019894</u>	<u>001</u>	Dec 26, 1989
			<u>DEXTROSE 60% IN PLASTIC CONTAINER</u>			
<u>AP</u>	<u>+</u>	BAXTER HLTHCARE	<u>60GM/100ML</u>	<u>N017521</u>	<u>005</u>	Mar 26, 1982
			<u>DEXTROSE 70% IN PLASTIC CONTAINER</u>			
<u>AP</u>	<u>+</u>	BAXTER HLTHCARE	<u>70GM/100ML</u>	<u>N017521</u>	<u>006</u>	Mar 26, 1982
<u>AP</u>	<u>+</u>		<u>70GM/100ML</u>	<u>N020047</u>	<u>003</u>	Jul 02, 1991
<u>AP</u>	<u>+</u>	HOSPIRA	<u>70GM/100ML</u>	<u>N018561</u>	<u>001</u>	Mar 23, 1982
<u>AP</u>	<u>+</u>		<u>70GM/100ML</u>	<u>N019893</u>	<u>001</u>	Dec 26, 1989
			<u>DEXTROSE 25%</u>			
		+ HOSPIRA	<u>250MG/ML</u>	<u>N019445</u>	<u>002</u>	Nov 23, 1998

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

DEXTROSE; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA 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, 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; 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

IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA 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

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

PRESCRIPTION DRUG PRODUCT LIST

3 - 128 (of 424)

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
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER	HOSPIRA	5GM/100ML;30MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML	N017609 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	

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;75MG/100ML</u>	<u>N017634 004</u>	
<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;150MG/100ML</u>	<u>N017634 001</u>	
<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;224MG/100ML</u>	<u>N017634 003</u>	
<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;300MG/100ML</u>	<u>N017634 002</u>	
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>B BRAUN</u>	<u>5GM/100ML;75MG/100ML</u>	<u>N018744 001</u>	Nov 09, 1982
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>B BRAUN</u>	<u>5GM/100ML;150MG/100ML</u>	<u>N018744 002</u>	Nov 09, 1982
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>B BRAUN</u>	<u>5GM/100ML;150MG/100ML</u>	<u>N019699 004</u>	Sep 29, 1989
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>B BRAUN</u>	<u>5GM/100ML;300MG/100ML</u>	<u>N018744 004</u>	Nov 09, 1982
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>B BRAUN</u>	<u>5GM/100ML;300MG/100ML</u>	<u>N019699 006</u>	Sep 29, 1989
<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>HOSPIRA</u>	<u>5GM/100ML;224MG/100ML</u>	<u>N018371 003</u>	
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER	B BRAUN	5GM/100ML;220MG/100ML	N018744 003	Nov 09, 1982
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER	HOSPIRA	5GM/100ML;149MG/100ML	N018371 001	
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER	HOSPIRA	5GM/100ML;298MG/100ML	N018371 002	

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
---	---------	--	-------------	--------------

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

<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ (K)</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;75MG/100ML;200MG/100ML</u>	<u>N018037 006</u>	Apr 13, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;150MG/100ML;200MG/100ML</u>	<u>N018037 007</u>	Apr 13, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)</u>	<u>BAXTER HLTHCARE</u>	<u>5GM/100ML;224MG/100ML;200MG/100ML</u>	<u>N018037 004</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 129 (of 424)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ</u>				
AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;200MG/100ML	N018037	008	Apr 13, 1982	
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)</u>				
AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;200MG/100ML	N018037	001		
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ</u>				
AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;200MG/100ML	N018037	005	Apr 13, 1982	
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ</u>				
AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;200MG/100ML	N018037	009	Apr 13, 1982	
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ</u>				
AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;200MG/100ML	N018037	002		
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)</u>				
AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;200MG/100ML	N018037	003		
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER</u>				
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	
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ 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 20MEQ IN PLASTIC CONTAINER</u>				
AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;330MG/100ML	N018629	004	Mar 23, 1982	
5GM/100ML;300MG/100ML;330MG/100ML	N018629	006	Mar 23, 1982	
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ 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 40MEQ 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 5MEQ 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 20MEQ (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 10MEQ 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	
5GM/100ML;150MG/100ML;450MG/100ML	N018008	006	Apr 28, 1982	
AP HOSPIRA 5GM/100ML;74.5MG/100ML;450MG/100ML	N018362	005	Mar 28, 1988	
5GM/100ML;74.5MG/100ML;450MG/100ML	N018362	009	Jul 05, 1983	
<u>POTASSIUM CHLORIDE 10MEQ 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	

PRESCRIPTION DRUG PRODUCT LIST

3 - 130 (of 424)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 002	Apr 05, 1985
AP	HOSPIRA	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 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	HOSPIRA	5GM/100ML;224MG/100ML;450MG/100ML	N018362 006	Mar 28, 1988
<u>POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	HOSPIRA	5GM/100ML;224MG/100ML;900MG/100ML	N019691 006	Mar 24, 1988
<u>POTASSIUM CHLORIDE 20MEQ 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	HOSPIRA	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 20MEQ 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	HOSPIRA	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 30MEQ 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	HOSPIRA	5GM/100ML;224MG/100ML;450MG/100ML	N018362 002	
<u>POTASSIUM CHLORIDE 30MEQ 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	HOSPIRA	5GM/100ML;224MG/100ML;900MG/100ML	N019691 007	Mar 24, 1988
<u>POTASSIUM CHLORIDE 40MEQ 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	HOSPIRA	5GM/100ML;298MG/100ML;450MG/100ML	N018362 003	
<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;900MG/100ML	N019308 007	Apr 05, 1985
AP	HOSPIRA	5GM/100ML;298MG/100ML;900MG/100ML	N019691 009	Mar 24, 1988
<u>POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;450MG/100ML	N018008 004	
AP	HOSPIRA	5GM/100ML;74.5MG/100ML;450MG/100ML	N018362 008	Mar 28, 1988
AP		5GM/100ML;149MG/100ML;450MG/100ML	N018362 004	Mar 28, 1988
<u>POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;900MG/100ML	N019308 001	Apr 05, 1985
AP	HOSPIRA	5GM/100ML;74.5MG/100ML;900MG/100ML	N019691 001	Mar 24, 1988
AP		5GM/100ML;149MG/100ML;900MG/100ML	N019691 003	Mar 24, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;37MG/100ML;200MG/100ML	N019630 031	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;37MG/100ML;450MG/100ML	N019630 037	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;37MG/100ML;900MG/100ML	N019630 043	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;37MG/100ML;110MG/100ML	N019630 001	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;37MG/100ML;200MG/100ML	N019630 007	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;37MG/100ML;330MG/100ML	N019630 013	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;37MG/100ML;450MG/100ML	N019630 019	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	5GM/100ML;37MG/100ML;900MG/100ML	N019630 025	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;75MG/100ML;200MG/100ML	N019630 032	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;75MG/100ML;450MG/100ML	N019630 038	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
	B BRAUN	10GM/100ML;75MG/100ML;900MG/100ML	N019630 044	Feb 17, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 131 (of 424)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

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
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

PRESCRIPTION DRUG PRODUCT LIST

3 - 132 (of 424)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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			
HOSPIRA	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 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP B BRAUN	<u>10GM/100ML;900MG/100ML</u>	<u>N019631 015</u>	Feb 24, 1988
AP BAXTER HLTHCARE	<u>10GM/100ML;900MG/100ML</u>	<u>N016696 001</u>	
<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP B BRAUN	<u>2.5GM/100ML;450MG/100ML</u>	<u>N019631 004</u>	Feb 24, 1988
AP BAXTER HLTHCARE	<u>2.5GM/100ML;450MG/100ML</u>	<u>N016697 001</u>	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP B BRAUN	<u>5GM/100ML;200MG/100ML</u>	<u>N019631 007</u>	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP B BRAUN	<u>5GM/100ML;330MG/100ML</u>	<u>N019631 008</u>	Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP B BRAUN	<u>5GM/100ML;450MG/100ML</u>	<u>N019631 009</u>	Feb 24, 1988
AP HOSPIRA	<u>5GM/100ML;450MG/100ML</u>	<u>N017607 001</u>	
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP B BRAUN	<u>5GM/100ML;900MG/100ML</u>	<u>N019631 010</u>	Feb 24, 1988
AP HOSPIRA	<u>5GM/100ML;900MG/100ML</u>	<u>N017585 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;200MG/100ML</u>	<u>N016689 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;330MG/100ML</u>	<u>N016687 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;450MG/100ML</u>	<u>N016683 001</u>	
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;900MG/100ML</u>	<u>N016678 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 133 (of 424)

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

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 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 0.2% IN PLASTIC CONTAINER B BRAUN	2.5GM/100ML;200MG/100ML	N019631 002	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 HOSPIRA	5GM/100ML;225MG/100ML	N017606 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER HOSPIRA	5GM/100ML;300MG/100ML	N017799 001	

DIATRIZOATE MEGLUMINE

SOLUTION; URETHRAL

CYSTOGRAFIN

BRACCO	30%	N010040 018
CYSTOGRAFIN DILUTE BRACCO	18%	N010040 022 Nov 09, 1982

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

MD-76R

<u>AP</u> + MALLINCKRODT	<u>66%;10%</u>	<u>N019292 001</u>	Sep 29, 1989
<u>AP</u> + BRACCO	<u>66%;10%</u>	<u>N010040 001</u>	

SOLUTION; ORAL, RECTAL

GASTROGRAFIN

<u>AA</u> + BRACCO	<u>66%;10%</u>	<u>N011245 003</u>	
<u>AA</u> MALLINCKRODT	<u>66%;10%</u>	<u>A087388 001</u>	

DIATRIZOATE MEGLUMINE; IODIPAMIDE MEGLUMINE

SOLUTION; INTRAUTERINE

SINOGRAPHIN

+ BRACCO	52.7%;26.8%	N011324 002
----------	-------------	-------------

DIAZEPAM

CONCENTRATE; ORAL

DIAZEPAM INTENSOL

+ ROXANE	5MG/ML	A071415 001	Apr 03, 1987
----------	--------	-------------	--------------

GEL; RECTAL

DIASTAT

VALEANT	2.5MG/0.5ML (5MG/ML)	N020648 001	Jul 29, 1997
---------	----------------------	-------------	--------------

DIASTAT ACUDIAL

VALEANT	10MG/2ML (5MG/ML)	N020648 007	Sep 15, 2005
---------	-------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 134 (of 424)

DIAZEPAM

GEL; RECTAL DIASTAT ACUDIAL + VALEANT	20MG/4ML (5MG/ML)	N020648 006	Sep 15, 2005
INJECTABLE; INJECTION			
<u>DIAZEPAM</u>			
<u>AP</u> + HOSPIRA	<u>5MG/ML</u>	<u>A071583 001</u>	Oct 13, 1987
<u>AP</u>	<u>5MG/ML</u>	<u>A071584 001</u>	Oct 13, 1987
<u>AP</u>	<u>5MG/ML</u>	<u>A072079 001</u>	Dec 20, 1988
<u>AP</u> WATSON LABS	<u>5MG/ML</u>	<u>A070296 001</u>	Feb 12, 1986
SOLUTION; ORAL DIAZEPAM + ROXANE	5MG/5ML	A070928 001	Apr 03, 1987
TABLET; ORAL			
<u>DIAZEPAM</u>			
<u>AB</u> BARR	<u>2MG</u>	<u>A070152 001</u>	Nov 01, 1985
<u>AB</u>	<u>10MG</u>	<u>A070154 001</u>	Nov 01, 1985
<u>AB</u> DAVA PHARMS INC	<u>2MG</u>	<u>A070226 001</u>	Sep 26, 1985
<u>AB</u> IVAX SUB TEVA PHARMS	<u>2MG</u>	<u>A071307 001</u>	Dec 10, 1986
<u>AB</u>	<u>5MG</u>	<u>A071321 001</u>	Dec 10, 1986
<u>AB</u>	<u>10MG</u>	<u>A071322 001</u>	Dec 10, 1986
<u>AB</u> MYLAN	<u>2MG</u>	<u>A070325 002</u>	Sep 04, 1985
<u>AB</u>	<u>5MG</u>	<u>A070325 003</u>	Sep 04, 1985
<u>AB</u>	<u>10MG</u>	<u>A070325 001</u>	Sep 04, 1985
<u>AB</u> VINTAGE PHARMS	<u>2MG</u>	<u>A077749 001</u>	Mar 31, 2006
<u>AB</u>	<u>5MG</u>	<u>A077749 002</u>	Mar 31, 2006
<u>AB</u>	<u>10MG</u>	<u>A077749 003</u>	Mar 31, 2006
<u>AB</u> WATSON LABS	<u>2MG</u>	<u>A071134 001</u>	Feb 03, 1987
<u>AB</u>	<u>5MG</u>	<u>A071135 001</u>	Feb 03, 1987
<u>AB</u>	<u>10MG</u>	<u>A071136 001</u>	Feb 03, 1987
<u>VALIUM</u>			
<u>AB</u> ROCHE	<u>2MG</u>	<u>N013263 002</u>	
<u>AB</u>	<u>5MG</u>	<u>N013263 004</u>	
<u>AB</u> +	<u>10MG</u>	<u>N013263 006</u>	

DIAZOXIDE

SUSPENSION; ORAL PROGLYCEM + TEVA GLOBAL	50MG/ML	N017453 001
--	---------	-------------

DICLOFENAC EPOLAMINE

PATCH; TOPICAL FLECTOR + INST BIOCHEM	1.3%	N021234 001	Jan 31, 2007
---	------	-------------	--------------

DICLOFENAC POTASSIUM

CAPSULE; ORAL ZIPSOR + XANODYNE PHARM	25MG	N022202 001	Jun 16, 2009
FOR SOLUTION; ORAL			
CAMBIA + NAUTILUS NEUROSCIENC	50MG	N022165 001	Jun 17, 2009
TABLET; ORAL			
<u>CATAFLAM</u>			
<u>AB</u> + NOVARTIS	<u>50MG</u>	<u>N020142 002</u>	Nov 24, 1993
<u>DICLOFENAC POTASSIUM</u>			
<u>AB</u> APOTEX	<u>50MG</u>	<u>A076561 001</u>	Mar 18, 2004
<u>AB</u> MYLAN	<u>50MG</u>	<u>A075463 001</u>	Jul 26, 1999
<u>AB</u> SANDOZ	<u>50MG</u>	<u>A075229 001</u>	Nov 20, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 135 (of 424)

DICLOFENAC POTASSIUM

TABLET; ORAL

DICLOFENAC POTASSIUMAB TEVA 50MG A075219 001 Aug 06, 1998DICLOFENAC SODIUM

GEL; TOPICAL

SOLARAZE

+ FOUGERA PHARMS 3% N021005 001 Oct 16, 2000
VOLTAREN

+ NOVARTIS 1% N022122 001 Oct 17, 2007

SOLUTION; TOPICAL

PENNNSAID

+ MALLINCKRODT 1.5% N020947 001 Nov 04, 2009

SOLUTION/DROPS; OPHTHALMIC

DICLOFENAC SODIUMAT AKORN 0.1% A077845 001 Apr 17, 2008
AT ALCON PHARMS LTD 0.1% A078031 001 Feb 06, 2008
AT APOTEX INC 0.1% A077600 001 Nov 13, 2008
AT BAUSCH AND LOMB 0.1% A078792 001 Dec 28, 2007
AT NEXUS PHARMS 0.1% A078553 001 Dec 28, 2007VOLTARENAT + NOVARTIS 0.1% N020037 001 Mar 28, 1991

TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUMAB ACTAVIS ELIZABETH 50MG A074514 001 Mar 26, 1996
AB 75MG A074514 002 Mar 26, 1996
AB ALPHAPHARM 50MG A075281 002 Feb 12, 2002
AB 75MG A075281 003 Feb 12, 2002
AB CARLSBAD 25MG A075185 002 Nov 13, 1998
AB 50MG A075185 003 Nov 13, 1998
AB 75MG A075185 001 Nov 13, 1998
AB + SANDOZ 25MG A074376 001 Sep 28, 1995
AB + 50MG A074376 002 Sep 28, 1995
AB + 75MG A074394 001 Nov 30, 1995
AB UNIQUE PHARM LABS 25MG A090066 001 Dec 01, 2010
AB 50MG A090066 002 Dec 01, 2010
AB 75MG A077863 003 Jun 08, 2007

TABLET, EXTENDED RELEASE; ORAL

DICLOFENAC SODIUMAB ACTAVIS ELIZABETH 100MG A075910 001 Jan 07, 2002
AB DEXCEL LTD 100MG A076201 001 Nov 06, 2002
AB MYLAN 100MG A076152 001 Dec 13, 2001
AB VALEANT INTL 100MG A075492 001 Feb 11, 2000VOLTAREN-XRAB + NOVARTIS 100MG N020254 001 Mar 08, 1996DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE; ORAL

ARTHROTEC

GD SEARLE LLC 50MG;0.2MG N020607 001 Dec 24, 1997
+ 75MG;0.2MG N020607 002 Dec 24, 1997DICLOXACILLIN SODIUM

CAPSULE; ORAL

DICLOXACILLIN SODIUMAB SANDOZ EQ 250MG BASE A061454 001
AB + EQ 500MG BASE A061454 003
AB TEVA EQ 250MG BASE A062286 001 Jun 03, 1982

PRESCRIPTION DRUG PRODUCT LIST

3 - 136 (of 424)

DICLOXACILLIN SODIUM

CAPSULE; ORAL

DICLOXACILLIN SODIUM

<u>AB</u>	TEVA	<u>EQ 500MG BASE</u>	<u>A062286 002</u>	Jun 03, 1982
	DICLOXACILLIN SODIUM SANDOZ	EQ 125MG BASE	A061454 002	

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

BENTYL

<u>AB</u>	+ APTALIS PHARMA US	<u>10MG</u>	<u>N007409 003</u>	Oct 15, 1984
	<u>DICYCLOMINE HYDROCHLORIDE</u>			
<u>AB</u>	LANNETT	<u>10MG</u>	<u>A084285 001</u>	
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A040319 001</u>	Sep 07, 1999
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A085082 001</u>	Jun 19, 1986
<u>AB</u>	WEST WARD	<u>10MG</u>	<u>A040204 001</u>	Feb 28, 1997

INJECTABLE; INJECTION

BENTYL

<u>AP</u>	+ APTALIS PHARMA US	<u>10MG/ML</u>	<u>N008370 001</u>	Oct 15, 1984
	<u>BENTYL PRESERVATIVE FREE</u>			
<u>AP</u>	+ APTALIS PHARMA US	<u>10MG/ML</u>	<u>N008370 002</u>	Oct 15, 1984
	<u>DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE-FREE)</u>			

<u>AP</u>	BEDFORD	<u>10MG/ML</u>	<u>A040465 001</u>	Jun 30, 2003
-----------	---------	----------------	--------------------	--------------

SYRUP; ORAL

BENTYL

<u>AA</u>	+ APTALIS PHARMA US	<u>10MG/5ML</u>	<u>N007961 002</u>	Oct 15, 1984
	<u>DICYCLOMINE HYDROCHLORIDE</u>			

<u>AA</u>	MIKART	<u>10MG/5ML</u>	<u>A040169 001</u>	Mar 24, 2005
-----------	--------	-----------------	--------------------	--------------

TABLET; ORAL

BENTYL

<u>AB</u>	+ APTALIS PHARMA US	<u>20MG</u>	<u>N007409 001</u>	Oct 15, 1984
	<u>DICYCLOMINE HYDROCHLORIDE</u>			
<u>AB</u>	LANNETT	<u>20MG</u>	<u>A040230 001</u>	Feb 26, 1999
<u>AB</u>	MYLAN	<u>20MG</u>	<u>A040317 001</u>	Sep 07, 1999
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>A085223 001</u>	Jul 30, 1986
<u>AB</u>	WEST WARD	<u>20MG</u>	<u>A040161 001</u>	Oct 01, 1996

DIDANOSINE

CAPSULE, DELAYED REL PELLETS; ORAL

DIDANOSINE

<u>AB</u>	AUROBINDO PHARMA	<u>125MG</u>	<u>A090094 001</u>	Sep 24, 2008
<u>AB</u>		<u>200MG</u>	<u>A090094 002</u>	Sep 24, 2008
<u>AB</u>		<u>250MG</u>	<u>A090094 003</u>	Sep 24, 2008
<u>AB</u>		<u>400MG</u>	<u>A090094 004</u>	Sep 24, 2008
<u>AB</u>	BARR	<u>200MG</u>	<u>A077167 001</u>	Dec 03, 2004
<u>AB</u>		<u>250MG</u>	<u>A077167 002</u>	Dec 03, 2004
<u>AB</u>		<u>400MG</u>	<u>A077167 003</u>	Dec 03, 2004
<u>AB</u>	MATRIX LABS LTD	<u>125MG</u>	<u>A090788 001</u>	Apr 08, 2010
<u>AB</u>		<u>200MG</u>	<u>A090788 002</u>	Apr 08, 2010
<u>AB</u>		<u>250MG</u>	<u>A090788 003</u>	Apr 08, 2010
<u>AB</u>		<u>400MG</u>	<u>A090788 004</u>	Apr 08, 2010

VIDEX EC

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>125MG</u>	<u>N021183 001</u>	Oct 31, 2000
<u>AB</u>		<u>200MG</u>	<u>N021183 002</u>	Oct 31, 2000
<u>AB</u>		<u>250MG</u>	<u>N021183 003</u>	Oct 31, 2000
<u>AB</u>	+	<u>400MG</u>	<u>N021183 004</u>	Oct 31, 2000

FOR SOLUTION; ORAL

DIDANOSINE

<u>AA</u>	AUROBINDO PHARMA	<u>10MG/ML</u>	<u>A078112 001</u>	Mar 08, 2007
-----------	------------------	----------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 137 (of 424)

DIDANOSINE

FOR SOLUTION; ORAL

VIDEXAA + BRISTOL MYERS SQUIBB 10MG/ML N020156 001 Oct 09, 1991DIENOGEST; ESTRADIOL VALERATE

TABLET; ORAL-28

NATAZIA

+ BAYER HLTHCARE N/A, 2MG, 3MG, N/A, N/A; 3MG, 2MG, 2MG, 1MG, N/A N022252 001 May 06, 2010

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDEAA AVANTHI INC 25MG A201212 001 Dec 22, 2010
AA COREPHARMA 25MG A040828 001 Nov 05, 2008
AA LANNETT HOLDINGS INC 25MG A200177 001 Jul 18, 2011TENUATEAA + WATSON PHARMS 25MG N011722 002

TABLET, EXTENDED RELEASE; ORAL

DIETHYLPROPION HYDROCHLORIDEAB LANNETT HOLDINGS INC 75MG A091680 001 Oct 24, 2011
TENUATE DOSPAN
AB + WATSON PHARMS 75MG N012546 001DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATEAB1 + ALTANA 0.05% A075187 001 Mar 30, 1998
AB1 TARO 0.05% A075508 001 Apr 24, 2000
DIFLORASONE DIACETATE
BX + ALTANA 0.05% A076263 001 Dec 20, 2002

OINTMENT; TOPICAL

DIFLORASONE DIACETATEAB ALTANA 0.05% A075374 001 Apr 27, 1999
AB + TARO 0.05% A075331 001 May 14, 1999DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

+ TEVA 500MG A073673 001 Jul 31, 1992

DIFLUPREDNATE

EMULSION; OPHTHALMIC

DUREZOL

+ ALCON PHARMS LTD 0.05% N022212 001 Jun 23, 2008

DIGOXIN

ELIXIR; ORAL

DIGOXIN

+ ROXANE 0.05MG/ML N021648 001 Aug 26, 2004

INJECTABLE; INJECTION

DIGOXINAP BAXTER HLTHCARE 0.25MG/ML A083391 001
AP HOSPIRA 0.25MG/ML A040093 001 May 16, 1996
AP SANDOZ 0.25MG/ML A040481 001 Aug 21, 2003LANOXINAP + COVIS PHARMA 0.25MG/ML N009330 002
LANOXIN PEDIATRIC
+ COVIS PHARMA 0.1MG/ML N009330 004

PRESCRIPTION DRUG PRODUCT LIST

3 - 138 (of 424)

DIGOXIN

TABLET; ORAL

DIGOXIN

<u>AB</u>	CARACO	<u>0.125MG</u>	<u>A076363</u>	<u>001</u>	Jan 31, 2003
<u>AB</u>		<u>0.25MG</u>	<u>A076363</u>	<u>002</u>	Jan 31, 2003
<u>AB</u>	IMPAK LABS	<u>0.125MG</u>	<u>A078556</u>	<u>001</u>	Jul 20, 2009
<u>AB</u>		<u>0.25MG</u>	<u>A078556</u>	<u>002</u>	Jul 20, 2009
<u>AB</u>	STEVENS J	<u>0.125MG</u>	<u>A076268</u>	<u>001</u>	Jul 26, 2002
<u>AB</u>		<u>0.25MG</u>	<u>A076268</u>	<u>002</u>	Jul 26, 2002
<u>AB</u>	WEST WARD	<u>0.125MG</u>	<u>A077002</u>	<u>002</u>	Oct 30, 2007
<u>AB</u>		<u>0.25MG</u>	<u>A077002</u>	<u>001</u>	Oct 30, 2007
	<u>LANOXIN</u>				
<u>AB</u>	COVIS PHARMA	<u>0.125MG</u>	<u>N020405</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	<u>+</u>	<u>0.25MG</u>	<u>N020405</u>	<u>004</u>	Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

<u>AP</u>	<u>+</u>	VALEANT	<u>1MG/ML</u>	<u>N005929</u>	<u>001</u>
<u>DIHYDROERGOTAMINE MESYLATE</u>					
<u>AP</u>		BEDFORD LABS	<u>1MG/ML</u>	<u>A040453</u>	<u>001</u>
<u>AP</u>		PADDICK LLC	<u>1MG/ML</u>	<u>A040475</u>	<u>001</u>

SPRAY, METERED; NASAL

MIGRANAL

+ VALEANT	0.5MG/INH
-----------	-----------

N020148 001 Dec 08, 1997

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILACOR XR

<u>AB2</u>	WATSON LABS	<u>120MG</u>	<u>N020092</u>	<u>001</u>	May 29, 1992
<u>AB2</u>		<u>180MG</u>	<u>N020092</u>	<u>002</u>	May 29, 1992
<u>AB2</u>	<u>+</u>	<u>240MG</u>	<u>N020092</u>	<u>003</u>	May 29, 1992

DILTIAZEM HYDROCHLORIDE

<u>AB2</u>	APOTEX	<u>120MG</u>	<u>A074943</u>	<u>003</u>	Dec 19, 2000
<u>AB2</u>		<u>180MG</u>	<u>A074943</u>	<u>002</u>	Dec 19, 2000
<u>AB2</u>		<u>240MG</u>	<u>A074943</u>	<u>001</u>	Aug 06, 1998
<u>AB2</u>	MYLAN	<u>120MG</u>	<u>A075124</u>	<u>002</u>	Mar 18, 1998
<u>AB2</u>		<u>180MG</u>	<u>A075124</u>	<u>003</u>	Mar 18, 1998
<u>AB2</u>		<u>240MG</u>	<u>A075124</u>	<u>001</u>	Mar 18, 1998
<u>AB2</u>	WATSON LABS FLORIDA	<u>120MG</u>	<u>A074852</u>	<u>001</u>	Oct 10, 1997
<u>AB2</u>		<u>180MG</u>	<u>A074852</u>	<u>002</u>	Oct 10, 1997
<u>AB2</u>		<u>240MG</u>	<u>A074852</u>	<u>003</u>	Oct 10, 1997

CARDIZEM CD

<u>AB3</u>	VALEANT INTL	<u>120MG</u>	<u>N020062</u>	<u>001</u>	Aug 10, 1992
<u>AB3</u>		<u>180MG</u>	<u>N020062</u>	<u>002</u>	Dec 27, 1991
<u>AB3</u>		<u>240MG</u>	<u>N020062</u>	<u>003</u>	Dec 27, 1991
<u>AB3</u>		<u>300MG</u>	<u>N020062</u>	<u>004</u>	Dec 27, 1991
<u>AB3</u>	<u>+</u>	<u>360MG</u>	<u>N020062</u>	<u>005</u>	Aug 24, 1999

CARTIA XT

<u>AB3</u>	WATSON LABS FLORIDA	<u>120MG</u>	<u>A074752</u>	<u>002</u>	Jul 09, 1998
<u>AB3</u>		<u>180MG</u>	<u>A074752</u>	<u>001</u>	Jul 09, 1998
<u>AB3</u>		<u>240MG</u>	<u>A074752</u>	<u>003</u>	Jul 09, 1998
<u>AB3</u>		<u>300MG</u>	<u>A074752</u>	<u>004</u>	Jul 09, 1998

DILT-CD

<u>AB3</u>	APOTEX	<u>120MG</u>	<u>A076151</u>	<u>001</u>	May 20, 2004
<u>AB3</u>		<u>180MG</u>	<u>A076151</u>	<u>002</u>	May 20, 2004
<u>AB3</u>		<u>240MG</u>	<u>A076151</u>	<u>003</u>	May 20, 2004
<u>AB3</u>		<u>300MG</u>	<u>A076151</u>	<u>004</u>	May 20, 2004

DILTIAZEM HYDROCHLORIDE

<u>AB3</u>	ACTAVIS ELIZABETH	<u>120MG</u>	<u>A074984</u>	<u>001</u>	Dec 20, 1999
------------	-------------------	--------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 140 (of 424)

DILTIAZEM HYDROCHLORIDE

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

+ HOSPIRA	100MG/VIAL	A075853 001	Dec 17, 2002
+ TEVA PARENTERAL	10MG/ML	A074894 002	Apr 19, 2002

TABLET; ORAL

CARDIZEM

AB VALEANT INTL	<u>30MG</u>	N018602 001	Nov 05, 1982
AB	<u>60MG</u>	N018602 002	Nov 05, 1982
AB	<u>90MG</u>	N018602 003	Dec 08, 1986
AB +	<u>120MG</u>	N018602 004	Dec 08, 1986
<u>DILTIAZEM HYDROCHLORIDE</u>			
AB DAVA PHARMS INC	<u>30MG</u>	A074093 001	Nov 05, 1992
AB	<u>60MG</u>	A074093 002	Nov 05, 1992
AB	<u>90MG</u>	A074093 003	Nov 05, 1992
AB	<u>120MG</u>	A074093 004	Nov 05, 1992
AB MYLAN	<u>30MG</u>	A072838 004	Nov 05, 1992
AB	<u>60MG</u>	A072838 003	Nov 05, 1992
AB	<u>90MG</u>	A072838 002	Nov 05, 1992
AB	<u>120MG</u>	A072838 001	Nov 05, 1992
AB TEVA	<u>30MG</u>	A074185 001	May 31, 1995
AB	<u>60MG</u>	A074185 002	May 31, 1995
AB	<u>90MG</u>	A074185 003	May 31, 1995
AB	<u>120MG</u>	A074185 004	May 31, 1995

TABLET, EXTENDED RELEASE; ORAL

CARDIZEM LA

AB VALEANT INTL	<u>120MG</u>	N021392 001	Feb 06, 2003
AB	<u>180MG</u>	N021392 002	Feb 06, 2003
AB	<u>240MG</u>	N021392 003	Feb 06, 2003
AB	<u>300MG</u>	N021392 004	Feb 06, 2003
AB	<u>360MG</u>	N021392 005	Feb 06, 2003
AB +	<u>420MG</u>	N021392 006	Feb 06, 2003
<u>DILTIAZEM HYDROCHLORIDE</u>			
AB WATSON LABS FLORIDA	<u>120MG</u>	A077686 006	Mar 15, 2010
AB	<u>180MG</u>	A077686 005	Mar 15, 2010
AB	<u>240MG</u>	A077686 004	Mar 15, 2010
AB	<u>300MG</u>	A077686 003	Mar 15, 2010
AB	<u>360MG</u>	A077686 002	Mar 15, 2010
AB	<u>420MG</u>	A077686 001	Mar 15, 2010

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

AP APP PHARMS	<u>50MG/ML</u>	A040519 001	Jun 23, 2004
AP + WATSON LABS	<u>50MG/ML</u>	A080615 001	

DIMERCAPROL

INJECTABLE; INJECTION

BAL

+ AKORN	10%	N005939 001
---------	-----	-------------

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

DIMETHYL SULFOXIDE

AT BIONICHE PHARMA	<u>50%</u>	A076185 001	Nov 29, 2002
RIMSO-50			
AT + BIONICHE PHARMA	<u>50%</u>	N017788 001	

PRESCRIPTION DRUG PRODUCT LIST

3 - 141 (of 424)

DINOPROSTONE

GEL; ENDOCERVICAL PREPIDIL		
+ PHARMACIA AND UPJOHN 0.5MG/3GM	N019617 001	Dec 09, 1992
INSERT, EXTENDED RELEASE; VAGINAL CERVIDIL		
+ CONTROLLED THERAP 10MG	N020411 001	Mar 30, 1995
SUPPOSITORY; VAGINAL PROSTIN E2		
+ PHARMACIA AND UPJOHN 20MG	N017810 001	

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 <u>BENADRYL</u>		
<u>AP + MCNEIL CONS 50MG/ML</u>	<u>N006146 002</u>	
<u>BENADRYL PRESERVATIVE FREE</u>		
<u>AP + MCNEIL CONS 50MG/ML</u>	<u>N009486 001</u>	
<u>DIPHENHYDRAMINE HYDROCHLORIDE</u>		
<u>AP APP PHARMS 50MG/ML</u>	<u>A040466 001</u>	May 28, 2002
<u>AP BAXTER HLTHCARE 50MG/ML</u>	<u>A080817 002</u>	
<u>AP BIONICHE PHARMA 50MG/ML</u>	<u>A040498 001</u>	Jul 12, 2005
<u>AP HOSPIRA 50MG/ML</u>	<u>A040140 001</u>	Nov 20, 1998
<u>AP WATSON LABS 50MG/ML</u>	<u>A080873 002</u>	
<u>DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE</u>		
<u>AP INTL MEDICATION 50MG/ML</u>	<u>A084094 001</u>	
<u>AP WATSON LABS 50MG/ML</u>	<u>A080873 003</u>	
DIPHENHYDRAMINE HYDROCHLORIDE		
+ WATSON LABS 10MG/ML	A080873 001	

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC <u>AKPRO</u>		
<u>AT AKORN 0.1%</u>	<u>A074382 001</u>	Sep 29, 1995
<u>DIPIVEFRIN HYDROCHLORIDE</u>		
<u>AT FALCON PHARMS 0.1%</u>	<u>A073636 001</u>	Jun 30, 1994
<u>PROPINE</u>		
<u>AT + ALLERGAN 0.1%</u>	<u>N018239 001</u>	

DIPYRIDAMOLE

INJECTABLE; INJECTION <u>DIPYRIDAMOLE</u>		
<u>AP APP PHARMS 5MG/ML</u>	<u>A074956 001</u>	Sep 30, 1998
<u>AP BAXTER HLTHCARE 5MG/ML</u>	<u>A074521 001</u>	Oct 18, 1996
<u>AP + BEDFORD 5MG/ML</u>	<u>A074939 001</u>	Apr 13, 1998
<u>AP CLARIS LIFESCIENCES 5MG/ML</u>	<u>A075769 001</u>	Nov 27, 2002
<u>AP TEVA PARENTERAL 5MG/ML</u>	<u>A074952 001</u>	Nov 26, 1997
TABLET; ORAL <u>DIPYRIDAMOLE</u>		
<u>AB BARR 25MG</u>	<u>A087184 001</u>	Oct 03, 1990
<u>AB 50MG</u>	<u>A087716 001</u>	Oct 03, 1990
<u>AB 75MG</u>	<u>A087717 001</u>	Oct 03, 1990
<u>AB GLENMARK GENERICS 25MG</u>	<u>A088999 001</u>	Feb 05, 1991
<u>AB 50MG</u>	<u>A089000 001</u>	Feb 05, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 142 (of 424)

DIPYRIDAMOLE

TABLET; ORAL

DIPYRIDAMOLE

AB	GLENMARK GENERICS	<u>75MG</u>	<u>A089001</u> <u>001</u>	Feb 05, 1991
AB	IMPAK LABS	<u>25MG</u>	<u>A040782</u> <u>001</u>	Jul 18, 2007
AB		<u>50MG</u>	<u>A040782</u> <u>002</u>	Jul 18, 2007
AB		<u>75MG</u>	<u>A040782</u> <u>003</u>	Jul 18, 2007
AB	LANNETT	<u>25MG</u>	<u>A040898</u> <u>001</u>	Apr 23, 2008
AB		<u>50MG</u>	<u>A040898</u> <u>002</u>	Apr 23, 2008
AB		<u>75MG</u>	<u>A040898</u> <u>003</u>	Apr 23, 2008
AB	MURTY PHARMS	<u>25MG</u>	<u>A040733</u> <u>001</u>	Feb 13, 2007
AB		<u>50MG</u>	<u>A040733</u> <u>002</u>	Feb 13, 2007
AB		<u>75MG</u>	<u>A040733</u> <u>003</u>	Feb 13, 2007
AB	PROSAM LABS	<u>25MG</u>	<u>A040542</u> <u>001</u>	Apr 21, 2006
AB		<u>50MG</u>	<u>A040542</u> <u>002</u>	Apr 21, 2006
AB		<u>75MG</u>	<u>A040542</u> <u>003</u>	Apr 21, 2006
AB	PUREPAC PHARM	<u>25MG</u>	<u>A089425</u> <u>001</u>	Jul 12, 1990
AB	WATSON LABS	<u>50MG</u>	<u>A087160</u> <u>001</u>	Jun 07, 1996
AB	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A040874</u> <u>001</u>	Jan 28, 2008
AB		<u>50MG</u>	<u>A040874</u> <u>002</u>	Jan 28, 2008
AB		<u>75MG</u>	<u>A040874</u> <u>003</u>	Jan 28, 2008
<u>PERSANTINE</u>				
AB	BOEHRINGER INGELHEIM	<u>25MG</u>	<u>N012836</u> <u>003</u>	Dec 22, 1986
AB		<u>50MG</u>	<u>N012836</u> <u>004</u>	Feb 06, 1987
AB	+	<u>75MG</u>	<u>N012836</u> <u>005</u>	Feb 06, 1987

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

AB	TEVA	<u>EQ 100MG BASE</u>	<u>A070101</u> <u>001</u>	Feb 22, 1985
AB		<u>EQ 150MG BASE</u>	<u>A070102</u> <u>001</u>	Feb 22, 1985
AB	WATSON LABS	<u>EQ 100MG BASE</u>	<u>A070173</u> <u>001</u>	May 31, 1985
AB		<u>EQ 150MG BASE</u>	<u>A070174</u> <u>001</u>	May 31, 1985
<u>NORPACE</u>				
AB	GD SEARLE LLC	<u>EQ 100MG BASE</u>	<u>N017447</u> <u>001</u>	
AB	+	<u>EQ 150MG BASE</u>	<u>N017447</u> <u>002</u>	

CAPSULE, EXTENDED RELEASE; ORAL

DISOPYRAMIDE PHOSPHATE

AB	NESHER PHARMS	<u>EQ 150MG BASE</u>	<u>A071200</u> <u>001</u>	Dec 15, 1987
AB	<u>NORPACE CR</u>			
AB	+ GD SEARLE LLC	<u>EQ 150MG BASE</u>	<u>N018655</u> <u>002</u>	Jul 20, 1982

NORPACE CR

GD SEARLE LLC EQ 100MG BASE N018655 001 Jul 20, 1982

DISULFIRAM

TABLET; ORAL

ANTABUSE

AB	ODYSSEY PHARMS	<u>250MG</u>	<u>A088482</u> <u>001</u>	Dec 08, 1983
AB	+	<u>500MG</u>	<u>A088483</u> <u>001</u>	Dec 08, 1983
<u>DISULFIRAM</u>				
AB	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A091619</u> <u>001</u>	Mar 28, 2011
AB		<u>500MG</u>	<u>A091619</u> <u>002</u>	Mar 28, 2011

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS; ORAL

DEPAKOTE

AB	+ ABBOTT	<u>EQ 125MG VALPROIC ACID</u>	<u>N019680</u> <u>001</u>	Sep 12, 1989
<u>DIVALPROEX SODIUM</u>				
AB	DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078979</u> <u>001</u>	Jan 23, 2009
AB	MYLAN	<u>EQ 125MG VALPROIC ACID</u>	<u>A090407</u> <u>001</u>	Mar 28, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 143 (of 424)

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS; ORAL

DIVALPROEX SODIUM

<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A078919 001</u>	Jan 27, 2009
		TABLET, DELAYED RELEASE; ORAL		
		<u>DEPAKOTE</u>		
<u>AB</u>	ABBOTT	<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>	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>A077254 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 125MG VALPROIC ACID</u>	<u>A090062 001</u>	Mar 17, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077254 002</u>	Jul 29, 2008
<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>A077254 003</u>	Jul 29, 2008
<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
<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>	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	<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>	VINTAGE	<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>	WATSON LABS FLORIDA	<u>EQ 500MG VALPROIC ACID</u>	<u>A079080 001</u>	Feb 25, 2011
<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>	ABBOTT	<u>EQ 250MG VALPROIC ACID</u>	<u>N021168 002</u>	May 31, 2002
-----------	--------	-------------------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 144 (of 424)

DIVALPROEX SODIUM

TABLET, EXTENDED RELEASE; ORAL

DEPAKOTE ER

<u>AB</u> + ABBOTT	<u>EQ 500MG VALPROIC ACID</u>	<u>N021168</u> <u>001</u>	Aug 04, 2000
<u>DIVALPROEX SODIUM</u>			
<u>AB</u> ANCHEN PHARMS	<u>EQ 250MG VALPROIC ACID</u>	<u>A078445</u> <u>001</u>	Feb 26, 2009
<u>AB</u>	<u>EQ 500MG VALPROIC ACID</u>	<u>A078445</u> <u>002</u>	Aug 04, 2009
<u>AB</u> IMPAX LABS	<u>EQ 250MG VALPROIC ACID</u>	<u>A078791</u> <u>001</u>	May 06, 2009
<u>AB</u>	<u>EQ 500MG VALPROIC ACID</u>	<u>A078791</u> <u>002</u>	Aug 04, 2009
<u>AB</u> MYLAN	<u>EQ 250MG VALPROIC ACID</u>	<u>A077567</u> <u>001</u>	Jan 29, 2009
<u>AB</u>	<u>EQ 500MG VALPROIC ACID</u>	<u>A077567</u> <u>002</u>	Jan 29, 2009
<u>AB</u> TEVA PHARMS	<u>EQ 500MG VALPROIC ACID</u>	<u>A078700</u> <u>001</u>	Aug 03, 2009
<u>AB</u> WOCKHARDT	<u>EQ 250MG VALPROIC ACID</u>	<u>A078705</u> <u>002</u>	Feb 10, 2009
<u>AB</u>	<u>EQ 500MG VALPROIC ACID</u>	<u>A078705</u> <u>001</u>	Aug 04, 2009
<u>AB</u> ZYDUS PHARMS USA INC	<u>EQ 250MG VALPROIC ACID</u>	<u>A078239</u> <u>001</u>	Feb 27, 2009
<u>AB</u>	<u>EQ 500MG VALPROIC ACID</u>	<u>A078239</u> <u>002</u>	Aug 04, 2009

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

<u>AP</u> BEDFORD	<u>EQ 12.5MG BASE/ML</u>	<u>A074277</u> <u>001</u>	Oct 31, 1994
<u>AP</u> HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>A074086</u> <u>001</u>	Nov 29, 1993
<u>AP</u> +	<u>EQ 12.5MG BASE/ML</u>	<u>A074292</u> <u>001</u>	Feb 16, 1995
<u>AP</u> WATSON LABS	<u>EQ 12.5MG BASE/ML</u>	<u>A074114</u> <u>001</u>	Nov 30, 1993
<u>DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%</u>			
<u>AP</u> + HOSPIRA	<u>EQ 50MG BASE/100ML</u>	<u>N020269</u> <u>001</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 100MG BASE/100ML</u>	<u>N020269</u> <u>002</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 200MG BASE/100ML</u>	<u>N020269</u> <u>003</u>	Oct 19, 1993
<u>DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u> + BAXTER HLTHCARE	<u>EQ 50MG BASE/100ML</u>	<u>N020255</u> <u>001</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 100MG BASE/100ML</u>	<u>N020255</u> <u>003</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 200MG BASE/100ML</u>	<u>N020255</u> <u>004</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 400MG BASE/100ML</u>	<u>N020255</u> <u>005</u>	Oct 19, 1993
<u>AP</u> + HOSPIRA	<u>EQ 50MG BASE/100ML</u>	<u>N020201</u> <u>003</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 100MG BASE/100ML</u>	<u>N020201</u> <u>002</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 200MG BASE/100ML</u>	<u>N020201</u> <u>001</u>	Oct 19, 1993
<u>AP</u> +	<u>EQ 400MG BASE/100ML</u>	<u>N020201</u> <u>006</u>	Jul 07, 1994

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

<u>AP</u> + HOSPIRA INC	<u>20MG/2ML (10MG/ML)</u>	<u>N022234</u> <u>001</u>	Mar 08, 2011
<u>AP</u> +	<u>80MG/8ML (10MG/ML)</u>	<u>N022234</u> <u>002</u>	Mar 08, 2011
<u>AP</u> +	<u>160MG/16ML (10MG/ML)</u>	<u>N022234</u> <u>003</u>	Mar 08, 2011
<u>AP</u> SANDOZ	<u>20MG/2ML (10MG/ML)</u>	<u>N201525</u> <u>001</u>	Jun 29, 2011
<u>AP</u>	<u>160MG/16ML (10MG/ML)</u>	<u>N201525</u> <u>003</u>	Jun 29, 2011
<u>AP</u> SANDOZ INC	<u>80MG/8ML (10MG/ML)</u>	<u>N201525</u> <u>002</u>	Jun 29, 2011
DOCEFREZ			
+ SUN PHARMA GLOBAL	20MG/VIAL	N022534 001	May 03, 2011
+	80MG/VIAL	N022534 002	May 03, 2011
DOCETAXEL			
ACCORD HLTHCARE	20MG/0.5ML (40MG/ML) 80MG/2ML (40MG/ML)	N201195 001 N201195 002	Jun 08, 2011 Jun 08, 2011
TAXOTERE			
+ SANOFI AVENTIS US	20MG/ML (20MG/ML) 80MG/4ML (20MG/ML)	N020449 003 N020449 004	Aug 03, 2010 Aug 02, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 145 (of 424)

DOFETILIDE

CAPSULE; ORAL				
TIKOSYN				
PFIZER	0.125MG		N020931 001	Oct 01, 1999
	0.25MG		N020931 002	Oct 01, 1999
+	0.5MG		N020931 003	Oct 01, 1999

DOLASETRON MESYLATE

INJECTABLE; INJECTION				
ANZEMET				
+	SANOFI AVENTIS US	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
TABLET; ORAL				
ANZEMET				
SANOFI AVENTIS US	50MG		N020623 001	Sep 11, 1997
+	100MG		N020623 002	Sep 11, 1997

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL				
<u>ARICEPT</u>				
<u>AB</u>	EISAI INC	<u>5MG</u>	<u>N020690 002</u>	Nov 25, 1996
<u>AB</u>	+	<u>10MG</u>	<u>N020690 001</u>	Nov 25, 1996
	<u>DONEPEZIL HYDROCHLORIDE</u>			
<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A201335 001</u>	Aug 29, 2011
<u>AB</u>		<u>10MG</u>	<u>A201335 002</u>	Aug 29, 2011
<u>AB</u>	ACTAVIS PHARMA	<u>5MG</u>	<u>A090551 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090551 002</u>	May 31, 2011
<u>AB</u>	APOTEX	<u>5MG</u>	<u>A078841 001</u>	Jun 02, 2011
<u>AB</u>		<u>10MG</u>	<u>A078841 002</u>	Jun 02, 2011
<u>AB</u>	AUROBINDO	<u>5MG</u>	<u>A090056 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090056 002</u>	May 31, 2011
<u>AB</u>	CIPILA LTD	<u>5MG</u>	<u>A077518 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A077518 002</u>	May 31, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A201001 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A201001 002</u>	May 31, 2011
<u>AB</u>	HIKMA PHARMS	<u>5MG</u>	<u>A090247 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090247 002</u>	May 31, 2011
<u>AB</u>	HUAHAI US INC	<u>5MG</u>	<u>A200292 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A200292 002</u>	May 31, 2011
<u>AB</u>	JUBILANT LIFE	<u>5MG</u>	<u>A090768 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090768 002</u>	May 31, 2011
<u>AB</u>	MATRIX LABS LTD	<u>5MG</u>	<u>A090521 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090521 002</u>	May 31, 2011
<u>AB</u>	PLIVA HRVATSKA DOO	<u>5MG</u>	<u>A090425 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090425 002</u>	May 31, 2011
<u>AB</u>	RANBAXY	<u>5MG</u>	<u>A076786 001</u>	Nov 26, 2010
<u>AB</u>		<u>10MG</u>	<u>A076786 002</u>	Nov 26, 2010
<u>AB</u>	ROXANE	<u>5MG</u>	<u>A078662 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A078662 002</u>	May 31, 2011
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A090290 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090290 002</u>	May 31, 2011
<u>AB</u>	SUN PHARM INDs	<u>5MG</u>	<u>A090493 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090493 002</u>	May 31, 2011
<u>AB</u>	TEVA	<u>5MG</u>	<u>A077344 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A077344 002</u>	May 31, 2011
<u>AB</u>	TORRENT PHARMS	<u>5MG</u>	<u>A090686 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A090686 002</u>	May 31, 2011
<u>AB</u>	WOCKHARDT	<u>5MG</u>	<u>A091267 001</u>	May 31, 2011
<u>AB</u>		<u>10MG</u>	<u>A091267 002</u>	May 31, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 146 (of 424)

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL ARICEPT + EISAI INC	23MG	N022568 001	Jul 23, 2010
TABLET, ORALLY DISINTEGRATING; ORAL <u>ARICEPT ODT</u>			
<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
<u>DONEPEZIL HYDROCHLORIDE</u>			
<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> 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> 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

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION <u>DOPAMINE HYDROCHLORIDE</u>			
<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
<u>AP</u> +	<u>160MG/ML</u>	<u>A070826 001</u>	Feb 11, 1987
<u>DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%</u>			
<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
<u>DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u> + B BRAUN	<u>160MG/100ML</u>	<u>N019099 003</u>	Oct 15, 1986
<u>DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u> + BAXTER HLTHCARE	<u>80MG/100ML</u>	<u>N019615 001</u>	Mar 27, 1987
<u>AP</u> +	<u>160MG/100ML</u>	<u>N019615 002</u>	Mar 27, 1987
<u>AP</u> +	<u>320MG/100ML</u>	<u>N019615 003</u>	Mar 27, 1987
<u>AP</u> + HOSPIRA	<u>80MG/100ML</u>	<u>N018826 001</u>	Sep 30, 1983
<u>AP</u> +	<u>160MG/100ML</u>	<u>N018826 002</u>	Sep 30, 1983
<u>AP</u> +	<u>320MG/100ML</u>	<u>N018826 003</u>	Sep 30, 1983
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER			
+ B BRAUN	40MG/100ML	N019099 001	Oct 15, 1986
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER			
+ BAXTER HLTHCARE	640MG/100ML	N019615 004	Mar 27, 1987

DORIPENEM

INJECTABLE; IV (INFUSION)			
JANSSEN PHARMS	250MG/VIAL	N022106 002	Oct 05, 2010
+	500MG/VIAL	N022106 001	Oct 12, 2007

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC <u>DORZOLAMIDE HYDROCHLORIDE</u>			
AT ALCON PHARMS LTD	<u>EQ 2% BASE</u>	<u>A078981 001</u>	Apr 13, 2009
AT APOTEX INC	<u>EQ 2% BASE</u>	<u>A078395 001</u>	Oct 28, 2008
AT BAUSCH AND LOMB	<u>EQ 2% BASE</u>	<u>A090143 001</u>	Jun 25, 2009
AT HI TECH PHARMA	<u>EQ 2% BASE</u>	<u>A077846 001</u>	Oct 28, 2008
AT LUITPOLD	<u>EQ 2% BASE</u>	<u>A079186 001</u>	Nov 18, 2009
AT SANDOZ	<u>EQ 2% BASE</u>	<u>A078748 001</u>	Nov 06, 2008
AT TEVA PHARMS	<u>EQ 2% BASE</u>	<u>A078756 001</u>	Dec 04, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 147 (of 424)

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

TRUSOPTAT + MERCK EQ 2% BASE N020408 001 Dec 09, 1994DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COSOPT

AT + MERCK	<u>EQ 2% BASE;EQ 0.5% BASE</u>	N020869 001	Apr 07, 1998
<u>DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE</u>			
AT ALCON RES	<u>EQ 2% BASE;EQ 0.5% BASE</u>	A090604 001	Nov 18, 2009
AT APOTEX INC	<u>EQ 2% BASE;EQ 0.5% BASE</u>	A078201 001	Oct 28, 2008
AT BAUSCH AND LOMB	<u>EQ 2% BASE;EQ 0.5% BASE</u>	A090037 001	Jul 14, 2009
AT HI TECH PHARMA	<u>EQ 2% BASE;EQ 0.5% BASE</u>	A077847 001	Oct 28, 2008
AT SANDOZ	<u>EQ 2% BASE;EQ 0.5% BASE</u>	A078749 001	Nov 06, 2008
AT TEVA PARNTL	<u>EQ 2% BASE;EQ 0.5% BASE</u>	A078704 001	Sep 28, 2009

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOPRAM

AP + BAXTER HLTHCARE	<u>20MG/ML</u>	N014879 001	
<u>DOXAPRAM HYDROCHLORIDE</u>			
AP BEDFORD	<u>20MG/ML</u>	A076266 001	Jan 10, 2003
AP WATSON LABS	<u>20MG/ML</u>	A073529 001	Jan 30, 1992

DOXAZOSIN MESYLATE

TABLET; ORAL

CARDURA

AB + PFIZER	<u>EQ 1MG BASE</u>	N019668 001	Nov 02, 1990
AB	<u>EQ 2MG BASE</u>	N019668 002	Nov 02, 1990
AB	<u>EQ 4MG BASE</u>	N019668 003	Nov 02, 1990
AB	<u>EQ 8MG BASE</u>	N019668 004	Nov 02, 1990

DOXAZOSIN MESYLATE

AB APOTEX	<u>EQ 1MG BASE</u>	A075580 001	Oct 18, 2000
AB	<u>EQ 2MG BASE</u>	A075580 002	Oct 18, 2000
AB	<u>EQ 4MG BASE</u>	A075580 003	Oct 18, 2000
AB	<u>EQ 8MG BASE</u>	A075580 004	Oct 18, 2000
AB DAVA PHARMS INC	<u>EQ 1MG BASE</u>	A076161 001	Jun 10, 2004
AB	<u>EQ 2MG BASE</u>	A076161 002	Jun 10, 2004
AB	<u>EQ 4MG BASE</u>	A076161 003	Jun 10, 2004
AB	<u>EQ 8MG BASE</u>	A076161 004	Jun 10, 2004
AB MYLAN	<u>EQ 1MG BASE</u>	A075509 001	Oct 19, 2000
AB	<u>EQ 2MG BASE</u>	A075509 002	Oct 19, 2000
AB	<u>EQ 4MG BASE</u>	A075509 003	Oct 19, 2000
AB	<u>EQ 8MG BASE</u>	A075509 004	Oct 19, 2000
AB NESHER PHARMS	<u>EQ 1MG BASE</u>	A075609 001	Oct 18, 2000
AB	<u>EQ 2MG BASE</u>	A075609 002	Oct 18, 2000
AB	<u>EQ 4MG BASE</u>	A075609 003	Oct 18, 2000
AB	<u>EQ 8MG BASE</u>	A075609 004	Oct 18, 2000
AB PLIVA	<u>EQ 1MG BASE</u>	A075750 001	Jun 08, 2001
AB	<u>EQ 2MG BASE</u>	A075750 002	Jun 08, 2001
AB	<u>EQ 4MG BASE</u>	A075750 003	Jun 08, 2001
AB	<u>EQ 8MG BASE</u>	A075750 004	Jun 08, 2001
AB TEVA	<u>EQ 1MG BASE</u>	A075536 001	Oct 18, 2000
AB	<u>EQ 2MG BASE</u>	A075536 002	Oct 18, 2000
AB	<u>EQ 4MG BASE</u>	A075536 003	Oct 18, 2000
AB	<u>EQ 8MG BASE</u>	A075536 004	Oct 18, 2000
AB WATSON LABS	<u>EQ 1MG BASE</u>	A075426 001	Oct 18, 2000
AB	<u>EQ 2MG BASE</u>	A075426 002	Oct 18, 2000
AB	<u>EQ 4MG BASE</u>	A075426 003	Oct 18, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 148 (of 424)

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

<u>AB</u>	WATSON LABS	<u>EQ 8MG BASE</u>	<u>A075426 004</u>	Oct 18, 2000
TABLET, EXTENDED RELEASE; ORAL				
CARDURA XL				

PFIZER

+

EQ 4MG BASE

EQ 8MG BASE

N021269 001 Feb 22, 2005

N021269 002 Feb 22, 2005

DOXE PIN HYDROCHLORIDE

CAPSULE; ORAL

DOXE PIN HYDROCHLORIDE

<u>AB</u>	MYLAN	<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
<u>AB</u>	PAR PHARM	<u>EQ 10MG BASE</u>	<u>A071697 001</u>	Nov 09, 1987
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>A071437 001</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A071595 001</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A071608 001</u>	Nov 09, 1987
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>A071422 001</u>	Nov 09, 1987
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>A071485 001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A071486 001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A071238 001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A071326 001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A071239 001</u>	Apr 30, 1987
	DOXE PIN HYDROCHLORIDE			
	PAR PHARM	EQ 150MG BASE	A071669 001	Nov 09, 1987

CONCENTRATE; ORAL

DOXE PIN HYDROCHLORIDE

<u>AA</u>	PHARM ASSOC	<u>EQ 10MG BASE/ML</u>	<u>A075924 001</u>	Jan 15, 2004
<u>AA</u>	SILARX	<u>EQ 10MG BASE/ML</u>	<u>A074721 001</u>	Dec 29, 1998
<u>AA</u>	+	<u>TEVA PHARMS</u>	<u>A071609 001</u>	Nov 09, 1987
<u>AA</u>	WOCKHARDT	<u>EQ 10MG BASE/ML</u>	<u>A071918 001</u>	Jul 20, 1988

CREAM; TOPICAL

ZONALON

+ NYCOMED US

5%

N020126 001 Apr 01, 1994

TABLET; ORAL

SILENOR

SOMAXON

EQ 3MG BASE

N022036 001 Mar 17, 2010

+

EQ 6MG BASE

N022036 002 Mar 17, 2010

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

<u>AB</u>	ROXANE	<u>0.5MCG</u>	<u>A091433 001</u>	Sep 23, 2011
<u>AB</u>	HECTOROL	<u>0.5MCG</u>	<u>N020862 002</u>	Apr 23, 2004
<u>AB</u>	GENZYME CORP	<u>0.5MCG</u>		
	HECTOROL	1MCG		
	GENZYME CORP	2.5MCG		
	+			

INJECTABLE; INJECTION

HECTOROL

GENZYME CORP

2MCG/ML (2MCG/ML)

N021027 002 Apr 06, 2000

+

4MCG/2ML (2MCG/ML)

N021027 001 Apr 06, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 149 (of 424)

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>	APP PHARMS	<u>2MG/ML</u>	<u>A063277</u>	<u>001</u>	Oct 26, 1995
<u>AP</u>	+ BEDFORD	<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
<u>AP</u>	ONCO THERAPIES LTD	<u>10MG/VIAL</u>	<u>A200170</u>	<u>001</u>	Oct 28, 2011
<u>AP</u>		<u>50MG/VIAL</u>	<u>A200170</u>	<u>002</u>	Oct 28, 2011
<u>AP</u>	PHARMACHEMIE	<u>2MG/ML</u>	<u>A063336</u>	<u>001</u>	Feb 28, 1995
<u>AP</u>		<u>10MG/VIAL</u>	<u>A063097</u>	<u>001</u>	May 21, 1990
<u>AP</u>		<u>20MG/VIAL</u>	<u>A063097</u>	<u>002</u>	May 21, 1990
<u>AP</u>		<u>50MG/VIAL</u>	<u>A063097</u>	<u>003</u>	May 21, 1990
<u>AP</u>		<u>200MG/100ML</u>	<u>A063336</u>	<u>004</u>	Feb 28, 1995
<u>AP</u>	TEVA PARENTERAL	<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

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL

+ ORTHO BIOTECH	20MG/10ML (2MG/ML)	N050718	001	Nov 17, 1995
+ ORTHO BIOTECH	50MG/25ML (2MG/ML)	N050718	002	Jun 13, 2000

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

<u>AB</u>	IMPAK LABS INC	<u>EQ 150MG BASE</u>	<u>A200065</u>	<u>001</u>	Feb 17, 2011
<u>AB</u>	MYLAN	<u>40MG</u>	<u>A090855</u>	<u>001</u>	Jul 01, 2010
<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>	RANBAXY	<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>	SANDOZ	<u>EQ 50MG BASE</u>	<u>A065032</u>	<u>001</u>	Jun 30, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065032</u>	<u>002</u>	Jun 30, 2000
<u>AB</u>	WATSON LABS	<u>EQ 50MG BASE</u>	<u>A065041</u>	<u>001</u>	Apr 28, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065041</u>	<u>002</u>	Apr 28, 2000

MONODOX

<u>AB</u>	WATSON 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

<u>AB</u>	+ GALDERMA LABS LP	<u>40MG</u>	<u>N050805</u>	<u>001</u>	May 26, 2006
-----------	--------------------	-------------	----------------	------------	--------------

FOR SUSPENSION; ORAL

DOXYCYCLINE

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 25MG BASE/5ML</u>	<u>A065454</u>	<u>001</u>	Jul 16, 2008
<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
<u>AB</u>	MYLAN	<u>EQ 50MG BASE</u>	<u>A065377</u>	<u>001</u>	Nov 07, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 150 (of 424)

DOXYCYCLINE

TABLET; ORAL

DOXYCYCLINE

<u>AB</u>	MYLAN	<u>EQ 75MG BASE</u>	<u>A065377</u>	<u>002</u>	Nov 07, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065377</u>	<u>003</u>	Nov 07, 2006
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065427</u>	<u>001</u>	Jun 07, 2007
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A065070</u>	<u>001</u>	Dec 15, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065070</u>	<u>003</u>	Dec 30, 2002
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065070</u>	<u>002</u>	Dec 15, 2000
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>A065070</u>	<u>004</u>	Jul 14, 2005
<u>AB</u>	RANBAXY	<u>EQ 50MG BASE</u>	<u>A065356</u>	<u>001</u>	May 31, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065356</u>	<u>002</u>	May 31, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065356</u>	<u>003</u>	May 31, 2006
<u>AB</u>	SANDOZ	<u>EQ 50MG BASE</u>	<u>A065353</u>	<u>001</u>	Nov 27, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065353</u>	<u>002</u>	Nov 27, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065353</u>	<u>003</u>	Nov 27, 2006

DOXYCYCLINE CALCIUM

SUSPENSION; ORAL

VIBRAMYCIN

+ PFIZER EQ 50MG BASE/5ML

N050480 001

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 50MG BASE</u>	<u>A062500</u>	<u>001</u>	Sep 11, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062500</u>	<u>002</u>	Sep 11, 1984
<u>AB</u>	MUTUAL PHARM	<u>EQ 50MG BASE</u>	<u>A062675</u>	<u>001</u>	Jul 10, 1986
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062676</u>	<u>001</u>	Jul 10, 1986
<u>AB</u>	WATSON LABS FLORIDA	<u>EQ 50MG BASE</u>	<u>A062031</u>	<u>002</u>	Oct 13, 1982
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062031</u>	<u>001</u>	
<u>AB</u>	WEST WARD	<u>EQ 50MG BASE</u>	<u>A062396</u>	<u>002</u>	Nov 07, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062396</u>	<u>001</u>	May 07, 1984

VIBRAMYCIN

<u>AB</u>	PFIZER	<u>EQ 50MG BASE</u>	<u>N050007</u>	<u>001</u>
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N050007</u>	<u>002</u>

DOXYCYCLINE HYCLATE

+ WEST WARD EQ 20MG BASE

A065103 001 May 13, 2005

CAPSULE, DELAYED RELEASE; ORAL

DOXYCYCLINE HYCLATE

MEDICIS EQ 75MG BASE
+ EQ 100MG BASEA065281 001 Dec 21, 2005
A065281 002 Dec 21, 2005

INJECTABLE; INJECTION

DOXY 100

<u>AP</u>	+ APP PHARMS	<u>EQ 100MG BASE/VIAL</u>	<u>A062475</u>	<u>001</u>	Dec 09, 1983
-----------	--------------	---------------------------	----------------	------------	--------------

DOXYCYCLINE

<u>AP</u>	+ BEDFORD	<u>EQ 100MG BASE/VIAL</u>	<u>A062569</u>	<u>001</u>	Mar 09, 1988
	DOXY 200				
	+ APP PHARMS	<u>EQ 200MG BASE/VIAL</u>	<u>A062475</u>	<u>002</u>	Dec 09, 1983

SYSTEM, EXTENDED RELEASE; PERIODONTAL

ATRIDOX
+ TOLMAR 50MG

N050751 001 Sep 03, 1998

TABLET; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	COREPHARMA	<u>EQ 20MG BASE</u>	<u>A065182</u>	<u>001</u>	May 13, 2005
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A065163</u>	<u>001</u>	May 13, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062505</u>	<u>001</u>	Sep 11, 1984
<u>AB</u>	LANNETT	<u>EQ 20MG BASE</u>	<u>A065277</u>	<u>001</u>	Nov 10, 2005
<u>AB</u>	MUTUAL PHARM	<u>EQ 100MG BASE</u>	<u>A062677</u>	<u>001</u>	Jul 10, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 151 (of 424)

DOXYCYCLINE HYCLATE

TABLET; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	MUTUAL PHARMA	<u>EQ 20MG BASE</u>	<u>A065134 001</u>	May 13, 2005
<u>AB</u>	VINTAGE PHARMS	<u>EQ 100MG BASE</u>	<u>A062538 001</u>	Apr 07, 1986
<u>AB</u>	WATSON LABS	<u>EQ 100MG BASE</u>	<u>A062421 001</u>	Feb 02, 1983
<u>AB</u>	+ WEST-WARD PHARM CORP	<u>EQ 100MG BASE</u>	<u>A065095 001</u>	Jul 02, 2003
		<u>PERIOSTAT</u>		
<u>AB</u>	+ GALDERMA LABS LP	<u>EQ 20MG BASE</u>	<u>N050783 001</u>	Feb 02, 2001

TABLET, DELAYED RELEASE; ORAL

DORYX

<u>AB</u>	MAYNE PHARMA	<u>EQ 75MG BASE</u>	<u>N050795 001</u>	May 06, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N050795 002</u>	May 06, 2005

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>	IMPAK LABS INC	<u>EQ 75MG BASE</u>	<u>A090505 001</u>	Dec 28, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090505 002</u>	Dec 28, 2010
<u>AB</u>	MYLAN	<u>EQ 75MG BASE</u>	<u>A090431 001</u>	Dec 28, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090431 002</u>	Dec 28, 2010

DORYX

+ MAYNE PHARMA EQ 150MG BASE

N050795 003 Jun 20, 2008

DRONABINOL

CAPSULE; ORAL

DRONABINOL

<u>AB</u>	INSYS THERAP	<u>2.5MG</u>	<u>A078501 001</u>	Aug 19, 2011
<u>AB</u>		<u>5MG</u>	<u>A078501 002</u>	Aug 19, 2011
<u>AB</u>		<u>10MG</u>	<u>A078501 003</u>	Aug 19, 2011
<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

<u>AB</u>		<u>MARINOL</u>		
<u>AB</u>	ABBOTT PRODS	<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

DRONEDARONE HYDROCHLORIDE

TABLET; ORAL

MULTAQ

+ SANOFI AVENTIS US EQ 400MG BASE

N022425 001 Jul 01, 2009

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

<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>AP</u>				
<u>AP</u>	+ AKORN INC	<u>2.5MG/ML</u>	<u>N016796 001</u>	

DROSPIRENONE; ESTRADIOL

TABLET; ORAL

ANGELIQ

+ BAYER HLTHCARE 0.5MG;1MG

N021355 002 Sep 28, 2005

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

<u>AB</u>	BARR	<u>3MG;0.02MG</u>	<u>A078515 001</u>	Mar 30, 2009
-----------	------	-------------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 152 (of 424)

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

<u>AB</u>	WATSON LABS	<u>3MG; 0.02MG</u>	<u>A078833</u> <u>001</u>	Nov 28, 2011
<u>AB</u>	<u>LORYNA</u>			
<u>AB</u>	SANDOZ	<u>3MG; 0.02MG</u>	<u>A079221</u> <u>001</u>	Mar 28, 2011
<u>AB</u>	<u>YAZ</u>			
<u>AB</u> + BAYER HLTHCARE		<u>3MG; 0.02MG</u>	<u>N021676</u> <u>001</u>	Mar 16, 2006
TABLET; ORAL-28				
		<u>DROSPIRENONE AND ETHINYL ESTRADIOL</u>		
<u>AB</u>	BARR	<u>3MG; 0.03MG</u>	<u>A077527</u> <u>001</u>	May 09, 2008
<u>AB</u>	WATSON LABS	<u>3MG; 0.03MG</u>	<u>A090081</u> <u>001</u>	Sep 07, 2010
<u>AB</u>	<u>SYEDA</u>			
<u>AB</u>	SANDOZ	<u>3MG; 0.03MG</u>	<u>A090114</u> <u>001</u>	Mar 28, 2011
<u>AB</u>	<u>YASMIN</u>			
<u>AB</u> + BAYER HLTHCARE		<u>3MG; 0.03MG</u>	<u>N021098</u> <u>001</u>	May 11, 2001

DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET; ORAL

BEYAZ

+ BAYER HLTHCARE	3MG, N/A; 0.02MG, N/A; 0.451MG, 0.451MG	N022532	001	Sep 24, 2010
SAFYRAL				
BAYER HLTHCARE	3MG, N/A; 0.03MG, N/A; 0.451MG, 0.451MG	N022574	001	Dec 16, 2010

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

CYMBALTA

LILLY	EQ 20MG BASE	N021427	001	Aug 03, 2004
	EQ 30MG BASE	N021427	002	Aug 03, 2004
+	EQ 60MG BASE	N021427	004	Aug 03, 2004

DUTASTERIDE

CAPSULE; ORAL

AVODART

<u>AB</u> + GLAXOSMITHKLINE	<u>0.5MG</u>	<u>N021319</u> <u>001</u>	Nov 20, 2001
<u>AB</u>	<u>DUTASTERIDE</u>		
BARR	<u>0.5MG</u>	<u>A090095</u> <u>001</u>	Dec 21, 2010

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

JALYN

+ GLAXOSMITHKLINE	0.5MG; 0.4MG	N022460	001	Jun 14, 2010
-------------------	--------------	---------	-----	--------------

DYPHYLLINE

TABLET; ORAL

LUFYLLIN

MEDA PHARMS	200MG	A084566	001
+	400MG	A084566	002

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

+ WYETH PHARMS INC	0.125%	N011963	001
--------------------	--------	---------	-----

ECONAZOLE NITRATE

CREAM; TOPICAL

ECONAZOLE NITRATE

<u>AB</u> + ALTANA	<u>1%</u>	<u>A076075</u> <u>001</u>	Nov 26, 2002
PERRIGO NEW YORK	<u>1%</u>	<u>A076479</u> <u>001</u>	Jun 23, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 153 (of 424)

ECONAZOLE NITRATE

CREAM; TOPICAL

ECONAZOLE NITRATE

<u>AB</u>	PRASCO	<u>1%</u>	<u>A076574 001</u>	Dec 17, 2004
<u>AB</u>	TARO	<u>1%</u>	<u>A076005 001</u>	Nov 26, 2002

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

+ MEDICIS 200MG/ML

N008922 001

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON

<u>AP</u>	BIONICHE PHARMA	<u>10MG/ML</u>	<u>A088873 001</u>	Aug 06, 1985
<u>AP</u>	+ VALEANT PHARM INTL	<u>10MG/ML</u>	<u>N007959 001</u>	
		<u>TENSTILON PRESERVATIVE FREE</u>		
<u>AP</u>	+ VALEANT PHARM INTL	<u>10MG/ML</u>	<u>N007959 002</u>	

EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

+ BRISTOL MYERS SQUIBB 50MG
200MGN020972 001 Sep 17, 1998
N020972 003 Sep 17, 1998

TABLET; ORAL

SUSTIVA

+ BRISTOL MYERS SQUIBB 600MG

N021360 002 Feb 01, 2002

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

ATRIPLA

+ GILEAD 600MG;200MG;300MG

N021937 001 Jul 12, 2006

EFLORNITHINE HYDROCHLORIDE

CREAM; TOPICAL

VANIQA

+ SKINMEDICA 13.9%

N021145 001 Jul 27, 2000

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

RELPAX

+ PFIZER IRELAND EQ 20MG BASE
EQ 40MG BASEN021016 001 Dec 26, 2002
N021016 002 Dec 26, 2002ELTROMBOPAG OLAMINE

TABLET; ORAL

PROMACTA

+ GLAXOSMITHKLINE EQ 12.5MG ACID
EQ 25MG ACID
EQ 50MG ACID
EQ 75MG ACIDN022291 004 Oct 20, 2011
N022291 001 Nov 20, 2008
N022291 002 Nov 20, 2008
N022291 003 Sep 08, 2009EMEDASTINE DIFUMARATE

SOLUTION/DROPS; OPHTHALMIC

+ ALCON 0.05%

N020706 001 Dec 29, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 154 (of 424)

EMTRICITABINE

CAPSULE; ORAL EMTRIVA + GILEAD	200MG	N021500 001	Jul 02, 2003
SOLUTION; ORAL EMTRIVA + GILEAD	10MG/ML	N021896 001	Sep 28, 2005

EMTRICITABINE; RILPIVIRINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL COMPLERA + GILEAD SCIENCES INC	200MG;25MG;300MG	N202123 001	Aug 10, 2011
---	------------------	-------------	--------------

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL TRUVADA + GILEAD	200MG;300MG	N021752 001	Aug 02, 2004
-------------------------------------	-------------	-------------	--------------

ENALAPRIL MALEATE

<u>ENALAPRIL MALEATE</u>			
<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>	LEK PHARMS	<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>	MYLAN	<u>2.5MG</u>	<u>A075472 001</u> Aug 22, 2000
<u>AB</u>		<u>2.5MG</u>	<u>A075480 001</u> Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075472 002</u> Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075480 002</u> Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075472 003</u> Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075480 003</u> Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075472 004</u> Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075480 004</u> Aug 22, 2000
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>A075621 001</u> Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075621 002</u> Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075621 003</u> Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075621 004</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>	WATSON LABS	<u>2.5MG</u>	<u>A075501 001</u> Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075501 002</u> Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075501 003</u> Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075501 004</u> Aug 22, 2000
<u>AB</u>	WOCKHARDT USA	<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 INTL	<u>2.5MG</u>	<u>N018998 005</u> Jul 26, 1988
<u>AB</u>		<u>5MG</u>	<u>N018998 001</u> Dec 24, 1985

PREScription DRUG PRODUCT LIST

3 - 155 (of 424)

ENALAPRIL MALEATE

TABLET; ORAL

VASOTEC

AB VALEANT INTL 10MG N018998 002 Dec 24, 1985
AB + 20MG N018998 003 Dec 24, 1985

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX INC	<u>5MG;12.5MG</u>	<u>A076486</u>	<u>001</u>	Oct 27, 2004
<u>AB</u>		<u>10MG;25MG</u>	<u>A076486</u>	<u>002</u>	Oct 27, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG;12.5MG</u>	<u>A075909</u>	<u>001</u>	Oct 15, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075909</u>	<u>002</u>	Oct 15, 2001
<u>AB</u>	MYLAN	<u>5MG;12.5MG</u>	<u>A075624</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075624</u>	<u>002</u>	Sep 18, 2001
<u>AB</u>	SANDOZ	<u>5MG;12.5MG</u>	<u>A076116</u>	<u>001</u>	Sep 19, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A076116</u>	<u>002</u>	Sep 19, 2001
<u>AB</u>	TARO PHARM IND'S	<u>5MG;12.5MG</u>	<u>A075788</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075788</u>	<u>002</u>	Sep 18, 2001
<u>AB</u>	TEVA	<u>5MG;12.5MG</u>	<u>A075727</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075727</u>	<u>002</u>	Sep 18, 2001
VASERETIC					
<u>AB</u>	VALEANT INT'L	<u>5MG;12.5MG</u>	<u>N019221</u>	<u>003</u>	Jul 12, 1995
<u>AB</u>	+	<u>10MG;25MG</u>	<u>N019221</u>	<u>001</u>	Oct 31, 1986

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

<u>AP</u>	+	BEDFORD	<u>1.25MG/ML</u>	<u>A075634</u>	<u>001</u>	Aug 22, 2000
<u>AP</u>		HIKMA FARMACEUTICA	<u>1.25MG/ML</u>	<u>A078687</u>	<u>001</u>	Dec 23, 2008
<u>AP</u>	+	HOSPIRA	<u>1.25MG/ML</u>	<u>A075458</u>	<u>001</u>	Aug 22, 2000
<u>AP</u>		TEVA PARENTERAL	<u>1.25MG/ML</u>	<u>A075578</u>	<u>001</u>	Aug 22, 2000

ENFLURANE

LIQUID; INHALATION

ENFLURANE

AN PIRAMAL CRITICAL 99.9% A074396 001 Jul 29, 1994
AN ETHRANE BAXTER HEALTHCARE CORP. 99.9% N017087 001

ENELIVIRTIDE

INJECTABLE: SUBCUTANEOUS

FUZEON

+ BOCHÉ 80MG/VIAL

N021481_001 Mar 13 2003

ENOXAPARIN SODIUM

INJECTABLE: INTRAVENOUS SUBCUTANEOUS

ENOXAPARIN SODIUM

AB ENOXAPARIN SODIUM SANDOZ INC 300MG/3ML (100MG/ML) A078660 001 Nov 28, 2011
AB LOVENOX Sandoz Nutentis US 300MG/3ML (100MG/ML) N000154 000 7-22-2002

INJECTABLE: SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

<u>ENOXAPARIN SODIUM (PRESERVATIVE FREE)</u>					
<u>AP</u>	AMPHASTAR PHARM	30MG/0.3ML (100MG/ML)	<u>A076684</u>	<u>001</u>	Sep 19, 2011
<u>AP</u>		40MG/0.4ML (100MG/ML)	<u>A076684</u>	<u>002</u>	Sep 19, 2011
<u>AP</u>		60MG/0.6ML (100MG/ML)	<u>A076684</u>	<u>003</u>	Sep 19, 2011
<u>AP</u>		80MG/0.8ML (100MG/ML)	<u>A076684</u>	<u>004</u>	Sep 19, 2011
<u>AP</u>		100MG/ML (100MG/ML)	<u>A076684</u>	<u>005</u>	Sep 19, 2011
<u>AP</u>		120MG/0.8ML (150MG/ML)	<u>A076684</u>	<u>006</u>	Sep 19, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 156 (of 424)

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

<u>AP</u>	AMPHASTAR PHARM	<u>150MG/ML (150MG/ML)</u>	<u>A076684</u>	<u>007</u>	Sep 19, 2011
<u>AP</u>	SANDOZ	<u>30MG/0.3ML (100MG/ML)</u>	<u>A077857</u>	<u>002</u>	Jul 23, 2010
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>A077857</u>	<u>003</u>	Jul 23, 2010
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>A077857</u>	<u>004</u>	Jul 23, 2010
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>A077857</u>	<u>005</u>	Jul 23, 2010
<u>AP</u>		<u>100MG/ML (100MG/ML)</u>	<u>A077857</u>	<u>001</u>	Jul 23, 2010
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>A077857</u>	<u>006</u>	Jul 23, 2010
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>A077857</u>	<u>007</u>	Jul 23, 2010
		<u>LOVENOX (PRESERVATIVE FREE)</u>			
<u>AP</u>	SANOFI AVENTIS US	<u>30MG/0.3ML (100MG/ML)</u>	<u>N020164</u>	<u>001</u>	Mar 29, 1993
<u>AP</u>		<u>40MG/0.4ML (100MG/ML)</u>	<u>N020164</u>	<u>002</u>	Jan 30, 1998
<u>AP</u>		<u>60MG/0.6ML (100MG/ML)</u>	<u>N020164</u>	<u>003</u>	Mar 27, 1998
<u>AP</u>		<u>80MG/0.8ML (100MG/ML)</u>	<u>N020164</u>	<u>004</u>	Mar 27, 1998
<u>AP</u>	+	<u>100MG/ML (100MG/ML)</u>	<u>N020164</u>	<u>005</u>	Mar 27, 1998
<u>AP</u>		<u>120MG/0.8ML (150MG/ML)</u>	<u>N020164</u>	<u>007</u>	Jun 02, 2000
<u>AP</u>		<u>150MG/ML (150MG/ML)</u>	<u>N020164</u>	<u>008</u>	Jun 02, 2000

ENTACAPONE

TABLET; ORAL

COMTAN

+ ORION

200MG

N020796 001 Oct 19, 1999

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

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ELESTAT

<u>AT</u>	+ ALLERGAN	<u>0.05%</u>	<u>N021565</u>	<u>001</u>	Oct 16, 2003
	<u>EPINASTINE HYDROCHLORIDE</u>				
<u>AT</u>	APOTEX	<u>0.05%</u>	<u>A090919</u>	<u>001</u>	Oct 31, 2011
<u>AT</u>	CYPRESS PHARM	<u>0.05%</u>	<u>A090870</u>	<u>001</u>	Mar 14, 2011
<u>AT</u>	PHARMAFORCE	<u>0.05%</u>	<u>A090951</u>	<u>001</u>	Oct 31, 2011
<u>AT</u>	SUN PHARM INDs	<u>0.05%</u>	<u>A091626</u>	<u>001</u>	Oct 31, 2011

EPINEPHRINEINJECTABLE; IM-SC
ADRENACLICK

BX	+ SHIONOGI INC	EQ 0.15MG /DELIVERY	N020800	003	Nov 25, 2009
BX	+	EQ 0.3MG /DELIVERY	N020800	004	Nov 25, 2009
	TWINJECT 0.15				
BX	+ SHIONOGI INC	EQ 0.15MG /DELIVERY	N020800	002	May 28, 2004
	TWINJECT 0.3				
BX	+ SHIONOGI INC	EQ 0.3MG /DELIVERY	N020800	001	May 30, 2003

INJECTABLE; INTRAMUSCULAR
EPIPEN

BX	+ MERIDIAN MEDCL TECHN	0.3MG/DELIVERY	N019430	001	Dec 22, 1987
	EPIPEN JR.				

BX + MERIDIAN MEDCL TECHN 0.15MG/DELIVERY

N019430 002 Dec 22, 1987

PRESCRIPTION DRUG PRODUCT LIST

3 - 157 (of 424)

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 <u>CITANEST FORTE DENTAL</u> AP + DENTSPLY PHARM	<u>0.005MG/ML; 4%</u>	<u>N021383 001</u>	
<u>PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE</u> AP SEPTODONT INC	<u>0.005MG/ML; 4%</u>	<u>A078959 001</u>	Aug 30, 2011

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION <u>LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE</u> AP EASTMAN KODAK	<u>0.01MG/ML; 2%</u>	<u>A040057 002</u>	Feb 26, 1993
	<u>0.02MG/ML; 2%</u>	<u>A040057 001</u>	Feb 26, 1993
AP HOSPIRA	<u>0.005MG/ML; 0.5%</u>	<u>A089635 001</u>	Jun 21, 1988
	<u>0.005MG/ML; 1.5%</u>	<u>A088571 001</u>	Sep 13, 1985
	<u>0.005MG/ML; 1.5%</u>	<u>A089645 001</u>	Jun 21, 1988
	<u>0.005MG/ML; 2%</u>	<u>A089651 001</u>	Jun 21, 1988
	<u>0.01MG/ML; 1%</u>	<u>A089644 001</u>	Jun 21, 1988
	<u>0.01MG/ML; 2%</u>	<u>A078772 001</u>	May 12, 2008
	<u>0.01MG/ML; 2%</u>	<u>A089646 001</u>	Jun 21, 1988
	<u>0.02MG/ML; 2%</u>	<u>A078772 002</u>	May 12, 2008
<u>OCTOCAIN</u> AP SEPTODONT	<u>0.01MG/ML; 2%</u>	<u>A084048 001</u>	
	<u>0.02MG/ML; 2%</u>	<u>A084048 002</u>	
<u>XYLOCAINE DENTAL WITH EPINEPHRINE</u> AP + DENTSPLY PHARM	<u>0.01MG/ML; 2%</u>	<u>N021381 001</u>	
	<u>0.02MG/ML; 2%</u>	<u>N021381 002</u>	
<u>XYLOCAINE W/ EPINEPHRINE</u> AP + APP PHARMS	<u>0.005MG/ML; 0.5%</u>	<u>N006488 012</u>	
	<u>0.005MG/ML; 1%</u>	<u>N006488 018</u>	Nov 13, 1986
	<u>0.005MG/ML; 1.5%</u>	<u>N006488 017</u>	
	<u>0.005MG/ML; 2%</u>	<u>N006488 019</u>	Nov 13, 1986
	<u>0.01MG/ML; 1%</u>	<u>N006488 004</u>	
	<u>0.02MG/ML; 2%</u>	<u>N006488 005</u>	

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION <u>ELLENCE</u> AP + PFIZER INC	<u>200MG/100ML (2MG/ML)</u>	<u>N050778 001</u>	Sep 15, 1999
	<u>50MG/25ML (2MG/ML)</u>	<u>N050778 002</u>	Sep 15, 1999
<u>EPIRUBICIN HYDROCHLORIDE</u> AP ACTAVIS TOTOWA	<u>10MG/5ML (2MG/ML)</u>	<u>A065445 001</u>	Sep 18, 2008
	<u>50MG/25ML (2MG/ML)</u>	<u>A065445 002</u>	Sep 18, 2008
	<u>200MG/100ML (2MG/ML)</u>	<u>A065445 003</u>	Sep 18, 2008
AP AKORN INC	<u>50MG/25ML (2MG/ML)</u>	<u>A090163 001</u>	Jun 24, 2009
AP APP PHARMS	<u>10MG/5ML (2MG/ML)</u>	<u>A065408 001</u>	Oct 15, 2007
	<u>150MG/75ML (2MG/ML)</u>	<u>A065408 003</u>	Oct 15, 2007
	<u>200MG/100ML (2MG/ML)</u>	<u>A065408 004</u>	Oct 15, 2007
	<u>50MG/25ML (2MG/ML)</u>	<u>A065408 002</u>	Oct 15, 2007
AP BEDFORD	<u>200MG/100ML (2MG/ML)</u>	<u>A065289 002</u>	Jun 27, 2007
	<u>50MG/25ML (2MG/ML)</u>	<u>A065289 001</u>	Jun 27, 2007
AP BIONICHE PHARMA USA	<u>50MG/25ML (2MG/ML)</u>	<u>A065371 001</u>	Nov 28, 2007
	<u>200MG/100ML (2MG/ML)</u>	<u>A065371 002</u>	Nov 28, 2007
AP EBWEWE PHARMA	<u>50MG/25ML (2MG/ML)</u>	<u>A065339 001</u>	Dec 22, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 158 (of 424)

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE

<u>AP</u>	EBEWE PHARMA	<u>200MG/100ML (2MG/ML)</u>	<u>A065339</u>	<u>002</u>	Dec 22, 2009
<u>AP</u>	FRESENIUS KABI ONCOL	<u>200MG/100ML (2MG/ML)</u>	<u>A065411</u>	<u>001</u>	Aug 20, 2007
<u>AP</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A065411</u>	<u>002</u>	Aug 20, 2007
<u>AP</u>	HOSPIRA	<u>200MG/100ML (2MG/ML)</u>	<u>A065343</u>	<u>004</u>	Apr 19, 2007
<u>AP</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A065343</u>	<u>002</u>	Apr 19, 2007
<u>AP</u>		<u>10MG/5ML (2MG/ML)</u>	<u>A065343</u>	<u>001</u>	Apr 19, 2007
<u>AP</u>		<u>150MG/75ML (2MG/ML)</u>	<u>A065343</u>	<u>003</u>	Apr 19, 2007
<u>AP</u>	MUSTAFA NEVSAT	<u>200MG/100ML (2MG/ML)</u>	<u>A090266</u>	<u>002</u>	Apr 15, 2011
<u>AP</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A090266</u>	<u>001</u>	Apr 15, 2011
<u>AP</u>	TEVA PARENTERAL	<u>50MG/25ML (2MG/ML)</u>	<u>A065331</u>	<u>001</u>	Aug 09, 2007
<u>AP</u>		<u>200MG/100ML (2MG/ML)</u>	<u>A065331</u>	<u>002</u>	Aug 09, 2007
<u>AP</u>	WATSON LABS	<u>200MG/100ML (2MG/ML)</u>	<u>A065361</u>	<u>002</u>	Oct 22, 2007
<u>AP</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A065361</u>	<u>001</u>	Oct 22, 2007
<u>AP</u>	X GEN PHARMS	<u>50MG/25ML (2MG/ML)</u>	<u>A090075</u>	<u>001</u>	Mar 25, 2010
<u>AP</u>		<u>200MG/100ML (2MG/ML)</u>	<u>A090075</u>	<u>002</u>	Mar 25, 2010

INJECTABLE; IV (INFUSION)

EPIRUBICIN HYDROCHLORIDE

+ HOSPIRA 50MG/VIAL

N050807 001 Sep 15, 2006

EPLERENONE

TABLET; ORAL

EPLERENONE

<u>AB</u>	APOTEX	<u>25MG</u>	<u>A078482</u>	<u>001</u>	Jul 30, 2008
<u>AB</u>		<u>50MG</u>	<u>A078482</u>	<u>002</u>	Jul 30, 2008
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A078510</u>	<u>001</u>	Aug 01, 2008
<u>AB</u>		<u>50MG</u>	<u>A078510</u>	<u>002</u>	Aug 01, 2008
	<u>INSPIRA</u>				
<u>AB</u>	GD SEARLE LLC	<u>25MG</u>	<u>N021437</u>	<u>001</u>	Sep 27, 2002
<u>AB</u>	+	<u>50MG</u>	<u>N021437</u>	<u>002</u>	Sep 27, 2002

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUM

<u>AP</u>	TEVA PARENTERAL	<u>EQ 0.5MG BASE/VIAL</u>	<u>A078396</u>	<u>001</u>	Apr 23, 2008
<u>AP</u>		<u>EQ 1.5MG BASE/VIAL</u>	<u>A078396</u>	<u>002</u>	Apr 23, 2008
	<u>FLOLAN</u>				
<u>AP</u>	+ GLAXOSMITHKLINE LLC	<u>EQ 0.5MG BASE/VIAL</u>	<u>N020444</u>	<u>001</u>	Sep 20, 1995
<u>AP</u>	+	<u>EQ 1.5MG BASE/VIAL</u>	<u>N020444</u>	<u>002</u>	Sep 20, 1995
	<u>VELETRI</u>				
<u>AP</u>	+ ACTELION	<u>EQ 1.5MG BASE/VIAL</u>	<u>N022260</u>	<u>001</u>	Jun 27, 2008

EPROSARTAN MESYLATE

TABLET; ORAL

EPROSARTAN MESYLATE

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 400MG BASE</u>	<u>A202012</u>	<u>001</u>	Nov 16, 2011
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A202012</u>	<u>002</u>	Nov 16, 2011
	<u>TEVETEN</u>				
<u>AB</u>	ABBOTT	<u>EQ 400MG BASE</u>	<u>N020738</u>	<u>005</u>	Dec 22, 1997
<u>AB</u>	+	<u>EQ 600MG BASE</u>	<u>N020738</u>	<u>006</u>	May 27, 1999

EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEVETEN HCT

ABBOTT 600MG;12.5MG

+ 600MG;25MG

N021268 001 Nov 01, 2001
N021268 002 Nov 01, 2001

PRESCRIPTION DRUG PRODUCT LIST

3 - 159 (of 424)

EPTIFIBATIDE

INJECTABLE; INJECTION
 INTEGRILIN
 + SCHERING 2MG/ML N020718 001 May 18, 1998
 + 75MG/100ML N020718 002 May 18, 1998

ERGOCALCIFEROL

CAPSULE; ORAL
DRISDOL
AA + SANOFI AVENTIS US 50,000 IU N003444 001
ERGOCALCIFEROL
AA ORIT LABS LLC 50,000 IU A040833 001 May 20, 2009
AA SIGMAPHARM LABS LLC 50,000 IU A091004 001 Jul 14, 2010
AA STRIDES ARCOLAB LTD 50,000 IU A090455 001 Aug 03, 2010
AA SUN PHARM INDs INC 50,000 IU A040865 001 Dec 29, 2009
VITAMIN D
AA BANNER PHARMACAPS 50,000 IU A080704 001

ERGOLOID MESYLATES

TABLET; ORAL
ERGOLOID MESYLATES
AB MUTUAL PHARM 1MG A081113 001 Oct 31, 1991
HYDERGINE
AB + NOVARTIS 1MG N017993 001
 TABLET; SUBLINGUAL
 ERGOLOID MESYLATES
 WATSON LABS 0.5MG A087233 001

ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL
 ERGOMAR
 + ROSEDALE THERAPEUTIC 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)
 + MERCK EQ 1GM BASE/VIAL N021337 001 Nov 21, 2001

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL
ERYC
AB + HOSPIRA 250MG N050536 001
AB WARNER CHILCOTT 250MG A062338 001
ERYTHROMYCIN
AB ARBOR PHARMS INC 250MG A062746 001 Dec 22, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 160 (of 424)

ERYTHROMYCIN

GEL; TOPICAL			
<u>E-GLADES</u>			
AT	COREPHARMA	<u>2%</u>	A065009 001 Mar 18, 2002
<u>ERYGEL</u>			
AT	+ MERZ PHARMS	<u>2%</u>	N050617 001 Oct 21, 1987
<u>ERYTHROMYCIN</u>			
AT	ALTANA	<u>2%</u>	A064184 001 Sep 30, 1997
AT	PERRIGO	<u>2%</u>	A063211 001 Jan 29, 1993
OINTMENT; OPHTHALMIC			
<u>ERYTHROMYCIN</u>			
AT	AKORN	<u>0.5%</u>	A064030 001 Jul 18, 1996
AT	BAUSCH AND LOMB	<u>0.5%</u>	A064067 001 Jul 29, 1994
AT	+ FERA PHARMS	<u>0.5%</u>	A062447 001 Sep 26, 1983
OINTMENT; TOPICAL			
AKNE-MYCIN			
+ DOW PHARM SCIENCES		2%	N050584 001 Jan 10, 1985
SOLUTION; TOPICAL			
<u>ERYTHRA-DERM</u>			
AT	PADDOCK LLC	<u>2%</u>	A062687 001 Feb 05, 1988
<u>ERYTHROMYCIN</u>			
AT	+ FOUGERA PHARMS	<u>2%</u>	A064187 001 Sep 30, 1997
AT	PERRIGO NEW YORK	<u>2%</u>	A063038 001 Jan 11, 1991
AT	WOCKHARDT	<u>2%</u>	A062825 001 Oct 23, 1987
<u>ERYTHRO-STATIN</u>			
AT	HI TECH PHARMA	<u>2%</u>	A064101 001 Oct 22, 1996
SWAB; TOPICAL			
<u>ERYTHROMYCIN</u>			
AT	+ ALTANA	<u>2%</u>	A065320 001 Jul 25, 2006
AT	+ PERRIGO	<u>2%</u>	A064126 001 Jul 03, 1996
AT	VERSAPHARM	<u>2%</u>	A090215 001 May 12, 2010
TABLET; ORAL			
ERYTHROMYCIN			
ARBOR PHARMS INC 250MG			
+ ARBOR PHARMS INC		500MG	A061621 001 A061621 002
TABLET, COATED PARTICLES; ORAL			
PCE			
ARBOR PHARMS INC 333MG			
+ ARBOR PHARMS INC		500MG	N050611 001 Sep 09, 1986 N050611 002 Aug 22, 1990
TABLET, DELAYED RELEASE; ORAL			
ERY-TAB			
+ ARBOR PHARMS INC	250MG		A062298 001
+ ARBOR PHARMS INC	333MG		A062298 003 Mar 29, 1982
+ ARBOR PHARMS INC	500MG		A062298 002

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL			
<u>E.E.S.</u>			
AB	ARBOR PHARMS INC	<u>EQ 200MG BASE/5ML</u>	N050207 001
<u>ERYPED</u>			
AB	ARBOR PHARMS INC	<u>EQ 200MG BASE/5ML</u>	N050207 003 Mar 30, 1987
ERYPED			
+ ARBOR PHARMS INC		EQ 400MG BASE/5ML	N050207 002
SUSPENSION; ORAL			
<u>E.E.S. 200</u>			
AB	ARBOR PHARMS INC	<u>EQ 200MG BASE/5ML</u>	A061639 001
<u>E.E.S. 400</u>			
AB	+ ARBOR PHARMS INC	<u>EQ 400MG BASE/5ML</u>	A061639 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 161 (of 424)

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION; ORAL

PEDIAMYCIN

<u>AB</u>	ARBOR PHARMS INC	<u>EQ 200MG BASE/5ML</u>	<u>A062304 001</u>
	<u>PEDIAMYCIN 400</u>		
<u>AB</u>	ARBOR PHARMS INC	<u>EQ 400MG BASE/5ML</u>	<u>A062304 002</u>
	TABLET; ORAL		
	E.E.S. 400		
BX	+ ARBOR PHARMS INC	EQ 400MG BASE	A061905 002 Aug 12, 1982
	ERYTHROMYCIN ETHYLSUCCINATE		
BX	+ ARBOR PHARMS INC	EQ 400MG BASE	A061904 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
--------	-------------------------------------	--------------------------

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062638 001</u> Oct 31, 1986
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N050609 001</u> Sep 24, 1986
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>A062638 002</u> Oct 31, 1986

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

+ ARBOR PHARMS INC	EQ 250MG BASE	A060359 001
--------------------	---------------	-------------

ESCITALOPRAM OXALATE

SOLUTION; ORAL

LEXAPRO

+ FOREST LABS	EQ 5MG BASE/5ML	N021365 001 Nov 27, 2002
---------------	-----------------	--------------------------

TABLET; ORAL

LEXAPRO

FOREST LABS	EQ 5MG BASE	N021323 001 Aug 14, 2002
	EQ 10MG BASE	N021323 002 Aug 14, 2002
+	EQ 20MG BASE	N021323 003 Aug 14, 2002

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+ BAXTER HLTHCARE CORP	<u>10MG/ML</u>	<u>N019386 006</u> Feb 25, 2003
	<u>ESMOLOL HYDROCHLORIDE</u>		
<u>AP</u>	APP PHARMS	<u>10MG/ML</u>	<u>A076573 001</u> May 02, 2005
<u>AP</u>	BEDFORD LABS	<u>10MG/ML</u>	<u>A076323 001</u> Aug 10, 2004
<u>AP</u>	BIONICHE PHARMA	<u>10MG/ML</u>	<u>A076474 001</u> May 02, 2005
	BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER		
+ BAXTER HLTHCARE CORP	2GM/100ML	N019386 005	Jan 27, 2003
	BREVIBLOC IN PLASTIC CONTAINER		
+ BAXTER HLTHCARE CORP	1GM/100ML	N019386 004	Feb 16, 2001

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

NEXIUM

ASTRAZENECA	EQ 20MG BASE	N021153 001 Feb 20, 2001
+	EQ 40MG BASE	N021153 002 Feb 20, 2001

PRESCRIPTION DRUG PRODUCT LIST

3 - 162 (of 424)

ESOMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL
NEXIUM

ASTRAZENECA	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
VIMOVO

ASTRAZENECA LP	EQ 20MG BASE;375MG	N022511 002	Apr 30, 2010
+	EQ 20MG BASE;500MG	N022511 001	Apr 30, 2010

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS
NEXIUM IV

+	ASTRAZENECA	EQ 20MG BASE/VIAL	N021689 001	Mar 31, 2005
+		EQ 40MG BASE/VIAL	N021689 002	Mar 31, 2005

ESTAZOLAM

TABLET; ORAL

ESTAZOLAM

AB	PAR PHARM	<u>1MG</u>	A074826 001	Jul 03, 1997
AB		<u>2MG</u>	A074826 002	Jul 03, 1997
AB	TEVA	<u>1MG</u>	A074921 001	Jul 10, 1997
AB	+	<u>2MG</u>	A074921 002	Jul 10, 1997
AB	WATSON LABS	<u>1MG</u>	A074818 001	Aug 19, 1997
AB		<u>2MG</u>	A074818 002	Aug 19, 1997

ESTRADIOL

CREAM; VAGINAL
ESTRACE

+	WARNER CHILCOTT	0.01%	A086069 001	Jan 31, 1984
---	-----------------	-------	-------------	--------------

FILM, EXTENDED RELEASE; TRANSDERMAL

CLIMARA

AB	BAYER HLTHCARE	<u>0.0375MG/24HR</u>	N020375 005	May 27, 2003
AB		<u>0.06MG/24HR</u>	N020375 006	May 27, 2003

ESTRADIOL

AB	MYLAN TECHNOLOGIES	<u>0.0375MG/24HR</u>	A075182 004	Jul 20, 2006
AB		<u>0.06MG/24HR</u>	A075182 005	Jul 20, 2006

VIVELLE

AB1	NOVARTIS	<u>0.05MG/24HR</u>	N020323 002	Oct 28, 1994
AB1		<u>0.1MG/24HR</u>	N020323 004	Oct 28, 1994

VIVELLE-DOT

AB1	NOVARTIS	<u>0.05MG/24HR</u>	N020538 006	Jan 08, 1999
AB1	+	<u>0.1MG/24HR</u>	N020538 008	Jan 08, 1999

CLIMARA

AB2	BAYER HLTHCARE	<u>0.025MG/24HR</u>	N020375 004	Mar 05, 1999
AB2		<u>0.05MG/24HR</u>	N020375 001	Dec 22, 1994
AB2		<u>0.075MG/24HR</u>	N020375 003	Mar 23, 1998
AB2	+	<u>0.1MG/24HR</u>	N020375 002	Dec 22, 1994

ESTRADIOL

AB2	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	A075182 003	Jan 26, 2005
AB2		<u>0.05MG/24HR</u>	A075182 006	Feb 24, 2000
AB2		<u>0.075MG/24HR</u>	A075182 002	Jan 26, 2005
AB2	+	<u>0.1MG/24HR</u>	A075182 001	Feb 24, 2000

ALORA

BX	WATSON LABS	0.025MG/24HR	N020655 004	Apr 05, 2002
BX		0.05MG/24HR	N020655 001	Dec 20, 1996
BX		0.075MG/24HR	N020655 002	Dec 20, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 163 (of 424)

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL			
ALORA			
BX	WATSON LABS	0.1MG/24HR	N020655 003 Dec 20, 1996
	ESTRADERM		
BX	NOVARTIS	0.05MG/24HR	N019081 002 Sep 10, 1986
BX	+	0.1MG/24HR	N019081 003 Sep 10, 1986
	VIVELLE-DOT		
BX	NOVARTIS	0.025MG/24HR	N020538 009 May 03, 2002
BX		0.0375MG/24HR	N020538 005 Jan 08, 1999
BX		0.075MG/24HR	N020538 007 Jan 08, 1999
	MENOSTAR		
+	BAYER HLTHCARE	0.014MG/24HR	N021674 001 Jun 08, 2004
GEL; TRANSDERMAL			
	DIVIGEL		
	UPSHER SMITH	0.1% (0.5GM/PACKET)	N022038 002 Jun 04, 2007
		0.1% (0.25GM/PACKET)	N022038 001 Jun 04, 2007
+		0.1% (1GM/PACKET)	N022038 003 Jun 04, 2007
GEL, METERED; TRANSDERMAL			
	ELESTRIN		
+	AZUR PHARMA II	0.06% (0.87GM/ACTIVATION)	N021813 001 Dec 15, 2006
	ESTROGEL		
+	ASCEND	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		
+	KV PHARM	1.53MG/SPRAY	N022014 001 Jul 27, 2007
TABLET; ORAL			
	<u>ESTRACE</u>		
AB	BRISTOL MYERS SQUIBB	<u>1MG</u>	<u>A084499</u> <u>001</u>
AB	+	<u>2MG</u>	<u>A084500</u> <u>001</u>
	<u>ESTRADIOL</u>		
AB	BARR	<u>0.5MG</u>	<u>A040197</u> <u>001</u> Oct 22, 1997
AB		<u>1MG</u>	<u>A040197</u> <u>002</u> Oct 22, 1997
AB		<u>2MG</u>	<u>A040197</u> <u>003</u> Oct 22, 1997
AB	MYLAN	<u>0.5MG</u>	<u>A040326</u> <u>001</u> Apr 21, 1999
AB		<u>1MG</u>	<u>A040326</u> <u>002</u> Apr 21, 1999
AB		<u>2MG</u>	<u>A040326</u> <u>003</u> Apr 21, 1999
AB	USL PHARMA	<u>0.5MG</u>	<u>A040297</u> <u>001</u> Apr 17, 2002
AB		<u>1MG</u>	<u>A040297</u> <u>002</u> Apr 17, 2002
AB		<u>2MG</u>	<u>A040297</u> <u>003</u> Apr 17, 2002
AB	WATSON LABS	<u>0.5MG</u>	<u>A040114</u> <u>003</u> Mar 14, 1996
AB		<u>1MG</u>	<u>A040114</u> <u>001</u> Mar 14, 1996
AB		<u>2MG</u>	<u>A040114</u> <u>002</u> Mar 14, 1996
TABLET; VAGINAL			
	VAGIFEM		
+	NOVO NORDISK INC	10MCG	N020908 002 Nov 25, 2009
<u>ESTRADIOL ACETATE</u>			
INSERT, EXTENDED RELEASE; VAGINAL			
	FEMRING		
	GALEN LTD	EQ 0.05MG BASE/24HR	N021367 001 Mar 20, 2003
+		EQ 0.1MG BASE/24HR	N021367 002 Mar 20, 2003
TABLET; ORAL			
	FEMTRACE		
	WARNER CHILCOTT	0.45MG	N021633 001 Aug 20, 2004
		0.9MG	N021633 002 Aug 20, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 164 (of 424)

ESTRADIOL ACETATE

TABLET; ORAL
 FEMTRACE
 + WARNER CHILCOTT 1.8MG N021633 003 Aug 20, 2004

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION
DEPO-ESTRADOL
AO + PHARMACIA AND UPJOHN 5MG/ML A085470 003
ESTRADIOL CYPIONATE
AO WATSON LABS 5MG/ML A085620 001

ESTRADIOL HEMIHYDRATE

EMULSION; TOPICAL
 ESTRASORB
 + MEDICIS 0.25% N021371 001 Oct 09, 2003

ESTRADIOL VALERATE

INJECTABLE; INJECTION
DELESTROGEN
AO + JHP PHARMS 10MG/ML N009402 002
AO + 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
AO SANDOZ 10MG/ML A040628 001 Oct 04, 2007
AO 20MG/ML A040628 002 Oct 04, 2007
AO 40MG/ML A040628 003 Oct 04, 2007
AO WATSON LABS 20MG/ML A083547 001
AO 40MG/ML A083714 001

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
 COMBIPATCH
 NOVARTIS 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 NOVO NORDISK INC 0.5MG; 0.1MG N020907 002 Dec 28, 2006
AB + 1MG; 0.5MG N020907 001 Nov 18, 1998
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

ESTRADIOL; NORGESTIMATE

TABLET; ORAL
ESTRADIOL AND NORGESTIMATE
AB BARR 1MG, 1MG; N/A, 0.09MG A076812 001 Apr 29, 2005
PREFEST
AB + TEVA WOMENS 1MG, 1MG; N/A, 0.09MG N021040 001 Oct 22, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 165 (of 424)

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
 + 0.625MG N004782 004
 + 0.9MG N004782 005 Jan 26, 1984
 + 1.25MG N004782 001

ESTROGENS, CONJUGATED SYNTHETIC A

CREAM; VAGINAL
 SYNTHETIC CONJUGATED ESTROGENS A
 + TEVA WOMENS 0.625MG/GM N021788 001 Nov 28, 2008

TABLET; ORAL
 CENESTIN
 TEVA WOMENS 0.3MG N020992 001 Jun 21, 2002
 0.45MG N020992 005 Feb 05, 2004
 0.625MG N020992 002 Mar 24, 1999
 0.9MG N020992 003 Mar 24, 1999
 + 1.25MG N020992 004 Mar 13, 2000

ESTROGENS, CONJUGATED SYNTHETIC B

TABLET; ORAL
 ENJUVIA
 TEVA WOMENS 0.3MG N021443 001 Dec 20, 2004
 0.45MG N021443 002 Dec 20, 2004
 0.625MG N021443 003 May 10, 2004
 0.9MG N021443 005 Apr 27, 2007
 + 1.25MG N021443 004 May 10, 2004

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

PRESCRIPTION DRUG PRODUCT LIST

3 - 166 (of 424)

ESTRONE

INJECTABLE; INJECTION
 ESTRONE
 + WATSON LABS 5MG/ML A085239 001

ESTROPIPATE

TABLET; ORAL

ESTROPIPATE

<u>AB</u>	BARR	<u>0.75MG</u>	<u>A040135</u> 001	Nov 27, 1996
<u>AB</u>		<u>1.5MG</u>	<u>A040135</u> 002	Nov 27, 1996
<u>AB</u>		<u>3MG</u>	<u>A040135</u> 003	Nov 27, 1996
<u>AB</u>	MYLAN	<u>0.75MG</u>	<u>A040359</u> 001	Aug 26, 1999
<u>AB</u>		<u>1.5MG</u>	<u>A040359</u> 002	Aug 26, 1999
<u>AB</u>	WATSON LABS	<u>0.75MG</u>	<u>A081213</u> 001	Sep 23, 1993
<u>AB</u>		<u>1.5MG</u>	<u>A081214</u> 001	Sep 23, 1993
<u>AB</u>		<u>3MG</u>	<u>A081215</u> 001	Sep 23, 1993
<u>AB</u>		<u>6MG</u>	<u>A081216</u> 001	Sep 23, 1993
		<u>OGEN .625</u>		
<u>AB</u>	PHARMACIA AND UPJOHN	<u>0.75MG</u>	<u>A083220</u> 001	
		<u>OGEN 1.25</u>		
<u>AB</u>	PHARMACIA AND UPJOHN	<u>1.5MG</u>	<u>A083220</u> 002	
		<u>OGEN 2.5</u>		
<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>3MG</u>	<u>A083220</u> 003	
		<u>OGEN 5</u>		
<u>AB</u>	PHARMACIA AND UPJOHN	<u>6MG</u>	<u>A083220</u> 004	
		<u>ORTHO-EST</u>		
<u>AB</u>	SUN PHARM INDs (IN)	<u>0.75MG</u>	<u>A089567</u> 001	Feb 27, 1991
<u>AB</u>		<u>1.5MG</u>	<u>A089582</u> 001	Jul 17, 1991

ESZOPICLONE

TABLET; ORAL

LUNESTA

	SUNOVION PHARMS INC	1MG	N021476 001	Dec 15, 2004
		2MG	N021476 002	Dec 15, 2004
+		3MG	N021476 003	Dec 15, 2004

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECрин

+ ATON EQ 50MG BASE/VIAL N016093 001

ETHACRYNIC ACID

TABLET; ORAL

EDECрин

+ ATON 25MG N016092 001

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDE

<u>AB</u>	BARR	<u>400MG</u>	<u>A076057</u> 001	Nov 26, 2001
<u>AB</u>	LUPIN	<u>100MG</u>	<u>A078939</u> 001	Jun 17, 2009
<u>AB</u>		<u>400MG</u>	<u>A078939</u> 002	Jun 17, 2009
<u>AB</u>	WEST WARD	<u>100MG</u>	<u>A075095</u> 001	Nov 30, 1999
<u>AB</u>	+	<u>400MG</u>	<u>A075095</u> 002	Nov 30, 1999
		<u>MYAMBUTOL</u>		
<u>AB</u>	STI PHARMA LLC	<u>100MG</u>	<u>N016320</u> 001	
<u>AB</u>		<u>400MG</u>	<u>N016320</u> 003	

PRESCRIPTION DRUG PRODUCT LIST

3 - 167 (of 424)

ETHANOLAMINE OLEATE

INJECTABLE; INJECTION
 ETHAMOLIN
 + QOL MEDCL 50MG/ML N019357 001 Dec 22, 1988

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-28
KELNOR
AB BARR 0.035MG;1MG A076785 001 May 23, 2005
ZOVIA 1/35E-28
AB WATSON LABS 0.035MG;1MG A072721 001 Dec 30, 1991
 ZOVIA 1/50E-28
 + WATSON LABS 0.05MG;1MG A072723 001 Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL
 NUVARING
 + ORGANON USA INC 0.015MG;0.12MG N021187 001 Oct 03, 2001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL
INTROVALE
AB SANDOZ 0.03MG;0.15MG A079064 001 Sep 27, 2010
LEVONORGESTREL AND ETHINYL ESTRADIOL
AB LUPIN LTD 0.02MG,0.01MG;0.1MG,N/A A091674 001 Oct 26, 2011
AB VINTAGE PHARMS 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG A077502 001 Nov 23, 2011
AB WATSON LABS 0.02MG;0.09MG A079218 001 Jun 06, 2011
AB 0.02MG,0.01MG;0.1MG,N/A A200407 001 Oct 25, 2011
LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL
AB WATSON LABS 0.03MG,0.01MG;0.15MG,N/A A078834 001 May 31, 2011
LOSEASONIQUE
AB TEVA WOMENS 0.02MG,0.01MG;0.1MG,N/A N022262 001 Oct 24, 2008
LYBREL
AB + WYETH PHARMS INC 0.02MG;0.09MG N021864 001 May 22, 2007
QUASENSE
AB WATSON LABS 0.03MG;0.15MG A077101 001 Sep 06, 2006
SEASONALE
AB + DURAMED RES 0.03MG;0.15MG N021544 001 Sep 05, 2003
SEASONIQUE
AB + TEVA WOMENS 0.03MG,0.01MG;0.15MG,N/A N021840 001 May 25, 2006
 TABLET; ORAL-28
ALTAVERA
AB SANDOZ 0.03MG;0.15MG A079102 001 Aug 03, 2010
ENPRESSE-28
AB DURAMED PHARMS BARR 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG A075809 002 Jul 16, 2001
LEVONEST
AB NOVAST LABS LTD 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG A090719 001 Dec 29, 2010
LEVORA 0.15/30-28
AB WATSON LABS 0.03MG;0.15MG A073594 001 Dec 13, 1993
NORDETTE-28
AB + DURAMED 0.03MG;0.15MG N018782 001 Jul 21, 1982
PORTIA-28
AB BARR 0.03MG;0.15MG A075866 002 May 23, 2002
TRIVORA-28
AB + WATSON LABS 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG A074538 002 Dec 18, 1997
AVIANE-28
AB1 DURAMED PHARMS BARR 0.02MG;0.1MG A075796 001 Apr 30, 2001

PRESCRIPTION DRUG PRODUCT LIST

3 - 168 (of 424)

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

LEVONORGESTREL AND ETHINYL ESTRADIOL

<u>AB1</u> + WATSON LABS	<u>0.02MG;0.1MG</u>	<u>A076625</u> <u>001</u>	Nov 18, 2004
<u>ORSYTHIA</u>			
<u>AB1</u> VINTAGE PHARMS	<u>0.02MG;0.1MG</u>	<u>A077099</u> <u>001</u>	May 11, 2011
<u>LESSINA-28</u>			
<u>AB2</u> BARR	<u>0.02MG;0.1MG</u>	<u>A075803</u> <u>002</u>	Mar 20, 2002
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>			
<u>AB2</u> + WATSON LABS	<u>0.02MG;0.1MG</u>	<u>A077681</u> <u>001</u>	May 31, 2006

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

ORTHO EVRA

+ JANSSEN PHARMS	0.02MG/24HR;0.15MG/24HR	N021180 001	Nov 20, 2001
------------------	-------------------------	-------------	--------------

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORETHIN 1/35E-21

<u>AB</u> WATSON LABS	<u>0.035MG;1MG</u>	<u>A071480</u> <u>001</u>	Apr 12, 1988
<u>AB</u> WATSON LABS	<u>0.035MG;1MG</u>	<u>A070685</u> <u>001</u>	Jan 29, 1987
<u>NORINYL 1+35 21-DAY</u>			
<u>AB</u> WATSON LABS	<u>0.035MG;1MG</u>	<u>N017565</u> <u>001</u>	
<u>NORTREL 1/35-21</u>			
<u>AB</u> BARR	<u>0.035MG;1MG</u>	<u>A072693</u> <u>001</u>	Feb 28, 1992
NORETHINDRONE AND ETHINYL ESTRADIOL			
WATSON LABS	0.035MG;0.4MG	A078379 001	Feb 23, 2010
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	0.035MG,0.035MG;0.5MG,1MG	A071041 001	Sep 24, 1991
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

ARANELLE

<u>AB</u> BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG</u>	<u>A076783</u> <u>001</u>	Sep 29, 2004
<u>AB</u> BARR	<u>0.035MG;0.4MG</u>	<u>A076238</u> <u>001</u>	Apr 22, 2004
<u>BREVICON 28-DAY</u>			
<u>AB</u> WATSON LABS	<u>0.035MG;0.5MG</u>	<u>N017743</u> <u>001</u>	
<u>BRIELLYN</u>			
<u>AB</u> GLENMARK GENERICS	<u>0.035MG;0.4MG</u>	<u>A090538</u> <u>001</u>	Mar 22, 2011
<u>CYCLAFEM 1/35</u>			
<u>AB</u> VINTAGE	<u>0.035MG;1MG</u>	<u>A076337</u> <u>001</u>	Nov 12, 2010
<u>CYCLAFEM 7/7/7</u>			
<u>AB</u> VINTAGE	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A076338</u> <u>001</u>	Nov 16, 2010
G			
<u>DASETTA 1/35</u>			
<u>AB</u> NOVAST LABS LTD	<u>0.035MG;1MG</u>	<u>A090948</u> <u>001</u>	Dec 22, 2011
<u>DASETTA 7/7/7</u>			
<u>AB</u> NOVAST LABS LTD	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>A090946</u> <u>001</u>	Dec 22, 2011
G			
<u>MODICON 28</u>			
<u>AB</u> + JANSSEN PHARMS	<u>0.035MG;0.5MG</u>	<u>N017735</u> <u>001</u>	
<u>NORETHIN 1/35E-28</u>			
<u>AB</u> WATSON LABS	<u>0.035MG;1MG</u>	<u>A071481</u> <u>001</u>	Apr 12, 1988
<u>NORETHINDRONE AND ETHINYL ESTRADIOL</u>			
<u>AB</u> WATSON LABS	<u>0.035MG;0.4MG</u>	<u>A078323</u> <u>001</u>	Feb 04, 2010
	<u>0.035MG;0.5MG</u>	<u>A070686</u> <u>001</u>	Jan 29, 1987

PRESCRIPTION DRUG PRODUCT LIST

3 - 169 (of 424)

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

NORETHINDRONE AND ETHINYL ESTRADIOL

<u>AB</u>	WATSON LABS	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1M</u>	<u>A076393</u> <u>001</u>	Feb 04, 2010
<u>AB</u>		<u>G</u>		
<u>AB</u>		<u>0.035MG; 1MG</u>	<u>A070687</u> <u>001</u>	Jan 29, 1987
	<u>NORINYL 1+35 28-DAY</u>			
<u>AB</u>	WATSON LABS	<u>0.035MG; 1MG</u>	<u>N017565</u> <u>002</u>	
	<u>NORTREL 0.5/35-28</u>			
<u>AB</u>	BARR	<u>0.035MG; 0.5MG</u>	<u>A072695</u> <u>001</u>	Feb 28, 1992
	<u>NORTREL 1/35-28</u>			
<u>AB</u>	BARR	<u>0.035MG; 1MG</u>	<u>A072696</u> <u>001</u>	Feb 28, 1992
	<u>NORTREL 7/7/7</u>			
<u>AB</u>	BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1M</u>	<u>A075478</u> <u>002</u>	Aug 30, 2002
		<u>G</u>		
	<u>ORTHO-NOVUM 1/35-28</u>			
<u>AB</u>	+ JANSSEN PHARMS	<u>0.035MG; 1MG</u>	<u>N017919</u> <u>002</u>	
	<u>ORTHO-NOVUM 7/7/7-28</u>			
<u>AB</u>	+ JANSSEN PHARMS	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1M</u>	<u>N018985</u> <u>002</u>	Apr 04, 1984
		<u>G</u>		
	<u>OVCON-35</u>			
<u>AB</u>	+ WARNER CHILCOTT	<u>0.035MG; 0.4MG</u>	<u>N017716</u> <u>001</u>	
	<u>PHILITH</u>			
<u>AB</u>	NOVAST LABS LTD	<u>0.035MG; 0.4MG</u>	<u>A090947</u> <u>001</u>	Dec 22, 2011
	<u>TRI-NORINYL 28-DAY</u>			
<u>AB</u>	+ WATSON LABS	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MG</u>	<u>N018977</u> <u>002</u>	Apr 13, 1984
	NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)			
	WATSON LABS	0.035MG, 0.035MG; 0.5MG, 1MG	A071044 001	Apr 01, 1988
	NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)			
	WATSON LABS	0.035MG, 0.035MG; 0.5MG, 1MG	A071042 001	Sep 24, 1991
	OVCON-50			
	+ WARNER CHILCOTT	0.05MG; 1MG	N017576 001	
	TABLET, CHEWABLE; ORAL			
	<u>FEMCON FE</u>			
<u>AB</u>	+ WARNER CHILCOTT	<u>0.035MG; 0.4MG</u>	<u>N021490</u> <u>001</u>	Nov 14, 2003
	<u>NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>			
<u>AB</u>	BARR	<u>0.035MG; 0.4MG</u>	<u>A078965</u> <u>001</u>	Aug 05, 2010
<u>AB</u>	WATSON LABS	<u>0.035MG; 0.4MG</u>	<u>A078892</u> <u>001</u>	Sep 26, 2011
	NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE			
	+ WATSON LABS INC	0.025MG; 0.8MG	N022573 001	Dec 22, 2010

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

FEMHRT

<u>AB</u>	+ WARNER CHILCOTT	<u>0.005MG; 1MG</u>	<u>N021065</u> <u>002</u>	Oct 15, 1999
	<u>LOESTRIN 24 FE</u>			
<u>AB</u>	+ WARNER CHILCOTT	<u>0.02MG; 1MG</u>	<u>N021871</u> <u>001</u>	Feb 17, 2006
	<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>			
<u>AB</u>	BARR	<u>0.005MG; 1MG</u>	<u>A076221</u> <u>001</u>	Nov 06, 2009
	<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>			
<u>AB</u>	WATSON LABS	<u>0.02MG; 1MG</u>	<u>A078267</u> <u>001</u>	Sep 01, 2009
	FEMHRT			
	WARNER CHILCOTT	0.0025MG; 0.5MG	N021065 001	Jan 14, 2005
	LO LOESTRIN FE			
	+ WARNER CHILCOTT CO	0.01MG, 0.01MG; 1MG, N/A	N022501 001	Oct 21, 2010

TABLET; ORAL-21

JUNEL 1.5/30

<u>AB</u>	BARR	<u>0.03MG; 1.5MG</u>	<u>A076381</u> <u>001</u>	May 30, 2003
	<u>JUNEL 1/20</u>			
<u>AB</u>	BARR	<u>0.02MG; 1MG</u>	<u>A076380</u> <u>001</u>	May 30, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 170 (of 424)

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-21			
<u>LOESTRIN 21 1.5/30</u>			
AB WARNER CHILCOTT	<u>0.03MG;1.5MG</u>	<u>N017875</u>	<u>001</u>
<u>LOESTRIN 21 1/20</u>			
AB WARNER CHILCOTT	<u>0.02MG;1MG</u>	<u>N017876</u>	<u>001</u>
<u>MICROGESTIN 1.5/30</u>			
AB WATSON LABS	<u>0.03MG;1.5MG</u>	<u>A075548</u>	<u>002</u>
<u>MICROGESTIN 1/20</u>			Jul 30, 2003
AB WATSON LABS	<u>0.02MG;1MG</u>	<u>A075647</u>	<u>002</u>
TRI-LEGEST 21 BARR	0.02MG, 0.03MG, 0.035MG;1MG, 1MG, 1MG	A076405	001
Oct 26, 2007			
TABLET; ORAL-28			
<u>ESTROSTEP FE</u>			
AB + WARNER CHILCOTT	<u>0.02MG, 0.03MG, 0.035MG;1MG, 1MG, 1MG</u>	<u>N020130</u>	<u>002</u>
<u>GILDESS FE 1.5/30</u>			
AB VINTAGE	<u>0.03MG;1.5MG</u>	<u>A077075</u>	<u>001</u>
<u>GILDESS FE 1/20</u>			Apr 28, 2005
AB VINTAGE	<u>0.02MG;1MG</u>	<u>A077077</u>	<u>001</u>
<u>JUNEL FE 1.5/30</u>			May 20, 2005
AB BARR	<u>0.03MG;1.5MG</u>	<u>A076064</u>	<u>001</u>
<u>JUNEL FE 1/20</u>			Sep 18, 2003
AB BARR	<u>0.02MG;1MG</u>	<u>A076081</u>	<u>001</u>
<u>LOESTRIN FE 1.5/30</u>			Sep 18, 2003
AB + WARNER CHILCOTT	<u>0.03MG;1.5MG</u>	<u>N017355</u>	<u>001</u>
<u>LOESTRIN FE 1/20</u>			
AB WARNER CHILCOTT	<u>0.02MG;1MG</u>	<u>N017354</u>	<u>001</u>
<u>MICROGESTIN FE 1.5/30</u>			
AB WATSON LABS	<u>0.03MG;1.5MG</u>	<u>A075548</u>	<u>001</u>
<u>MICROGESTIN FE 1/20</u>			Feb 05, 2001
AB WATSON LABS	<u>0.02MG;1MG</u>	<u>A075647</u>	<u>001</u>
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>			
AB WATSON LABS	<u>0.02MG, 0.03MG, 0.035MG;1MG, 1MG, 1MG</u>	<u>A076629</u>	<u>001</u>
<u>TRI-LEGEST FE</u>			Mar 18, 2010
AB BARR	<u>0.02MG, 0.03MG, 0.035MG;1MG, 1MG, 1MG</u>	<u>A076105</u>	<u>001</u>
Oct 26, 2007			

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL			
<u>NORGESTIMATE AND ETHINYL ESTRADIOL</u>			
AB GLENMARK GENERICS	<u>0.035MG, 0.035MG, 0.035MG;0.18MG, 0.215MG, 0.25MG</u>	<u>A200494</u>	<u>001</u>
Jun 17, 2011			
TABLET; ORAL-28			
<u>NORGESTIMATE AND ETHINYL ESTRADIOL</u>			
AB WATSON LABS	<u>0.025MG, 0.025MG, 0.025MG;0.18MG, 0.215MG, 0.25MG</u>	<u>A090479</u>	<u>001</u>
Mar 09, 2011			
AB	<u>0.035MG, 0.035MG, 0.035MG;0.18MG, 0.215MG, 0.25MG</u>	<u>A076626</u>	<u>001</u>
Aug 17, 2006			
AB	<u>0.035MG;0.25MG</u>	<u>A076627</u>	<u>001</u>
Aug 17, 2006			
<u>ORTHO CYCLEN-28</u>			
AB + JANSSEN PHARMS	<u>0.035MG;0.25MG</u>	<u>N019653</u>	<u>002</u>
<u>ORTHO TRI-CYCLEN</u>			Dec 29, 1989
AB + JANSSEN PHARMS	<u>0.035MG, 0.035MG, 0.035MG;0.18MG, 0.215MG, 0.25MG</u>	<u>N019697</u>	<u>001</u>
Jul 03, 1992			
<u>ORTHO TRI-CYCLEN LO</u>			
AB + JANSSEN PHARMS	<u>0.025MG, 0.025MG, 0.025MG;0.18MG, 0.215MG, 0.25MG</u>	<u>N021241</u>	<u>001</u>
Aug 22, 2002			
<u>PREVIFEM</u>			
AB VINTAGE	<u>0.035MG;0.25MG</u>	<u>A076334</u>	<u>001</u>
Jan 09, 2004			
<u>SPRINTEC</u>			
AB BARR	<u>0.035MG;0.25MG</u>	<u>A075804</u>	<u>001</u>
Sep 25, 2002			

PRESCRIPTION DRUG PRODUCT LIST

3 - 171 (of 424)

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

TRI LO SPRINTEC

<u>AB</u>	BARR	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076784 001</u>	Jun 29, 2009
<u>AB</u>	VINTAGE	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076335 001</u>	Mar 26, 2004
<u>AB</u>	<u>TRI-SPRINTEC</u>	<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>AB</u>	+ WATSON LABS	<u>0.03MG; 0.3MG</u>	<u>A075288 001</u>	Jul 28, 1999

TABLET; ORAL-28

CRYSELLE

<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 002</u>	Nov 30, 2001
<u>AB</u>	<u>LO/OVRAL-28</u>	<u>0.03MG; 0.3MG</u>	<u>N017802 001</u>	
<u>AB</u>	WYETH PHARMS	<u>0.03MG; 0.3MG</u>		
<u>AB</u>	<u>LOW-OGESTREL-28</u>	<u>0.03MG; 0.3MG</u>	<u>A075288 002</u>	Jul 28, 1999
<u>AB</u>	WATSON LABS	<u>0.03MG; 0.3MG</u>		
	OGESTREL 0.5/50-28			
+ WATSON LABS		0.05MG; 0.5MG	A075406 002	Dec 15, 1999

ETHIONAMIDE

TABLET; ORAL

TRECATOR

+ WYETH PHARMS INC 250MG

N013026 002

ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDE

<u>AB</u>	BANNER PHARMACAPS	<u>250MG</u>	<u>A040430 001</u>	Oct 28, 2002
<u>AB</u>	VERSAPHARM	<u>250MG</u>	<u>A040686 001</u>	May 28, 2008
<u>AB</u>	<u>ZARONTIN</u>	<u>250MG</u>	<u>N012380 001</u>	

SYRUP; ORAL

ETHOSUXIMIDE

<u>AA</u>	MIKART	<u>250MG/5ML</u>	<u>A040506 001</u>	Dec 22, 2003
<u>AA</u>	PHARM ASSOC	<u>250MG/5ML</u>	<u>A040253 001</u>	Nov 22, 2000
<u>AA</u>	TEVA PHARMS	<u>250MG/5ML</u>	<u>A081306 001</u>	Jul 30, 1993
<u>AB</u>	<u>ZARONTIN</u>	<u>250MG/5ML</u>	<u>A080258 001</u>	
<u>AA</u>	+ PARKE DAVIS	<u>250MG/5ML</u>		

ETHOTOIN

TABLET; ORAL

PEGANONE

+ LUNDBECK INC 250MG

N010841 001

ETIDRONATE DISODIUM

TABLET; ORAL

DIDRONEL

<u>AB</u>	PROCTER AND GAMBLE	<u>200MG</u>	<u>N017831 001</u>	
<u>AB</u>	+	<u>400MG</u>	<u>N017831 002</u>	
<u>AB</u>	<u>ETIDRONATE DISODIUM</u>	<u>200MG</u>	<u>A075800 001</u>	Jan 24, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 172 (of 424)

ETIDRONATE DISODIUM

TABLET; ORAL

ETIDRONATE DISODIUM

<u>AB</u>	MYLAN	<u>400MG</u>	<u>A075800</u>	<u>002</u>	Jan 24, 2003
-----------	-------	--------------	----------------	------------	--------------

ETODOLAC

CAPSULE; ORAL

ETODOLAC

<u>AB</u>	APOTEX	<u>200MG</u>	<u>A075419</u>	<u>001</u>	Jul 28, 2000
<u>AB</u>		<u>300MG</u>	<u>A075419</u>	<u>002</u>	Jul 28, 2000
<u>AB</u>	TARO	<u>200MG</u>	<u>A075078</u>	<u>001</u>	Apr 30, 1998
<u>AB</u>	+	<u>300MG</u>	<u>A075078</u>	<u>002</u>	Apr 30, 1998
<u>AB</u>	TEVA	<u>300MG</u>	<u>A075126</u>	<u>002</u>	Sep 16, 1999

TABLET; ORAL

ETODOLAC

<u>AB</u>	APOTEX INC	<u>400MG</u>	<u>A076004</u>	<u>001</u>	Dec 03, 2002
<u>AB</u>		<u>500MG</u>	<u>A076004</u>	<u>002</u>	Dec 03, 2002
<u>AB</u>	MYLAN	<u>400MG</u>	<u>A075104</u>	<u>001</u>	Feb 06, 1998
<u>AB</u>		<u>500MG</u>	<u>A075104</u>	<u>002</u>	Nov 20, 1998
<u>AB</u>	SANDOZ	<u>400MG</u>	<u>A074903</u>	<u>001</u>	Apr 11, 1997
<u>AB</u>		<u>500MG</u>	<u>A074903</u>	<u>002</u>	Apr 19, 1999
<u>AB</u>	TARO PHARM IND	<u>400MG</u>	<u>A075074</u>	<u>001</u>	Mar 11, 1998
<u>AB</u>	+	<u>500MG</u>	<u>A075074</u>	<u>002</u>	Apr 25, 2000
<u>AB</u>	TEVA	<u>400MG</u>	<u>A075009</u>	<u>001</u>	Nov 26, 1997
<u>AB</u>		<u>500MG</u>	<u>A075009</u>	<u>002</u>	Dec 28, 1999

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

<u>AB</u>	TARO	<u>400MG</u>	<u>A076174</u>	<u>001</u>	Mar 13, 2003
<u>AB</u>		<u>500MG</u>	<u>A076174</u>	<u>002</u>	Mar 13, 2003
<u>AB</u>		<u>600MG</u>	<u>A076174</u>	<u>003</u>	Mar 13, 2003
<u>AB</u>	TEVA	<u>400MG</u>	<u>A075665</u>	<u>003</u>	Feb 05, 2001
<u>AB</u>		<u>500MG</u>	<u>A075665</u>	<u>002</u>	Jul 31, 2000
<u>AB</u>	+	<u>600MG</u>	<u>A075665</u>	<u>001</u>	Jul 31, 2000

ETOMIDATE

INJECTABLE; INJECTION

AMIDATE

<u>AP</u>	+	HOSPIRA	<u>2MG/ML</u>	<u>N018227</u>	<u>001</u>	Sep 07, 1982
		<u>ETOMIDATE</u>				
<u>AP</u>		BEDFORD	<u>2MG/ML</u>	<u>A074593</u>	<u>001</u>	Nov 04, 1996
<u>AP</u>		LUITPOLD	<u>2MG/ML</u>	<u>A078867</u>	<u>001</u>	Dec 22, 2009
<u>AP</u>		STRIDES ARCOLAB LTD	<u>2MG/ML</u>	<u>A078289</u>	<u>001</u>	Jan 02, 2009

ETONOGESTREL

IMPLANT; IMPLANTATION

IMPLANON

+ ORGANON USA INC	68MG/IMPLANT	<u>N021529</u>	<u>001</u>	Jul 17, 2006
NEXPLANON				
+ ORGANON USA INC	68MG/IMPLANT	<u>N021529</u>	<u>002</u>	May 31, 2011

ETOPOSIDE

CAPSULE; ORAL

ETOPOSIDE

+ MYLAN	50MG	<u>A075635</u>	<u>001</u>	Sep 19, 2001
---------	------	----------------	------------	--------------

INJECTABLE; INJECTION

ETOPOSIDE

<u>AP</u>	ACCORD HLTHCARE INC	<u>20MG/ML</u>	<u>A074513</u>	<u>001</u>	Mar 14, 1996	
<u>AP</u>	APP PHARMS	<u>20MG/ML</u>	<u>A074983</u>	<u>001</u>	Sep 30, 1998	
<u>AP</u>	+	<u>BEDFORD</u>	<u>20MG/ML</u>	<u>A074290</u>	<u>001</u>	Jul 17, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 173 (of 424)

ETOPOSIDE

INJECTABLE; INJECTION

ETOPOSIDE

<u>AP</u>	PHARMACHEMIE	<u>20MG/ML</u>	<u>A074227 001</u>	Feb 22, 1996
<u>AP</u>	TEVA PARENTERAL	<u>20MG/ML</u>	<u>A074284 001</u>	Feb 10, 1994
<u>AP</u>		<u>20MG/ML</u>	<u>A074529 001</u>	Jul 24, 1996

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

+ BRISTOL MYERS SQUIBB EQ 100MG BASE/VIAL

N020457 001 May 17, 1996

ETRAVIRINE

TABLET; ORAL

INTELENCE

TIBOTEC

100MG

N022187 001 Jan 18, 2008

+ 200MG

N022187 002 Dec 22, 2010

EVEROLIMUS

TABLET; ORAL

AFINITOR

NOVARTIS

2.5MG

N022334 003 Jul 09, 2010

5MG

N022334 001 Mar 30, 2009

+ 10MG

N022334 002 Mar 30, 2009

ZORTRESS

NOVARTIS

0.25MG

N021560 001 Apr 20, 2010

0.5MG

N021560 002 Apr 20, 2010

+ 0.75MG

N021560 003 Apr 20, 2010

EXEMESTANE

TABLET; ORAL

AROMASINAB + PHARMACIA AND UPJOHN 25MGN020753 001 Oct 21, 1999EXEMESTANEAB ROXANE 25MGA077431 001 Apr 01, 2011EXENATIDE SYNTHETIC

INJECTABLE; SUBCUTANEOUS

BYETTA

+ AMYLIN 300MCG/1.2ML (250MCG/ML)

N021773 001 Apr 28, 2005

+ 600MCG/2.4ML (250MCG/ML)

N021773 002 Apr 28, 2005

EZETIMIBE

TABLET; ORAL

ZETIA

+ MSP SINGAPORE 10MG

N021445 001 Oct 25, 2002

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

VYTORIN

MSD INTL

10MG;10MG

N021687 001 Jul 23, 2004

10MG;20MG

N021687 002 Jul 23, 2004

10MG;40MG

N021687 003 Jul 23, 2004

+ 10MG;80MG

N021687 004 Jul 23, 2004

EZOGABINE

TABLET; ORAL

POTIGA

VALEANT PHARMS

50MG

N022345 001 Jun 10, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 174 (of 424)

EZOGABINE

TABLET; ORAL POTIGA				
VALEANT PHARMS	200MG	N022345 002	Jun 10, 2011	
	300MG	N022345 003	Jun 10, 2011	
+	400MG	N022345 004	Jun 10, 2011	

FAMCICLOVIR

TABLET; ORAL FAMCICLOVIR				
AB APOTEX	<u>125MG</u>	<u>A091480 001</u>	Jul 22, 2011	
AB	<u>250MG</u>	<u>A091480 002</u>	Jul 22, 2011	
AB	<u>500MG</u>	<u>A091480 003</u>	Jul 22, 2011	
AB AUROBINDO PHARMA LTD	<u>125MG</u>	<u>A091114 001</u>	Mar 21, 2011	
AB	<u>250MG</u>	<u>A091114 002</u>	Mar 21, 2011	
AB	<u>500MG</u>	<u>A091114 003</u>	Mar 21, 2011	
AB MYLAN	<u>125MG</u>	<u>A201333 001</u>	Mar 24, 2011	
AB	<u>250MG</u>	<u>A201333 002</u>	Mar 24, 2011	
AB	<u>500MG</u>	<u>A201333 003</u>	Mar 24, 2011	
AB ROXANE	<u>125MG</u>	<u>A090128 001</u>	Mar 21, 2011	
AB	<u>250MG</u>	<u>A090128 002</u>	Mar 21, 2011	
AB	<u>500MG</u>	<u>A090128 003</u>	Mar 21, 2011	
AB TEVA PHARMS	<u>125MG</u>	<u>A077487 001</u>	Aug 24, 2007	
AB	<u>250MG</u>	<u>A077487 002</u>	Aug 24, 2007	
AB	<u>500MG</u>	<u>A077487 003</u>	Aug 24, 2007	
AB WATSON LABS	<u>125MG</u>	<u>A078278 001</u>	Mar 21, 2011	
AB	<u>250MG</u>	<u>A078278 002</u>	Mar 21, 2011	
AB	<u>500MG</u>	<u>A078278 003</u>	Mar 21, 2011	
FAMVIR				
AB NOVARTIS	<u>125MG</u>	<u>N020363 003</u>	Dec 11, 1995	
AB	<u>250MG</u>	<u>N020363 001</u>	Apr 26, 1996	
AB +	<u>500MG</u>	<u>N020363 002</u>	Jun 29, 1994	

FAMOTIDINE

FOR SUSPENSION; ORAL FAMOTIDINE				
AB LUPIN LTD	<u>40MG/5ML</u>	<u>A090440 001</u>	Jun 29, 2010	
AB NAVINTA LLC	<u>40MG/5ML</u>	<u>A091020 001</u>	May 27, 2010	
PEPCID				
AB + SALIX PHARMS	<u>40MG/5ML</u>	<u>N019527 001</u>	Feb 02, 1987	
INJECTABLE; INJECTION FAMOTIDINE				
AP APP PHARMS	<u>10MG/ML</u>	<u>A075709 001</u>	Apr 16, 2001	
AP + BAXTER HLTHCARE	<u>10MG/ML</u>	<u>A075488 001</u>	Apr 16, 2001	
AP +	<u>10MG/ML</u>	<u>A075799 001</u>	Apr 30, 2002	
AP BEDFORD	<u>10MG/ML</u>	<u>A075651 001</u>	Apr 16, 2001	
AP	<u>10MG/ML</u>	<u>A075684 001</u>	Apr 16, 2001	
AP PFIZER	<u>10MG/ML</u>	<u>A078641 001</u>	Jun 25, 2008	
FAMOTIDINE PRESERVATIVE FREE				
AP APP PHARMS	<u>10MG/ML</u>	<u>A075813 001</u>	Apr 16, 2001	
AP + BAXTER HLTHCARE	<u>10MG/ML</u>	<u>A075486 001</u>	Apr 16, 2001	
AP +	<u>10MG/ML</u>	<u>A075789 001</u>	Apr 30, 2002	
AP BEDFORD	<u>10MG/ML</u>	<u>A075622 001</u>	Apr 16, 2001	
AP BEN VENUE	<u>10MG/ML</u>	<u>A075825 001</u>	Apr 17, 2001	
AP CLARIS LIFESCIENCES	<u>10MG/ML</u>	<u>A076324 001</u>	Nov 27, 2002	
AP PFIZER	<u>10MG/ML</u>	<u>A078642 001</u>	Jun 25, 2008	
FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)				
AP CLARIS LIFESCIENCES	<u>10MG/ML</u>	<u>A076322 001</u>	Nov 27, 2002	
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER				
AP + BAXTER HLTHCARE	<u>0.4MG/ML</u>	<u>A075591 001</u>	May 10, 2001	

PRESCRIPTION DRUG PRODUCT LIST

3 - 175 (of 424)

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>20MG</u>	<u>A078916</u>	<u>001</u>	May 22, 2009
<u>AB</u>		<u>40MG</u>	<u>A078916</u>	<u>002</u>	May 22, 2009
<u>AB</u>	APOTEX	<u>20MG</u>	<u>A075611</u>	<u>001</u>	Jul 23, 2001
<u>AB</u>		<u>40MG</u>	<u>A075611</u>	<u>002</u>	Jul 23, 2001
<u>AB</u>	CARLSBAD	<u>20MG</u>	<u>A075805</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075805</u>	<u>002</u>	Apr 16, 2001
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A075718</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075718</u>	<u>002</u>	Apr 16, 2001
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>20MG</u>	<u>A075511</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075511</u>	<u>002</u>	Apr 16, 2001
<u>AB</u>	MYLAN	<u>20MG</u>	<u>A075457</u>	<u>001</u>	Apr 18, 2001
<u>AB</u>		<u>40MG</u>	<u>A075704</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075457</u>	<u>002</u>	Apr 18, 2001
<u>AB</u>		<u>40MG</u>	<u>A075704</u>	<u>002</u>	Apr 16, 2001
<u>AB</u>	PERRIGO	<u>20MG</u>	<u>A077352</u>	<u>002</u>	Jul 27, 2005
<u>AB</u>		<u>40MG</u>	<u>A077352</u>	<u>001</u>	Jul 27, 2005
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>A075607</u>	<u>001</u>	May 10, 2001
<u>AB</u>		<u>40MG</u>	<u>A075607</u>	<u>002</u>	May 10, 2001
<u>AB</u>	TEVA	<u>20MG</u>	<u>A075311</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075311</u>	<u>002</u>	Apr 16, 2001
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>A075062</u>	<u>002</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075062</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>	WOCKHARDT	<u>20MG</u>	<u>A075786</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>A075786</u>	<u>002</u>	Apr 16, 2001
	<u>PEPCID</u>				
<u>AB</u>	MERCK	<u>20MG</u>	<u>N019462</u>	<u>001</u>	Oct 15, 1986
<u>AB</u>	+	<u>40MG</u>	<u>N019462</u>	<u>002</u>	Oct 15, 1986

FAMOTIDINE; IBUPROFEN

TABLET; ORAL

DUEXIS

+ HORIZON PHARMA

26.6MG;800MG

N022519 001 Apr 23, 2011

FEBUXOSTAT

TABLET; ORAL

ULORIC

+ TAKEDA PHARMS

40MG

N021856 001 Feb 13, 2009

+ 80MG

N021856 002 Feb 13, 2009

FELBAMATE

SUSPENSION; ORAL

FELBAMATE

<u>AB</u>	AMNEAL PHARMS	<u>600MG/5ML</u>	<u>A202385</u>	<u>001</u>	Dec 16, 2011
-----------	---------------	------------------	----------------	------------	--------------

FELBATOL

<u>AB</u>	+ MEDA PHARMS	<u>600MG/5ML</u>	<u>N020189</u>	<u>003</u>	Jul 29, 1993
-----------	---------------	------------------	----------------	------------	--------------

TABLET; ORAL

FELBAMATE

<u>AB</u>	AMNEAL PHARMS	<u>400MG</u>	<u>A201680</u>	<u>001</u>	Sep 13, 2011
-----------	---------------	--------------	----------------	------------	--------------

600MGA201680 002 Sep 13, 2011FELBATOL

<u>AB</u>	MEDA PHARMS	<u>400MG</u>	<u>N020189</u>	<u>001</u>	Jul 29, 1993
-----------	-------------	--------------	----------------	------------	--------------

600MGN020189 002 Jul 29, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 176 (of 424)

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

<u>AB</u>	ENDO PHARMS	<u>2.5MG</u>	<u>A200815</u>	<u>001</u>	Oct 28, 2011
<u>AB</u>		<u>5MG</u>	<u>A200815</u>	<u>002</u>	Oct 28, 2011
<u>AB</u>		<u>10MG</u>	<u>A200815</u>	<u>003</u>	Oct 28, 2011
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A090365</u>	<u>001</u>	Dec 17, 2010
<u>AB</u>		<u>5MG</u>	<u>A090365</u>	<u>002</u>	Dec 17, 2010
<u>AB</u>		<u>10MG</u>	<u>A090365</u>	<u>003</u>	Dec 17, 2010
<u>AB</u>	MUTUAL PHARM	<u>2.5MG</u>	<u>A075896</u>	<u>001</u>	Nov 02, 2004
<u>AB</u>		<u>5MG</u>	<u>A075896</u>	<u>002</u>	Nov 02, 2004
<u>AB</u>		<u>10MG</u>	<u>A075896</u>	<u>003</u>	Nov 02, 2004
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A078855</u>	<u>001</u>	Apr 17, 2008
<u>AB</u>		<u>5MG</u>	<u>A078855</u>	<u>002</u>	Apr 17, 2008
<u>AB</u>	<u>+</u>	<u>10MG</u>	<u>A078855</u>	<u>003</u>	Apr 17, 2008
<u>AB</u>	TORRENT PHARMS LTD	<u>2.5MG</u>	<u>A202170</u>	<u>001</u>	Nov 28, 2011
<u>AB</u>		<u>5MG</u>	<u>A202170</u>	<u>002</u>	Nov 28, 2011
<u>AB</u>		<u>10MG</u>	<u>A202170</u>	<u>003</u>	Nov 28, 2011
	<u>PLENDIL</u>				
<u>AB</u>	ASTRAZENECA	<u>2.5MG</u>	<u>N019834</u>	<u>004</u>	Sep 22, 1994
<u>AB</u>		<u>5MG</u>	<u>N019834</u>	<u>001</u>	Jul 25, 1991
<u>AB</u>		<u>10MG</u>	<u>N019834</u>	<u>002</u>	Jul 25, 1991

FENOFIBRATE

CAPSULE; ORAL

FENOFIBRATE (MICRONIZED)

<u>AB</u>	IMPAK LABS	<u>67MG</u>	<u>A075868</u>	<u>001</u>	Oct 27, 2003
<u>AB</u>		<u>134MG</u>	<u>A075868</u>	<u>002</u>	Oct 27, 2003
<u>AB</u>		<u>200MG</u>	<u>A075868</u>	<u>003</u>	Oct 27, 2003
<u>AB</u>	TEVA	<u>67MG</u>	<u>A075753</u>	<u>001</u>	Sep 03, 2002
<u>AB</u>		<u>134MG</u>	<u>A075753</u>	<u>002</u>	Apr 09, 2002
<u>AB</u>	<u>+</u>	<u>200MG</u>	<u>A075753</u>	<u>003</u>	Apr 09, 2002
	ANTARA (MICRONIZED)				
	LUPIN ATLANTIS	43MG	N021695	001	Nov 30, 2004
	<u>+</u>	130MG	N021695	003	Nov 30, 2004
	LIPOFEN				
	CIPHER PHARMS INC	50MG	N021612	001	Jan 11, 2006
	<u>+</u>	150MG	N021612	003	Jan 11, 2006

TABLET; ORAL

FENOFIBRATE

<u>AB</u>	IMPAK LABS	<u>54MG</u>	<u>A076509</u>	<u>001</u>	Mar 26, 2008
<u>AB</u>		<u>160MG</u>	<u>A076509</u>	<u>002</u>	Mar 26, 2008
<u>AB</u>	LUPIN LTD	<u>48MG</u>	<u>A090856</u>	<u>001</u>	Dec 23, 2011
<u>AB</u>		<u>145MG</u>	<u>A090856</u>	<u>002</u>	Dec 23, 2011
<u>AB</u>	MYLAN	<u>54MG</u>	<u>A076520</u>	<u>001</u>	Oct 25, 2007
<u>AB</u>		<u>160MG</u>	<u>A076520</u>	<u>003</u>	Oct 25, 2007
<u>AB</u>	RANBAXY	<u>54MG</u>	<u>A076635</u>	<u>001</u>	Oct 31, 2005
<u>AB</u>		<u>160MG</u>	<u>A076635</u>	<u>003</u>	Oct 31, 2005
<u>AB</u>	TEVA	<u>54MG</u>	<u>A076433</u>	<u>001</u>	May 13, 2005
<u>AB</u>	<u>+</u>	<u>160MG</u>	<u>A076433</u>	<u>002</u>	May 13, 2005
	<u>TRICOR</u>				
<u>AB</u>	ABBOTT LABS PHARM	<u>48MG</u>	<u>N021656</u>	<u>001</u>	Nov 05, 2004
<u>AB</u>	<u>+</u>	<u>145MG</u>	<u>N021656</u>	<u>002</u>	Nov 05, 2004
	TRIGLIDE				
BX	+ SKYEPHARMA AG	160MG	N021350	002	May 07, 2005
	FENOFIBRATE				
	RANBAXY	107MG	A076635	002	Oct 31, 2005
	FENOGLIDE				
	SANTARUS	40MG	N022118	001	Aug 10, 2007
	<u>+</u>	120MG	N022118	002	Aug 10, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 177 (of 424)

FENOFIBRATE

TABLET; ORAL
 TRIGLIDE
 SKYEPHARMA AG 50MG N021350 001 May 07, 2005

FENOFIBRIC ACID

TABLET; ORAL
 FIBRICOR
 AR HOLDING CO INC 35MG N022418 001 Aug 14, 2009
 + 105MG N022418 002 Aug 14, 2009

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION
CORLOPAM
AP + HOSPIRA EQ 10MG BASE/ML N019922 001 Sep 23, 1997
FENOLDOPAM MESYLATE
AP BEDFORD LABS EQ 10MG BASE/ML A076582 001 Oct 12, 2004
AP SANDOZ EQ 10MG BASE/ML A077155 001 Feb 15, 2005

FENOPROFEN CALCIUM

CAPSULE; ORAL
 NALFON
 + PEDINOL EQ 200MG BASE N017604 003
 EQ 400MG BASE N017604 004 Jul 21, 2009

TABLET; ORAL

FENOPROFEN CALCIUM
AB IVAX SUB TEVA PHARMS EQ 600MG BASE A072557 001 Aug 29, 1988
AB + MYLAN EQ 600MG BASE A072267 001 Aug 17, 1988

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL
DURAGESIC-100
AB JANSSEN PHARMS 100MCG/HR N019813 001 Aug 07, 1990
DURAGESIC-12
AB JANSSEN PHARMS 12.5MCG/HR N019813 005 Feb 04, 2005
DURAGESIC-25
AB + JANSSEN PHARMS 25MCG/HR N019813 004 Aug 07, 1990
DURAGESIC-50
AB JANSSEN PHARMS 50MCG/HR N019813 003 Aug 07, 1990
DURAGESIC-75
AB JANSSEN PHARMS 75MCG/HR N019813 002 Aug 07, 1990
FENTANYL-100
AB ACTAVIS 100MCG/HR A077062 004 Aug 20, 2007
AB LAVIPHARM LABS 100MCG/HR A077051 004 Aug 04, 2006
AB MALLINCKRODT INC 100MCG/HR A077154 004 Feb 09, 2011
AB MYLAN TECHNOLOGIES 100MCG/HR A076258 004 Jan 28, 2005
AB NOVEN 100MCG/HR A077775 004 Oct 16, 2009
AB TEVA PHARMS 100MCG/HR A077449 004 Oct 20, 2008
AB WATSON LABS 100MCG/HR A076709 004 Aug 20, 2007
FENTANYL-12
AB MYLAN TECHNOLOGIES 12.5MCG/HR A076258 005 Jan 23, 2007
FENTANYL-25
AB ACTAVIS 25MCG/HR A077062 001 Aug 20, 2007
AB LAVIPHARM LABS 25MCG/HR A077051 001 Aug 04, 2006
AB MALLINCKRODT INC 25MCG/HR A077154 001 Feb 09, 2011
AB MYLAN TECHNOLOGIES 25MCG/HR A076258 001 Jan 28, 2005
AB NOVEN 25MCG/HR A077775 001 Oct 16, 2009
AB TEVA PHARMS 25MCG/HR A077449 001 Oct 20, 2008
AB WATSON LABS 25MCG/HR A076709 001 Aug 20, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 178 (of 424)

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL-50

<u>AB</u>	ACTAVIS	<u>50MCG/HR</u>	<u>A077062</u>	<u>002</u>	Aug 20, 2007
<u>AB</u>	LAVIPHARM LABS	<u>50MCG/HR</u>	<u>A077051</u>	<u>002</u>	Aug 04, 2006
<u>AB</u>	MALLINCKRODT INC	<u>50MCG/HR</u>	<u>A077154</u>	<u>002</u>	Feb 09, 2011
<u>AB</u>	MYLAN TECHNOLOGIES	<u>50MCG/HR</u>	<u>A076258</u>	<u>002</u>	Jan 28, 2005
<u>AB</u>	NOVEN	<u>50MCG/HR</u>	<u>A077775</u>	<u>002</u>	Oct 16, 2009
<u>AB</u>	TEVA PHARMS	<u>50MCG/HR</u>	<u>A077449</u>	<u>002</u>	Oct 20, 2008
<u>AB</u>	WATSON LABS	<u>50MCG/HR</u>	<u>A076709</u>	<u>002</u>	Aug 20, 2007

FENTANYL-75

<u>AB</u>	ACTAVIS	<u>75MCG/HR</u>	<u>A077062</u>	<u>003</u>	Aug 20, 2007
<u>AB</u>	LAVIPHARM LABS	<u>75MCG/HR</u>	<u>A077051</u>	<u>003</u>	Aug 04, 2006
<u>AB</u>	MALLINCKRODT INC	<u>75MCG/HR</u>	<u>A077154</u>	<u>003</u>	Feb 09, 2011
<u>AB</u>	MYLAN TECHNOLOGIES	<u>75MCG/HR</u>	<u>A076258</u>	<u>003</u>	Jan 28, 2005
<u>AB</u>	NOVEN	<u>75MCG/HR</u>	<u>A077775</u>	<u>003</u>	Oct 16, 2009
<u>AB</u>	TEVA PHARMS	<u>75MCG/HR</u>	<u>A077449</u>	<u>003</u>	Oct 20, 2008
<u>AB</u>	WATSON LABS	<u>75MCG/HR</u>	<u>A076709</u>	<u>003</u>	Aug 20, 2007

FENTANYL CITRATE

FILM; Buccal

ONSOLIS

MEDA PHARMS

EQ 0.2MG BASE
+ EQ 0.4MG BASE
EQ 0.6MG BASE
EQ 0.8MG BASE
EQ 1.2MG BASEN022266 001 Jul 16, 2009
N022266 002 Jul 16, 2009
N022266 003 Jul 16, 2009
N022266 004 Jul 16, 2009
N022266 005 Jul 16, 2009

INJECTABLE; INJECTION

FENTANYL CITRATE

<u>AP</u>	HOSPIRA	<u>EQ 0.05MG BASE/ML</u>	<u>N019115</u>	<u>001</u>	Jan 12, 1985
<u>AP</u>	+ BAXTER HLTHCARE	<u>EQ 0.05MG BASE/ML</u>	<u>N019101</u>	<u>001</u>	Jul 11, 1984
<u>AP</u>	HOSPIRA	<u>EQ 0.05MG BASE/ML</u>	<u>A072786</u>	<u>001</u>	Sep 24, 1991
<u>AP</u>	+ AKORN	<u>EQ 0.05MG BASE/ML</u>	<u>N016619</u>	<u>001</u>	

SPRAY, METERED; NASAL

LAZANDA

ARCHIMEDES
+ 400MCGN022569 001 Jun 30, 2011
N022569 002 Jun 30, 2011

TABLET; Buccal

FENTANYL CITRATE

<u>AB</u>	WATSON LABS	<u>EQ 0.1MG BASE</u>	<u>A079075</u>	<u>001</u>	Jan 07, 2011
<u>AB</u>		<u>EQ 0.2MG BASE</u>	<u>A079075</u>	<u>002</u>	Jan 07, 2011
<u>AB</u>		<u>EQ 0.4MG BASE</u>	<u>A079075</u>	<u>003</u>	Jan 07, 2011
<u>AB</u>		<u>EQ 0.6MG BASE</u>	<u>A079075</u>	<u>004</u>	Jan 07, 2011
<u>AB</u>		<u>EQ 0.8MG BASE</u>	<u>A079075</u>	<u>005</u>	Jan 07, 2011

FENTORA

<u>AB</u>	CEPHALON	<u>EQ 0.1MG BASE</u>	<u>N021947</u>	<u>001</u>	Sep 25, 2006
<u>AB</u>		<u>EQ 0.2MG BASE</u>	<u>N021947</u>	<u>002</u>	Sep 25, 2006
<u>AB</u>	+	<u>EQ 0.4MG BASE</u>	<u>N021947</u>	<u>003</u>	Sep 25, 2006
<u>AB</u>		<u>EQ 0.6MG BASE</u>	<u>N021947</u>	<u>004</u>	Sep 25, 2006
<u>AB</u>		<u>EQ 0.8MG BASE</u>	<u>N021947</u>	<u>005</u>	Sep 25, 2006

TABLET; SUBLINGUAL

ABSTRAL

PROSTRAKAN INC
+
EQ 0.1MG BASE
EQ 0.2MG BASE
EQ 0.3MG BASE
EQ 0.4MG BASE
EQ 0.6MG BASEN022510 001 Jan 07, 2011
N022510 002 Jan 07, 2011
N022510 003 Jan 07, 2011
N022510 004 Jan 07, 2011
N022510 005 Jan 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 179 (of 424)

FENTANYL CITRATE

TABLET; SUBLINGUAL ABSTRAL PROSTRAKAN INC	EQ 0.8MG BASE	N022510 006	Jan 07, 2011
TROCHE / LOZENGE; TRANSMUCOSAL			
<u>ACTIQ</u> <u>CEPHALON</u>	<u>EQ 0.2MG BASE</u>	<u>N020747 001</u>	Nov 04, 1998
<u>+</u>	<u>EQ 0.4MG BASE</u>	<u>N020747 002</u>	Nov 04, 1998
<u>CEPHALON</u>	<u>EQ 0.6MG BASE</u>	<u>N020747 003</u>	Nov 04, 1998
<u>+</u>	<u>EQ 0.8MG BASE</u>	<u>N020747 004</u>	Nov 04, 1998
<u>CEPHALON</u>	<u>EQ 1.2MG BASE</u>	<u>N020747 005</u>	Nov 04, 1998
<u>+</u>	<u>EQ 1.6MG BASE</u>	<u>N020747 006</u>	Nov 04, 1998
<u>FENTANYL CITRATE</u>			
<u>BARR</u>	<u>EQ 0.2MG BASE</u>	<u>A077312 001</u>	Oct 30, 2009
<u>+</u>	<u>EQ 0.4MG BASE</u>	<u>A077312 002</u>	Oct 30, 2009
<u>BARR</u>	<u>EQ 0.6MG BASE</u>	<u>A077312 003</u>	Oct 30, 2009
<u>+</u>	<u>EQ 0.8MG BASE</u>	<u>A077312 004</u>	Oct 30, 2009
<u>BARR</u>	<u>EQ 1.2MG BASE</u>	<u>A077312 005</u>	Oct 30, 2009
<u>+</u>	<u>EQ 1.6MG BASE</u>	<u>A077312 006</u>	Oct 30, 2009
<u>MALLINCKRODT</u>	<u>EQ 0.2MG BASE</u>	<u>A078907 001</u>	Oct 30, 2009
<u>+</u>	<u>EQ 0.4MG BASE</u>	<u>A078907 002</u>	Oct 30, 2009
<u>MALLINCKRODT</u>	<u>EQ 0.6MG BASE</u>	<u>A078907 003</u>	Oct 30, 2009
<u>+</u>	<u>EQ 0.8MG BASE</u>	<u>A078907 004</u>	Oct 30, 2009
<u>MALLINCKRODT</u>	<u>EQ 1.2MG BASE</u>	<u>A078907 005</u>	Oct 30, 2009
<u>+</u>	<u>EQ 1.6MG BASE</u>	<u>A078907 006</u>	Oct 30, 2009

FERRIC HEXACYANOFERRATE (II)

CAPSULE; ORAL RADIOGARDASE (PRUSSIAN BLUE) + HEYL CHEMISCH	500MG	N021626 001	Oct 02, 2003
--	-------	-------------	--------------

FERUMOXIDES

INJECTABLE; INJECTION FERIDEX I.V. + AMAG PHARMS INC	EQ 11.2MG IRON/ML	N020416 001	Aug 30, 1996
--	-------------------	-------------	--------------

FERUMOXSIL

SUSPENSION; ORAL GASTROMARK + AMAG PHARMS INC	EQ 0.175MG IRON/ML	N020410 001	Dec 06, 1996
---	--------------------	-------------	--------------

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 TOVIAZ PFIZER +	4MG	N022030 001	Oct 31, 2008
	8MG	N022030 002	Oct 31, 2008

FEXOFENADINE HYDROCHLORIDE

SUSPENSION; ORAL ALLEGRA + SANOFI AVENTIS US	30MG/5ML	N021963 001	Oct 16, 2006
--	----------	-------------	--------------

TABLET; ORAL

<u>FEXOFENADINE HYDROCHLORIDE</u> <u>BARR</u>	<u>30MG</u>	<u>A076191 001</u>	Aug 31, 2005
--	-------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 180 (of 424)

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>60MG</u>	<u>A076191</u>	<u>002</u>	Aug 31, 2005
<u>AB</u>		<u>180MG</u>	<u>A076191</u>	<u>003</u>	Aug 31, 2005
<u>AB</u>	DR REDDYS LABS LTD	<u>30MG</u>	<u>A076502</u>	<u>001</u>	Apr 11, 2006
<u>AB</u>		<u>60MG</u>	<u>A076502</u>	<u>002</u>	Apr 11, 2006
<u>AB</u>		<u>180MG</u>	<u>A076502</u>	<u>003</u>	Apr 11, 2006
<u>AB</u>	MYLAN	<u>30MG</u>	<u>A077081</u>	<u>002</u>	Apr 11, 2008
<u>AB</u>		<u>60MG</u>	<u>A077081</u>	<u>003</u>	Apr 11, 2008
<u>AB</u>		<u>180MG</u>	<u>A077081</u>	<u>001</u>	Apr 16, 2007
<u>AB</u>	TEVA	<u>30MG</u>	<u>A076447</u>	<u>001</u>	Sep 01, 2005
<u>AB</u>		<u>60MG</u>	<u>A076447</u>	<u>002</u>	Sep 01, 2005
<u>AB</u>		<u>180MG</u>	<u>A076447</u>	<u>003</u>	Sep 01, 2005

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>60MG;120MG</u>	<u>A076236</u>	<u>001</u>	Apr 14, 2005
<u>AB</u>	IMPAX PHARMS	<u>60MG;120MG</u>	<u>A076298</u>	<u>001</u>	Nov 12, 2010

FIDAXOMICIN

TABLET; ORAL

DIFICID

+ OPTIMER PHARMS 200MG

N201699 001 May 27, 2011

FINASTERIDE

TABLET; ORAL

FINASTERIDE

<u>AB</u>	ACCORD HLTHCARE INC	<u>5MG</u>	<u>A090121</u>	<u>001</u>	Feb 23, 2010
<u>AB</u>	ACTAVIS TOTOWA	<u>5MG</u>	<u>A077914</u>	<u>001</u>	Mar 28, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078341</u>	<u>001</u>	Oct 30, 2007
<u>AB</u>	DR REDDYS LABS INC	<u>1MG</u>	<u>A076436</u>	<u>001</u>	Jul 28, 2006
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076437</u>	<u>001</u>	Feb 28, 2007
<u>AB</u>	GEDEON RICHTER USA	<u>5MG</u>	<u>A077251</u>	<u>001</u>	Dec 22, 2006
<u>AB</u>	HETERO DRUGS LTD	<u>5MG</u>	<u>A090061</u>	<u>001</u>	Jun 07, 2010
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A077578</u>	<u>001</u>	Dec 18, 2006
<u>AB</u>	SUN PHARMA GLOBAL	<u>5MG</u>	<u>A090507</u>	<u>001</u>	Aug 16, 2011
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076511</u>	<u>001</u>	Dec 15, 2006
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078900</u>	<u>001</u>	Dec 28, 2009
<u>AB</u>	<u>PROPECIA</u>				
<u>AB</u>	+ MERCK	<u>1MG</u>	<u>N020788</u>	<u>001</u>	Dec 19, 1997
<u>AB</u>	<u>PROSCAR</u>				
<u>AB</u>	+ MERCK	<u>5MG</u>	<u>N020180</u>	<u>001</u>	Jun 19, 1992

FINGOLIMOD

CAPSULE; ORAL

GILENYA

+ NOVARTIS 0.5MG

N022527 001 Sep 21, 2010

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA	<u>100MG</u>	<u>A076835</u>	<u>001</u>	Nov 30, 2005
<u>AB</u>	IMPAX PHARMS	<u>100MG</u>	<u>A076234</u>	<u>001</u>	Aug 28, 2003
<u>AB</u>	+ PADDICK LLC	<u>100MG</u>	<u>A076831</u>	<u>001</u>	Dec 16, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 181 (of 424)

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

<u>AB</u>	AMNEAL PHARM	<u>50MG</u>	<u>A075442</u>	<u>001</u>	Jul 31, 2001
<u>AB</u>		<u>100MG</u>	<u>A075442</u>	<u>002</u>	Jul 31, 2001
<u>AB</u>		<u>150MG</u>	<u>A075442</u>	<u>003</u>	Jul 31, 2001
<u>AB</u>	APOTEX INC	<u>50MG</u>	<u>A079164</u>	<u>001</u>	Jul 09, 2009
<u>AB</u>		<u>100MG</u>	<u>A079164</u>	<u>002</u>	Jul 09, 2009
<u>AB</u>		<u>150MG</u>	<u>A079164</u>	<u>003</u>	Jul 09, 2009
<u>AB</u>	BARR	<u>50MG</u>	<u>A075882</u>	<u>001</u>	Oct 28, 2002
<u>AB</u>		<u>100MG</u>	<u>A075882</u>	<u>002</u>	Oct 28, 2002
<u>AB</u>		<u>150MG</u>	<u>A075882</u>	<u>003</u>	Oct 28, 2002
<u>AB</u>	RANBAXY	<u>50MG</u>	<u>A076421</u>	<u>001</u>	Mar 28, 2003
<u>AB</u>		<u>100MG</u>	<u>A076421</u>	<u>002</u>	Mar 28, 2003
<u>AB</u>		<u>150MG</u>	<u>A076421</u>	<u>003</u>	Mar 28, 2003
<u>AB</u>	ROXANE	<u>50MG</u>	<u>A076278</u>	<u>001</u>	Jan 14, 2003
<u>AB</u>		<u>100MG</u>	<u>A076278</u>	<u>002</u>	Jan 14, 2003
<u>AB</u>		<u>150MG</u>	<u>A076278</u>	<u>003</u>	Jan 14, 2003
	<u>TAMBOCOR</u>				
<u>AB</u>	MEDICIS	<u>50MG</u>	<u>N018830</u>	<u>004</u>	Aug 23, 1988
<u>AB</u>		<u>100MG</u>	<u>N018830</u>	<u>001</u>	Oct 31, 1985
<u>AB</u>	+	<u>150MG</u>	<u>N018830</u>	<u>003</u>	Jun 03, 1988

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

<u>AP</u>	APP PHARMS	<u>500MG/VIAL</u>	<u>A075837</u>	<u>001</u>	Feb 22, 2001
<u>AP</u>	+	<u>500MG/VIAL</u>	<u>A075387</u>	<u>001</u>	Apr 16, 2000

FLUCONAZOLE

FOR SUSPENSION; ORAL

DIFLUCAN

<u>AB</u>	PFIZER	<u>50MG/5ML</u>	<u>N020090</u>	<u>001</u>	Dec 23, 1993
<u>AB</u>	+	<u>200MG/5ML</u>	<u>N020090</u>	<u>002</u>	Dec 23, 1993

FLUCONAZOLE

<u>AB</u>	AUROBINDO PHARM	<u>50MG/5ML</u>	<u>A079150</u>	<u>001</u>	Sep 18, 2009
<u>AB</u>		<u>200MG/5ML</u>	<u>A079150</u>	<u>002</u>	Sep 18, 2009
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>50MG/5ML</u>	<u>A077523</u>	<u>001</u>	Sep 12, 2007
<u>AB</u>		<u>200MG/5ML</u>	<u>A077523</u>	<u>002</u>	Sep 12, 2007
<u>AB</u>	RANBAXY	<u>50MG/5ML</u>	<u>A076332</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>200MG/5ML</u>	<u>A076332</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>	ROXANE	<u>50MG/5ML</u>	<u>A076246</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>200MG/5ML</u>	<u>A076246</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>	TARO PHARM IND	<u>50MG/5ML</u>	<u>A076918</u>	<u>001</u>	Dec 18, 2006
<u>AB</u>		<u>200MG/5ML</u>	<u>A076918</u>	<u>002</u>	Dec 18, 2006

INJECTABLE; INJECTION

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	PFIZER	<u>200MG/100ML (2MG/ML)</u>	<u>N019950</u>	<u>003</u>	Sep 29, 1992
<u>AP</u>	+		<u>400MG/200ML (2MG/ML)</u>	<u>N019950</u>	<u>005</u>	Jul 08, 1994

DIFLUCAN IN SODIUM CHLORIDE 0.9%

<u>AP</u>	+	PFIZER	<u>200MG/100ML (2MG/ML)</u>	<u>N019950</u>	<u>001</u>	Jan 29, 1990
<u>AP</u>	+		<u>400MG/200ML (2MG/ML)</u>	<u>N019950</u>	<u>006</u>	Jan 29, 1990

DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+	PFIZER	<u>200MG/100ML (2MG/ML)</u>	<u>N019950</u>	<u>002</u>	Jan 29, 1990
<u>AP</u>	+		<u>400MG/200ML (2MG/ML)</u>	<u>N019950</u>	<u>004</u>	Jan 29, 1990

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	ACS DOBFAR INFO SA	<u>200MG/100ML (2MG/ML)</u>	<u>A079104</u>	<u>001</u>	Jul 30, 2009
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A079104</u>	<u>002</u>	Jul 30, 2009

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	APOTEX INC	<u>200MG/100ML (2MG/ML)</u>	<u>A076888</u>	<u>001</u>	Mar 25, 2005
-----------	------------	-----------------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 182 (of 424)

FLUCONAZOLE

INJECTABLE; INJECTION

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	APOTEX INC	<u>400MG/200ML (2MG/ML)</u>	<u>A076888</u> <u>002</u>	Mar 25, 2005
<u>AP</u>	HOSPIRA	<u>200MG/100ML (2MG/ML)</u>	<u>A076304</u> <u>001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076304</u> <u>002</u>	Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

<u>AP</u>	APP PHARMS	<u>200MG/100ML (2MG/ML)</u>	<u>A076145</u> <u>001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076145</u> <u>002</u>	Jul 29, 2004
<u>AP</u>	BEDFORD	<u>200MG/100ML (2MG/ML)</u>	<u>A076087</u> <u>001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076087</u> <u>003</u>	Jul 29, 2004
<u>AP</u>	CLARIS LIFESCIENCES	<u>200MG/100ML (2MG/ML)</u>	<u>A077947</u> <u>001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077947</u> <u>002</u>	May 26, 2010
<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML (2MG/ML)</u>	<u>A076736</u> <u>001</u>	Aug 23, 2005
<u>AP</u>	TEVA PARENTERAL	<u>200MG/100ML (2MG/ML)</u>	<u>A076653</u> <u>001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076653</u> <u>002</u>	Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML (2MG/ML)</u>	<u>A076766</u> <u>001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076766</u> <u>002</u>	Jul 29, 2004
<u>AP</u>	BEDFORD LABS	<u>200MG/100ML (2MG/ML)</u>	<u>A078107</u> <u>001</u>	Jul 30, 2008
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A078107</u> <u>002</u>	Jul 30, 2008
<u>AP</u>	CLARIS LIFESCIENCES	<u>200MG/100ML (2MG/ML)</u>	<u>A077909</u> <u>001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077909</u> <u>002</u>	May 26, 2010
<u>AP</u>	HOSPIRA	<u>200MG/100ML (2MG/ML)</u>	<u>A076303</u> <u>001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076303</u> <u>002</u>	Jul 29, 2004
<u>AP</u>	TEVA PARENTERAL	<u>200MG/100ML (2MG/ML)</u>	<u>A076837</u> <u>001</u>	Jan 13, 2005
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076837</u> <u>002</u>	Jan 13, 2005

FULCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	CLARIS LIFESCIENCES	<u>200MG/100ML (2MG/ML)</u>	<u>A077988</u> <u>001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077988</u> <u>002</u>	May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%BEDFORD 100MG/50ML (2MG/ML)A076087 002 Sep 26, 2008

TABLET; ORAL

DIFLUJCAN

<u>AB</u>	PFIZER	<u>50MG</u>	<u>N019949</u> <u>001</u>	Jan 29, 1990
<u>AB</u>		<u>100MG</u>	<u>N019949</u> <u>002</u>	Jan 29, 1990
<u>AB</u>		<u>150MG</u>	<u>N019949</u> <u>004</u>	Jun 30, 1994
<u>AB</u>	+	<u>200MG</u>	<u>N019949</u> <u>003</u>	Jan 29, 1990

FLUCONAZOLE

<u>AB</u>	AMNEAL PHARM	<u>50MG</u>	<u>A078423</u> <u>001</u>	Mar 07, 2011
<u>AB</u>		<u>100MG</u>	<u>A078423</u> <u>002</u>	Mar 07, 2011
<u>AB</u>		<u>150MG</u>	<u>A078423</u> <u>003</u>	Mar 07, 2011
<u>AB</u>		<u>200MG</u>	<u>A078423</u> <u>004</u>	Mar 07, 2011
<u>AB</u>	APOTEX	<u>50MG</u>	<u>A076665</u> <u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076665</u> <u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076665</u> <u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076665</u> <u>004</u>	Jul 29, 2004
<u>AB</u>	AUROBINDO PHARMA	<u>50MG</u>	<u>A077731</u> <u>001</u>	Oct 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A077731</u> <u>002</u>	Oct 07, 2008
<u>AB</u>		<u>150MG</u>	<u>A077731</u> <u>003</u>	Oct 07, 2008
<u>AB</u>		<u>200MG</u>	<u>A077731</u> <u>004</u>	Oct 07, 2008
<u>AB</u>	GLENMARK GENERICS	<u>50MG</u>	<u>A077253</u> <u>001</u>	Jan 25, 2006
<u>AB</u>		<u>100MG</u>	<u>A077253</u> <u>002</u>	Jan 25, 2006
<u>AB</u>		<u>150MG</u>	<u>A077253</u> <u>003</u>	Jan 25, 2006
<u>AB</u>		<u>200MG</u>	<u>A077253</u> <u>004</u>	Jan 25, 2006
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>50MG</u>	<u>A076077</u> <u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076077</u> <u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076077</u> <u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076077</u> <u>004</u>	Jul 29, 2004
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A076042</u> <u>001</u>	Jul 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 183 (of 424)

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

<u>AB</u>	MYLAN	<u>50MG</u>	<u>A076351</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076042</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076351</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076042</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076351</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076042</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076351</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	RANBAXY	<u>50MG</u>	<u>A076386</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076386</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076386</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076386</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	TARO	<u>50MG</u>	<u>A076507</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A076507</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A076507</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A076507</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	TEVA	<u>50MG</u>	<u>A074681</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>A074681</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>A074681</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>A074681</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	UNIQUE PHARM LABS	<u>50MG</u>	<u>A076957</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076957</u>	<u>002</u>	Sep 28, 2005
<u>AB</u>		<u>200MG</u>	<u>A076957</u>	<u>003</u>	Sep 28, 2005
	<u>FLUCONAZOLE</u>				
<u>BX</u>	DR REDDYS LABS INC	50MG			A076658 001 Jul 29, 2004
<u>BX</u>		100MG			A076658 002 Jul 29, 2004
<u>BX</u>		150MG			A076658 003 Jul 29, 2004
<u>BX</u>		200MG			A076658 004 Jul 29, 2004
<u>BX</u>	PLIVA	50MG			A076424 001 Jul 29, 2004
<u>BX</u>		100MG			A076424 002 Jul 29, 2004
<u>BX</u>		150MG			A076424 003 Jul 29, 2004
<u>BX</u>		200MG			A076424 004 Jul 29, 2004

FLUCYTOSINE

CAPSULE; ORAL

ANCOCOBON

<u>AB</u>	VALEANT	<u>250MG</u>	<u>N017001</u>	<u>001</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N017001</u>	<u>002</u>	
	<u>FLUCYTOSINE</u>				
<u>AB</u>	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A201566</u>	<u>001</u>	Jun 28, 2011
<u>AB</u>		<u>500MG</u>	<u>A201566</u>	<u>002</u>	Jun 28, 2011

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

<u>AP</u>	+ GENZYME	<u>50MG/VIAL</u>	<u>N020038</u>	<u>001</u>	Apr 18, 1991
	<u>FLUDARABINE PHOSPHATE</u>				
<u>AP</u>	ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078610</u>	<u>001</u>	Feb 11, 2009
<u>AP</u>	APP PHARMS	<u>50MG/2ML (25MG/ML)</u>	<u>A078393</u>	<u>001</u>	Oct 15, 2007
<u>AP</u>		<u>50MG/VIAL</u>	<u>A078544</u>	<u>001</u>	Oct 15, 2007
<u>AP</u>	BIONICHE PHARMA USA	<u>50MG/2ML (25MG/ML)</u>	<u>A090724</u>	<u>001</u>	Sep 27, 2010
<u>AP</u>	HOSPIRA	<u>50MG/VIAL</u>	<u>A077790</u>	<u>001</u>	Apr 06, 2007
<u>AP</u>	ONCO THERAPIES LTD	<u>50MG/2ML (25MG/ML)</u>	<u>A200647</u>	<u>001</u>	Dec 21, 2011
<u>AP</u>	+	<u>SANDOZ</u>	<u>N022137</u>	<u>001</u>	Sep 21, 2007
<u>AP</u>	TEVA PARENTERAL	<u>50MG/VIAL</u>	<u>A076349</u>	<u>001</u>	Aug 28, 2003
<u>AP</u>	+	<u>50MG/2ML (25MG/ML)</u>	<u>A076661</u>	<u>001</u>	Apr 28, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 184 (of 424)

FLUDARABINE PHOSPHATE

TABLET; ORAL
OFORTA
+ SANOFI AVENTIS US 10MG N022273 001 Dec 18, 2008

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS
FLUDEOXYGLUCOSE F18

<u>AP</u> + FEINSTEIN	<u>20-200mCi/ML</u>	<u>N021870 001</u>	Aug 19, 2005
<u>AP</u> PETNET	<u>20-200mCi/ML</u>	<u>A079086 001</u>	Feb 25, 2011
FLUDEOXYGLUCOSE F 18			
+ WEILL MEDCL COLL	10-100mCi/ML	N021768 001	Aug 05, 2004
FLUDEOXYGLUCOSE F18			
+ FEINSTEIN	20-300mCi/ML	N021870 002	Nov 21, 2008

FLUDROCORTISONE ACETATE

TABLET; ORAL
FLUDROCORTISONE ACETATE

<u>AB</u> BARR	<u>0.1MG</u>	<u>A040425 001</u>	Jan 21, 2003
<u>AB</u> + IMPAX LABS	<u>0.1MG</u>	<u>A040431 001</u>	Mar 18, 2002
<u>AB</u> WEST-WARD PHARM CORP	<u>0.1MG</u>	<u>A091302 001</u>	Jul 22, 2011

FLUMAZENIL

INJECTABLE; INJECTION
FLUMAZENIL

<u>AP</u> APP PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076955 002</u>	Oct 12, 2004
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A076955 001</u>	Oct 12, 2004
<u>AP</u> BAXTER HLTHCARE	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076787 002</u>	Oct 12, 2004
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A076787 001</u>	Oct 12, 2004
<u>AP</u> BEDFORD LABS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076256 002</u>	Oct 12, 2004
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A076256 001</u>	Oct 12, 2004
<u>AP</u> CLARIS LIFESCIENCES	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076755 002</u>	Oct 12, 2004
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A076755 001</u>	Oct 12, 2004
<u>AP</u> HIKMA FARMACEUTICA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078527 001</u>	Mar 23, 2009
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A078527 002</u>	Mar 23, 2009
<u>AP</u> PFIZER	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078595 001</u>	May 13, 2008
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A078595 002</u>	May 13, 2008
<u>AP</u> SANDOZ	<u>1MG/10ML (0.1MG/ML)</u>	<u>A077071 002</u>	May 03, 2005
<u>AP</u>	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A077071 001</u>	May 03, 2005
<u>AP</u> TEVA PARENTERAL	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076589 002</u>	Oct 12, 2004
<u>AP</u>	<u>1MG/10ML (0.1MG/ML)</u>	<u>A076589 001</u>	Oct 12, 2004
<u>ROMAZICON</u>			
<u>AP</u> + HOFFMANN LA ROCHE	<u>1MG/10ML (0.1MG/ML)</u>	<u>N020073 001</u>	Dec 20, 1991
<u>AP</u>	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N020073 002</u>	Dec 20, 1991

FLUNISOLIDE

AEROSOL, METERED; INHALATION
AEROBID

+ ROCHE PALO	0.25MG/INH	N018340 001	Aug 17, 1984
AEROSPAN HFA			
+ ACTON PHARMS	EQ 78MCG BASE/INH	N021247 001	Jan 27, 2006

SPRAY, METERED; NASAL
FLUNISOLIDE

<u>AB</u> + BAUSCH AND LOMB	<u>0.025MG/SPRAY</u>	<u>A074805 001</u>	Feb 20, 2002
<u>AB</u> HH AND P	<u>0.025MG/SPRAY</u>	<u>A077704 001</u>	Aug 03, 2006
FLUNISOLIDE			
+ APOTEX INC	0.029MG/SPRAY	A077436 001	Aug 09, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 185 (of 424)

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<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
	<u>SYNALAR</u>				
<u>AT</u>	+ MEDIMETRIKS PHARMS	<u>0.01%</u>	<u>N012787</u>	<u>004</u>	
<u>AT</u>	+	<u>0.025%</u>	<u>N012787</u>	<u>002</u>	
<u>AT</u>	+	<u>0.025%</u>	<u>N012787</u>	<u>005</u>	

IMPLANT; INTRAVITREAL

RETISERT

+ BAUSCH AND LOMB 0.59MG

N021737 001 Apr 08, 2005

OIL; TOPICAL

DERMA-SMOOTH/EFS

<u>AT</u>	+ HILL DERMAC	<u>0.01%</u>	<u>N019452</u>	<u>001</u>	Feb 03, 1988
<u>AT</u>	+	<u>0.01%</u>	<u>N019452</u>	<u>002</u>	Nov 09, 2005
	<u>FLUOCINOLONE ACETONIDE</u>				
<u>AT</u>	IDENTI PHARMS INC	<u>0.01%</u>	<u>A201759</u>	<u>001</u>	Oct 17, 2011
<u>AT</u>		<u>0.01%</u>	<u>A201764</u>	<u>001</u>	Oct 17, 2011

OIL/DROPS; OTIC

DERMOTIC

<u>AT</u>	+ HILL DERMAC	<u>0.01%</u>	<u>N019452</u>	<u>003</u>	Nov 09, 2005
	<u>FLUOCINOLONE ACETONIDE</u>				
<u>AT</u>	IDENTI PHARMS INC	<u>0.1%</u>	<u>A091306</u>	<u>001</u>	Oct 17, 2011

OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>0.025%</u>	<u>A088168</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>	G AND W LABS	<u>0.025%</u>	<u>A089524</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>	TARO	<u>0.025%</u>	<u>A040041</u>	<u>001</u>	Sep 15, 1994
	<u>SYNALAR</u>				
<u>AT</u>	+ MEDIMETRIKS PHARMS	<u>0.025%</u>	<u>N013960</u>	<u>001</u>	

SHAMPOO; TOPICAL

CAPEX

+ GALDERMA LABS LP 0.01%

N020001 001 Aug 27, 1990

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>0.01%</u>	<u>A088167</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>	TARO	<u>0.01%</u>	<u>A089124</u>	<u>001</u>	Sep 11, 1985
	<u>SYNALAR</u>				
<u>AT</u>	+ MEDIMETRIKS PHARMS	<u>0.01%</u>	<u>N015296</u>	<u>001</u>	

FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM; TOPICAL

TRI-LUMA

+ GALDERMA LABS LP 0.01%;4%;0.05%

N021112 001 Jan 18, 2002

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

<u>AB1</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A073085</u>	<u>001</u>	Feb 14, 1992
<u>AB1</u>	FOUGERA	<u>0.05%</u>	<u>A073030</u>	<u>001</u>	Oct 17, 1994
<u>AB1</u>	TARO	<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
	<u>LIDEX</u>				
<u>AB1</u>	+ MEDICIS	<u>0.05%</u>	<u>N016908</u>	<u>002</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 186 (of 424)

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE EMULSIFIED BASE

<u>AB2</u>	ALTANA	<u>0.05%</u>	<u>A076586</u>	<u>001</u>	Jun 23, 2004
<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
	<u>LIDEX-E</u>				
<u>AB2</u> + MEDICIS	VANOS	<u>0.05%</u>	<u>N016908</u>	<u>003</u>	
+ MEDICIS		0.1%	N021758	001	Feb 11, 2005

GEL; TOPICAL

FLUOCINONIDE

<u>AB</u>	FOUGERA	<u>0.05%</u>	<u>A072933</u>	<u>001</u>	Dec 30, 1994
<u>AB</u>	TARO	<u>0.05%</u>	<u>A074935</u>	<u>001</u>	Jul 29, 1997
<u>AB</u>	TEVA	<u>0.05%</u>	<u>A072537</u>	<u>001</u>	Feb 07, 1989
	<u>LIDEX</u>				
<u>AB</u> + MEDICIS		<u>0.05%</u>	<u>N017373</u>	<u>001</u>	

OINTMENT; TOPICAL

FLUOCINONIDE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A074905</u>	<u>001</u>	Aug 26, 1997
<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
	<u>LIDEX</u>				
<u>AB</u> + MEDICIS		<u>0.05%</u>	<u>N016909</u>	<u>002</u>	

SOLUTION; TOPICAL

FLUOCINONIDE

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>0.05%</u>	<u>A071535</u>	<u>001</u>	Dec 02, 1988
<u>AT</u>	FOUGERA	<u>0.05%</u>	<u>A072934</u>	<u>001</u>	Feb 27, 1995
<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
	<u>LIDEX</u>				
<u>AT</u> + MEDICIS		<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
<u>AP</u>	+ ALCON PHARMS LTD	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N021980</u>	<u>001</u>	Mar 28, 2006

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

+ ALCON	0.1%	N019079	001	Feb 11, 1986
---------	------	---------	-----	--------------

FLUOROURACIL

CREAM; TOPICAL

EFUDEX

<u>AB</u> + VALEANT PHARM INTL	<u>5%</u>	<u>N016831</u>	<u>003</u>
--------------------------------	-----------	----------------	------------

PREScription DRUG PRODUCT LIST

3 - 187 (of 424)

FLUOROURACIL

CREAM; TOPICAL

FLUOROURACIL

<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
	CARAC				
+	SANOFI AVENTIS US	0.5%	N020985	001	Oct 27, 2000
	FLUOROPLEX				
+	ALLERGAN HERBERT	1%	N016988	001	

INJECTABLE; INJECTION

FLUOROURACIL

<u>AP</u>	+	APP PHARMS	<u>500MG/10ML (50MG/ML)</u>	<u>A040279</u>	<u>002</u>	Sep 30, 1998
<u>AP</u>	+		<u>1GM/20ML (50MG/ML)</u>	<u>A040279</u>	<u>001</u>	Sep 30, 1998
<u>AP</u>	+		<u>2.5GM/50ML (50MG/ML)</u>	<u>A040278</u>	<u>001</u>	Sep 30, 1998
<u>AP</u>	+		<u>5GM/100ML (50MG/ML)</u>	<u>A040278</u>	<u>002</u>	Sep 30, 1998
<u>AP</u>	+	BIONICHE PHARMA	<u>500MG/10ML (50MG/ML)</u>	<u>A040743</u>	<u>002</u>	Apr 26, 2007
<u>AP</u>	+		<u>1GM/20ML (50MG/ML)</u>	<u>A040743</u>	<u>001</u>	Apr 26, 2007
<u>AP</u>	+		<u>2.5GM/50ML (50MG/ML)</u>	<u>A040798</u>	<u>002</u>	Apr 26, 2007
<u>AP</u>	+		<u>5GM/100ML (50MG/ML)</u>	<u>A040798</u>	<u>001</u>	Apr 26, 2007
<u>AP</u>		EBEWE PHARMA	<u>500MG/10ML (50MG/ML)</u>	<u>A040772</u>	<u>001</u>	Aug 11, 2008
<u>AP</u>		SANDOZ	<u>2.5GM/50ML (50MG/ML)</u>	<u>A091299</u>	<u>001</u>	May 02, 2011
<u>AP</u>			<u>5GM/100ML (50MG/ML)</u>	<u>A091299</u>	<u>002</u>	May 02, 2011
<u>AP</u>	+	TEVA PARENTERAL	<u>500MG/10ML (50MG/ML)</u>	<u>A040333</u>	<u>001</u>	Jan 27, 2000
<u>AP</u>	+		<u>2.5GM/50ML (50MG/ML)</u>	<u>A040334</u>	<u>001</u>	Feb 25, 2000
<u>AP</u>	+		<u>5GM/100ML (50MG/ML)</u>	<u>A040334</u>	<u>002</u>	Feb 25, 2000

SOLUTION; TOPICAL

EFUDEX

<u>AT</u>	+ VALEANT PHARM INTL	<u>2%</u>	<u>N016831</u>	<u>001</u>
<u>AT</u>	+	<u>5%</u>	<u>N016831</u>	<u>002</u>
<u>FLUOROURACIL</u>				
<u>AT</u>	TARO	<u>2%</u>	<u>A076526</u>	<u>001</u>
<u>AT</u>		<u>5%</u>	<u>A076526</u>	<u>002</u>

NOV 05, 2003

NOV 05, 2003

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 40MG BASE</u>	<u>A090223</u>	<u>003</u>	Mar 19, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 40MG BASE</u>	<u>A078619</u>	<u>003</u>	Jan 31, 2008
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 40MG BASE</u>	<u>A075465</u>	<u>003</u>	Aug 02, 2001
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 40MG BASE</u>	<u>A075245</u>	<u>003</u>	Sep 28, 2004
<u>AB</u>	MYLAN	<u>EQ 40MG BASE</u>	<u>A075207</u>	<u>003</u>	May 25, 2007
<u>AB</u>	RANBAXY	<u>EQ 40MG BASE</u>	<u>A076990</u>	<u>001</u>	Dec 13, 2004
<u>AB</u>	SANDOZ	<u>EQ 40MG BASE</u>	<u>A075049</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>	TEVA	<u>EQ 40MG BASE</u>	<u>A075452</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>	WOCKHARDT	<u>EQ 40MG BASE</u>	<u>A078143</u>	<u>003</u>	Jan 16, 2008

BROZAC

AB + LILLY FROZAC EO 40MG BASE NO18936-003 Jun 15 1999

ELI LIQUETINE HYDROCHLORIDE

FLUOXETINE HYDROCHLORIDE				
AB1	ALEMBIC PHARMS LTD	EQ 10MG BASE	A090223	001
AB1		EQ 20MG BASE	A090223	002
AB1	ALPHAPHARM	EQ 10MG BASE	A075577	001
AB1		EQ 20MG BASE	A075577	002
AB1	AUROBINDO PHARMA	EQ 10MG BASE	A078619	001
AB1		EQ 20MG BASE	A078619	002
AB1	BARR	EQ 10MG BASE	A074803	002
AB1		EQ 20MG BASE	A074803	001
AB1	BEIJING DOUBLE CRANE	EQ 10MG BASE	A076165	001
AB1		EQ 20MG BASE	A076165	002
AB1	CARLSBAD	EQ 10MG BASE	A076022	001
AB1		EQ 20MG BASE	A076022	002

PRESCRIPTION DRUG PRODUCT LIST

3 - 188 (of 424)

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB1</u>	DR REDDYS LABS INC	<u>EQ 10MG BASE</u>	<u>A075465 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075465 002</u>	Jan 29, 2002
<u>AB1</u>	IVAX SUB TEVA PHARMS	<u>EQ 10MG BASE</u>	<u>A075245 002</u>	Jan 31, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075245 001</u>	Jan 31, 2002
<u>AB1</u>	LANDELA PHARM	<u>EQ 10MG BASE</u>	<u>A075464 001</u>	Jan 30, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075464 002</u>	Jan 30, 2002
<u>AB1</u>	MALLINCKRODT	<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>	MYLAN	<u>EQ 10MG BASE</u>	<u>A075207 001</u>	Jan 30, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075207 002</u>	Jan 30, 2002
<u>AB1</u>	PLIVA	<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
<u>AB1</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>A075049 001</u>	Aug 02, 2001
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075049 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>	WOCKHARDT	<u>EQ 10MG BASE</u>	<u>A078143 001</u>	Jan 16, 2008
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A078143 002</u>	Jan 16, 2008
	<u>PROZAC</u>			
<u>AB1</u>	LILLY	<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
	<u>FLUOXETINE HYDROCHLORIDE</u>			
<u>AB2</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A078045 001</u>	Nov 17, 2008
<u>AB2</u>		<u>EQ 20MG BASE</u>	<u>A078045 002</u>	Nov 17, 2008
<u>AB2</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>A077469 001</u>	Nov 17, 2008
<u>AB2</u>		<u>EQ 20MG BASE</u>	<u>A077469 002</u>	Nov 17, 2008
<u>AB2</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A076287 001</u>	May 20, 2008
<u>AB2</u>		<u>EQ 20MG BASE</u>	<u>A076287 002</u>	May 20, 2008
	<u>SARAFEM</u>			
<u>AB2</u>	LILLY	<u>EQ 10MG BASE</u>	<u>N018936 007</u>	Jul 06, 2000
<u>AB2</u> +		<u>EQ 20MG BASE</u>	<u>N018936 008</u>	Jul 06, 2000

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
<u>AB</u>	<u>PROZAC WEEKLY</u>			
<u>AB</u> +	LILLY	<u>EQ 90MG BASE</u>	<u>N021235 001</u>	Feb 26, 2001

SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AA</u>	AUROBINDO PHARM	<u>EQ 20MG BASE/5ML</u>	<u>A079209 001</u>	Mar 20, 2009
<u>AA</u>	LANNETT	<u>EQ 20MG BASE/5ML</u>	<u>A076458 001</u>	May 14, 2004
<u>AA</u>	MALLINCKRODT	<u>EQ 20MG BASE/5ML</u>	<u>A075920 001</u>	Jan 29, 2002
<u>AA</u>	NOVEX	<u>EQ 20MG BASE/5ML</u>	<u>A075292 001</u>	Feb 07, 2002
<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>	TEVA	<u>EQ 20MG BASE/5ML</u>	<u>A075506 001</u>	Aug 02, 2001
<u>AA</u>	WOCKHARDT	<u>EQ 20MG BASE/5ML</u>	<u>A075514 001</u>	Aug 29, 2002

TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS INC	<u>EQ 10MG BASE</u>	<u>A076006 001</u>	Jan 30, 2002
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A075755 001</u>	Aug 02, 2001
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075872 001</u>	Jan 29, 2002
	<u>FLUOXETINE HYDROCHLORIDE</u>			
	EDGE MONT PHARMS LLC	<u>EQ 60MG BASE</u>	<u>N202133 001</u>	Oct 06, 2011
+ MYLAN		<u>EQ 20MG BASE</u>	<u>A075755 002</u>	Aug 02, 2001
	<u>SARAFEM</u>			
	WARNER CHILCOTT	<u>EQ 10MG BASE</u>	<u>N021860 001</u>	May 19, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 189 (of 424)

FLUOXETINE HYDROCHLORIDE

TABLET; ORAL SARAFEM			
WARNER CHILCOTT	EQ 15MG BASE	N021860 002	May 19, 2006
+	EQ 20MG BASE	N021860 003	May 19, 2006

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL SYMBYAX			
LILLY	EQ 25MG BASE;EQ 3MG BASE	N021520 001	Apr 09, 2007
	EQ 25MG BASE;EQ 6MG BASE	N021520 002	Dec 24, 2003
	EQ 25MG BASE;EQ 12MG BASE	N021520 004	Dec 24, 2003
+	EQ 50MG BASE;EQ 6MG BASE	N021520 003	Dec 24, 2003
	EQ 50MG BASE;EQ 12MG BASE	N021520 005	Dec 24, 2003

FLUOXYMESTERONE

TABLET; ORAL FLUOXYMESTERONE			
+ USL PHARMA	10MG	A088342 001	Oct 21, 1983

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION <u>FLUPHENAZINE DECANOATE</u>			
<u>AO</u> + APP PHARMS	<u>25MG/ML</u>	<u>A071413 001</u>	Jul 14, 1987
<u>AO</u> BEDFORD	<u>25MG/ML</u>	<u>A074531 001</u>	Aug 30, 1996
<u>AO</u> CLARIS LIFESCIENCES	<u>25MG/ML</u>	<u>A075918 001</u>	Aug 17, 2001

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			
+ APP PHARMS	2.5MG/ML	A089556 001	Apr 16, 1987

TABLET; ORAL <u>FLUPHENAZINE HYDROCHLORIDE</u>			
<u>AB</u> LANNETT	<u>1MG</u>	<u>A089740 001</u>	Aug 25, 1988
<u>AB</u>	<u>2.5MG</u>	<u>A089741 001</u>	Aug 25, 1988
<u>AB</u>	<u>5MG</u>	<u>A089742 001</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>A089583 001</u>	Oct 16, 1987
<u>AB</u>	<u>2.5MG</u>	<u>A089584 001</u>	Oct 16, 1987
<u>AB</u>	<u>5MG</u>	<u>A089585 001</u>	Oct 16, 1987
<u>AB</u>	<u>10MG</u>	<u>A089586 001</u>	Oct 16, 1987

FLURANDRENOLIDE

CREAM; TOPICAL CORDRAN SP			
WATSON PHARMS	0.025%	N012806 003	

PRESCRIPTION DRUG PRODUCT LIST

3 - 190 (of 424)

FLURANDRENOLIDE

LOTION; TOPICAL CORDRAN			
+ WATSON LABS	0.05%	N013790	001
TAPE; TOPICAL CORDRAN			
+ WATSON PHARMS	0.004MG/SQ CM	N016455	001

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL FLURAZEPAM HYDROCHLORIDE			
AB MYLAN PHARMS INC <u>15MG</u>		A070345	002 Nov 27, 1985
AB + <u>30MG</u>		A070345	001 Nov 27, 1985
AB WATSON LABS <u>15MG</u>		A071205	001 Nov 25, 1986
AB <u>30MG</u>		A071068	001 Nov 25, 1986
AB <u>30MG</u>		A072369	001 Mar 30, 1989
AB WEST WARD <u>15MG</u>		A071107	001 Dec 08, 1986
AB <u>30MG</u>		A071108	001 Dec 08, 1986

FLURBIPROFEN

TABLET; ORAL ANSAID			
AB PHARMACIA AND UPJOHN <u>50MG</u>		N018766	002 Oct 31, 1988
AB + <u>100MG</u>		N018766	003 Oct 31, 1988
FLURBIPROFEN			
AB CARACO <u>50MG</u>		A075058	001 Apr 27, 2001
AB <u>100MG</u>		A075058	002 Apr 27, 2001
AB MYLAN <u>50MG</u>		A074358	001 Jun 20, 1994
AB <u>100MG</u>		A074358	002 Jun 20, 1994
AB TEVA <u>100MG</u>		A074431	001 May 31, 1995

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC FLURBIPROFEN SODIUM			
AT BAUSCH AND LOMB <u>0.03%</u>		A074447	001 Jan 04, 1995
OCUFEN			
AT + ALLERGAN <u>0.03%</u>		N019404	001 Dec 31, 1986

FLUTAMIDE

CAPSULE; ORAL FLUTAMIDE			
AB IVAX SUB TEVA PHARMS <u>125MG</u>		A075780	001 Sep 19, 2001
AB MYLAN <u>125MG</u>		A076224	001 May 09, 2003
AB PAR PHARM <u>125MG</u>		A075298	001 Sep 18, 2001
AB + SANDOZ <u>125MG</u>		A075818	001 Sep 18, 2001
AB WATSON LABS <u>125MG</u>		A075820	001 Sep 18, 2001

FLUTICASONE FUROATE

SPRAY, METERED; NASAL VERAMYST			
+ GLAXOSMITHKLINE	0.0275MG/INH	N022051	001 Apr 27, 2007

FLUTICASONE PROPIONATE

AEROSOL, METERED; INHALATION FLOVENT HFA			
+ GLAXO GRP LTD	0.044MG/INH	N021433	003 May 14, 2004
+	0.11MG/INH	N021433	002 May 14, 2004
+	0.22MG/INH	N021433	001 May 14, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 191 (of 424)

FLUTICASONE PROPIONATE

CREAM; TOPICAL

CUTIVATE

<u>AB</u>	+ ALTANA	<u>0.05%</u>	<u>N019958</u>	<u>001</u>	Dec 18, 1990
<u>FLUTICASONE PROPIONATE</u>					
<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A076451</u>	<u>001</u>	May 14, 2004
<u>AB</u>	G AND W LABS	<u>0.05%</u>	<u>A077055</u>	<u>001</u>	Jun 30, 2006
<u>AB</u>	NESHER PHARMS	<u>0.05%</u>	<u>A076865</u>	<u>001</u>	Sep 10, 2004
<u>AB</u>	PERRIGO NEW YORK	<u>0.05%</u>	<u>A076793</u>	<u>001</u>	May 14, 2004
<u>AB</u>	TOLMAR	<u>0.05%</u>	<u>A076633</u>	<u>001</u>	May 14, 2004

LOTION; TOPICAL

CUTIVATE

<u>AB</u>	+ FOUGERA PHARMS	<u>0.05%</u>	<u>N021152</u>	<u>001</u>	Mar 31, 2005
<u>FLUTICASONE PROPIONATE</u>					
<u>AB</u>	GLENMARK GENERICS	<u>0.05%</u>	<u>A090759</u>	<u>001</u>	May 02, 2011

OINTMENT; TOPICAL

CUTIVATE

<u>AB</u>	+ FOUGERA PHARMS	<u>0.005%</u>	<u>N019957</u>	<u>001</u>	Dec 14, 1990
<u>FLUTICASONE PROPIONATE</u>					
<u>AB</u>	ALTANA	<u>0.005%</u>	<u>A076300</u>	<u>001</u>	May 14, 2004
<u>AB</u>	G AND W LABS	<u>0.005%</u>	<u>A077168</u>	<u>001</u>	Mar 03, 2006
<u>AB</u>	PERRIGO NEW YORK	<u>0.005%</u>	<u>A076668</u>	<u>001</u>	May 14, 2004

POWDER; INHALATION

FLOVENT DISKUS 100

+ GLAXOSMITHKLINE 0.1MG/INH

N020833 002 Sep 29, 2000

FLOVENT DISKUS 250

+ GLAXOSMITHKLINE 0.25MG/INH

N020833 003 Sep 29, 2000

FLOVENT DISKUS 50

+ GLAXOSMITHKLINE 0.05MG/INH

N020833 001 Sep 29, 2000

SPRAY, METERED; NASAL

FLONASE

<u>AB</u>	+ GLAXOSMITHKLINE	<u>0.05MG/SPRAY</u>	<u>N020121</u>	<u>001</u>	Oct 19, 1994
<u>FLUTICASONE PROPIONATE</u>					
<u>AB</u>	APOTEX INC	<u>0.05MG/SPRAY</u>	<u>A077538</u>	<u>001</u>	Sep 12, 2007
<u>AB</u>	HI TECH PHARMA	<u>0.05MG/SPRAY</u>	<u>A077570</u>	<u>001</u>	Jan 16, 2008
<u>AB</u>	ROXANE	<u>0.05MG/SPRAY</u>	<u>A076504</u>	<u>001</u>	Feb 22, 2006
<u>AB</u>	WOCKHARDT	<u>0.05MG/SPRAY</u>	<u>A078492</u>	<u>001</u>	Jan 09, 2012

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

ADVAIR HFA

+ GLAXOSMITHKLINE 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

+ GLAXOSMITHKLINE 0.1MG/INH;EQ 0.05MG BASE/INH

N021077 001 Aug 24, 2000

ADVAIR DISKUS 250/50

+ GLAXOSMITHKLINE 0.25MG/INH;EQ 0.05MG BASE/INH

N021077 002 Aug 24, 2000

ADVAIR DISKUS 500/50

+ GLAXOSMITHKLINE 0.5MG/INH;EQ 0.05MG BASE/INH

N021077 003 Aug 24, 2000

FLUVASTATIN SODIUM

CAPSULE; ORAL

LESCOL

NOVARTIS EQ 20MG BASE

N020261 001 Dec 31, 1993

+ EQ 40MG BASE

N020261 002 Dec 31, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 192 (of 424)

FLUVASTATIN SODIUM

TABLET, EXTENDED RELEASE; ORAL
 LESCOL XL
 + NOVARTIS 80MG N021192 001 Oct 06, 2000

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL
 LUVOX CR
 JAZZ 100MG N022033 001 Feb 28, 2008
 + 150MG N022033 002 Feb 28, 2008

TABLET; ORAL

FLUVOXAMINE MALEATE

<u>AB</u>	APOTEX	<u>25MG</u>	<u>A075902</u> <u>001</u>	May 07, 2001
<u>AB</u>		<u>50MG</u>	<u>A075902</u> <u>002</u>	May 07, 2001
<u>AB</u>		<u>100MG</u>	<u>A075902</u> <u>003</u>	May 07, 2001
<u>AB</u>	BARR	<u>25MG</u>	<u>A075897</u> <u>001</u>	Jan 25, 2001
<u>AB</u>		<u>50MG</u>	<u>A075897</u> <u>002</u>	Jan 25, 2001
<u>AB</u>		<u>100MG</u>	<u>A075897</u> <u>003</u>	Jan 25, 2001
<u>AB</u>	CARACO	<u>25MG</u>	<u>A075900</u> <u>001</u>	Feb 23, 2006
<u>AB</u>		<u>50MG</u>	<u>A075900</u> <u>002</u>	Feb 23, 2006
<u>AB</u>		<u>100MG</u>	<u>A075900</u> <u>003</u>	Feb 23, 2006
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A075889</u> <u>001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>A075889</u> <u>002</u>	Nov 29, 2000
<u>AB</u>		<u>100MG</u>	<u>A075889</u> <u>003</u>	Nov 29, 2000
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A075888</u> <u>001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>A075888</u> <u>002</u>	Nov 29, 2000
<u>AB</u>	+	<u>100MG</u>	<u>A075888</u> <u>003</u>	Nov 29, 2000
<u>AB</u>	TEVA	<u>25MG</u>	<u>A075893</u> <u>001</u>	Sep 10, 2002
<u>AB</u>		<u>50MG</u>	<u>A075893</u> <u>002</u>	Sep 10, 2002
<u>AB</u>		<u>100MG</u>	<u>A075893</u> <u>003</u>	Sep 10, 2002
	<u>LUVOX</u>			
<u>AB</u>	ANI PHARMS	<u>25MG</u>	<u>N021519</u> <u>001</u>	Dec 20, 2007
<u>AB</u>		<u>50MG</u>	<u>N021519</u> <u>002</u>	Dec 20, 2007
<u>AB</u>		<u>100MG</u>	<u>N021519</u> <u>003</u>	Dec 20, 2007

FOLIC ACID

INJECTABLE; INJECTION
 FOLIC ACID
 + APP PHARMS 5MG/ML A089202 001 Feb 18, 1986

TABLET; ORAL

FOLIC ACID

<u>AA</u>	+ AMNEAL PHARM	<u>1MG</u>	<u>A040625</u> <u>001</u>	Jul 21, 2005
<u>AA</u>	CONTRACT PHARMACAL	<u>1MG</u>	<u>A085061</u> <u>001</u>	
<u>AA</u>	EXCELLIUM	<u>1MG</u>	<u>A040796</u> <u>001</u>	Jan 12, 2009
<u>AA</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A090035</u> <u>001</u>	Jun 09, 2009
<u>AA</u>	JUBILANT CADISTA	<u>1MG</u>	<u>A040514</u> <u>001</u>	Jun 14, 2005
<u>AA</u>	VINTAGE	<u>1MG</u>	<u>A040756</u> <u>001</u>	Jun 04, 2010
<u>AA</u>	+	<u>WATSON LABS</u>	<u>1MG</u>	
<u>AA</u>	WEST WARD	<u>1MG</u>	<u>A080680</u> <u>001</u>	
			<u>A080600</u> <u>001</u>	

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS
 FOLLISTIM AQ
 + ORGANON USA INC 75 IU/0.5ML N021273 001 Aug 26, 2005
 + 150 IU/0.5ML N021273 002 Aug 26, 2005
 + 300 IU/0.36ML N021211 001 Mar 23, 2004
 + 600 IU/0.72ML N021211 002 Mar 23, 2004
 + 900 IU/1.08ML N021211 004 Feb 11, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 193 (of 424)

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS		
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 PEN		
+ EMD SERONO	300 IU/0.5ML	N021684 001 May 25, 2004
+ 450 IU/0.75ML		N021684 002 May 25, 2004
+ 900 IU/1.5ML		N021684 003 May 25, 2004

FOMEPIZOLE

INJECTABLE; INJECTION		
<u>ANTIZOL</u>		

<u>AP</u> + PALADIN LABS	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>N020696 001</u> Dec 04, 1997
<u>FOMEPIZOLE</u>		
<u>AP</u> BIONICHE PHARMA USA	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A079033 001</u> Apr 07, 2009
<u>AP</u> LUITPOLD	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078368 001</u> Dec 14, 2007
<u>AP</u> NAVINTA LLC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078537 001</u> Mar 06, 2008
<u>AP</u> SYNERX PHARMA	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078639 001</u> Mar 03, 2008

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS		
<u>ARIIXTRA</u>		

<u>AP</u> + GLAXOSMITHKLINE	<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
<u>FONDAPARINUX SODIUM</u>		
<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

FORMOTEROL FUMARATE

POWDER; INHALATION		
FORADIL		

+ NOVARTIS	0.012MG/INH	N020831 001 Feb 16, 2001
------------	-------------	--------------------------

SOLUTION; INHALATION		
PERFOROMIST		

+ DEY PHARMA	0.02MG/2ML	N022007 001 May 11, 2007
--------------	------------	--------------------------

FORMOTEROL FUMARATE; MOMETASONE FUROATE

AEROSOL, METERED; INHALATION		
DULERA		

+ SCHERING	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		
LEXIVA		

+ VIIV HLTHCARE	EQ 700MG BASE	N021548 001 Oct 20, 2003
-----------------	---------------	--------------------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 194 (of 424)

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS EMEND			
+ MERCK AND CO INC	EQ 115MG BASE/VIAL	N022023 001	Jan 25, 2008
+ EQ 150MG BASE/VIAL		N022023 002	Nov 12, 2010

FOSCARNET SODIUM

INJECTABLE; INJECTION <u>FOSCARNET SODIUM</u>			
<u>AP HOSPIRA</u>	<u>2.4GM/100ML</u>	<u>A077174 001</u>	May 31, 2005
<u>FOSCAVIR</u>			
<u>AP + CLINIGEN HLTHCARE</u>	<u>2.4GM/100ML</u>	<u>N020068 001</u>	Sep 27, 1991

FOSFOMYCIN TROMETHAMINE

FOR SUSPENSION; ORAL MONUROL			
+ ZAMBON SPA	EQ 3GM BASE/PACKET	N050717 001	Dec 19, 1996

FOSINOPRIL SODIUM

TABLET; ORAL <u>FOSINOPRIL SODIUM</u>			
<u>AB APOTEX INC</u>	<u>10MG</u>	<u>A076906 001</u>	May 17, 2005
	<u>20MG</u>	<u>A076906 002</u>	May 17, 2005
	<u>40MG</u>	<u>A076906 003</u>	May 17, 2005
<u>AB AUROBINDO PHARMA LTD</u>	<u>10MG</u>	<u>A091163 001</u>	Mar 30, 2011
	<u>20MG</u>	<u>A091163 002</u>	Mar 30, 2011
	<u>40MG</u>	<u>A091163 003</u>	Mar 30, 2011
<u>AB INVAGEN PHARMS</u>	<u>10MG</u>	<u>A077222 001</u>	Apr 20, 2005
	<u>20MG</u>	<u>A077222 002</u>	Apr 20, 2005
	<u>40MG</u>	<u>A077222 003</u>	Apr 20, 2005
<u>AB RANBAXY</u>	<u>10MG</u>	<u>A076580 001</u>	Apr 23, 2004
	<u>20MG</u>	<u>A076580 002</u>	Apr 23, 2004
	<u>40MG</u>	<u>A076580 003</u>	Apr 23, 2004
<u>AB SANDOZ</u>	<u>10MG</u>	<u>A076483 001</u>	Apr 23, 2004
	<u>20MG</u>	<u>A076483 002</u>	Apr 23, 2004
	<u>40MG</u>	<u>A076483 003</u>	Apr 23, 2004
<u>AB TEVA</u>	<u>10MG</u>	<u>A076139 001</u>	Nov 25, 2003
	<u>20MG</u>	<u>A076139 002</u>	Nov 25, 2003
	<u>40MG</u>	<u>A076139 003</u>	Nov 25, 2003
<u>AB + WATSON LABS</u>	<u>10MG</u>	<u>A076987 001</u>	Dec 23, 2004
	<u>10MG</u>	<u>A077531 001</u>	Aug 31, 2006
	<u>20MG</u>	<u>A076987 002</u>	Dec 23, 2004
	<u>20MG</u>	<u>A077531 002</u>	Aug 31, 2006
	<u>40MG</u>	<u>A076987 003</u>	Dec 23, 2004
	<u>40MG</u>	<u>A077531 003</u>	Aug 31, 2006
<u>AB WATSON LABS FLORIDA</u>	<u>10MG</u>	<u>A076620 001</u>	Oct 15, 2004
	<u>20MG</u>	<u>A076620 002</u>	Oct 15, 2004
	<u>40MG</u>	<u>A076620 003</u>	Oct 15, 2004
<u>MONOPRIL</u>			
<u>AB BRISTOL MYERS SQUIBB</u>	<u>10MG</u>	<u>N019915 002</u>	May 16, 1991
	<u>20MG</u>	<u>N019915 003</u>	May 16, 1991
<u>AB +</u>	<u>40MG</u>	<u>N019915 004</u>	Mar 28, 1995

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL <u>FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE</u>			
<u>AB AUROBINDO PHARMA</u>	<u>10MG;12.5MG</u>	<u>A079245 001</u>	Jul 09, 2009
	<u>20MG;12.5MG</u>	<u>A079245 002</u>	Jul 09, 2009
<u>AB INVAGEN PHARMS</u>	<u>10MG;12.5MG</u>	<u>A090228 001</u>	Jul 09, 2009
	<u>20MG;12.5MG</u>	<u>A090228 002</u>	Jul 09, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 195 (of 424)

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	MYLAN	<u>10MG;12.5MG</u>	<u>A077705</u>	<u>001</u>	Aug 14, 2006
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A077705</u>	<u>002</u>	Aug 14, 2006
<u>AB</u>	RANBAXY	<u>10MG;12.5MG</u>	<u>A076739</u>	<u>001</u>	Dec 17, 2004
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076739</u>	<u>002</u>	Dec 17, 2004
<u>AB</u>	SANDOZ	<u>10MG;12.5MG</u>	<u>A076961</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076961</u>	<u>002</u>	Sep 28, 2005
<u>AB</u>	WATSON LABS FLORIDA	<u>10MG;12.5MG</u>	<u>A076608</u>	<u>001</u>	Dec 03, 2004
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076608</u>	<u>002</u>	Dec 03, 2004

FOSINOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	EMCURE PHARMS USA	<u>10MG;12.5MG</u>	<u>A079025</u>	<u>001</u>	Sep 17, 2010
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A079025</u>	<u>002</u>	Sep 17, 2010

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

<u>AP</u>	APOTEX INC	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078126</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>	+ APP PHARMS	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078052</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077989</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>	BEDFORD	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077481</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078765</u>	<u>001</u>	Dec 02, 2009
<u>AP</u>	HOSPIRA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078158</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>	LUITPOLD	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078277</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>		<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A090099</u>	<u>001</u>	May 13, 2010
<u>AP</u>	PFIZER	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078476</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>	STRIDES ARCOLAB	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078736</u>	<u>001</u>	Jun 08, 2010
<u>AP</u>	SUN PHARMA GLOBAL	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078417</u>	<u>001</u>	Mar 18, 2008
<u>AP</u>	TEVA PARENTERAL	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A076886</u>	<u>001</u>	Aug 06, 2007
<u>AP</u>	WOCKHARDT	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078137</u>	<u>001</u>	Aug 06, 2007

FOSPROPOFOL DISODIUM

SOLUTION; INTRAVENOUS

LUSEDRA

+ EISAI INC	1050MG/30ML (35MG/ML)	N022244	001	Dec 12, 2008
-------------	-----------------------	---------	-----	--------------

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

+ ENDO PHARMS	EQ 2.5MG BASE	N021006	001	Nov 08, 2001
---------------	---------------	---------	-----	--------------

FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FASLODEX

+ ASTRAZENECA	50MG/ML	N021344	001	Apr 25, 2002
---------------	---------	---------	-----	--------------

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>	APP PHARMS	<u>10MG/ML</u>	<u>N018902</u>	<u>001</u>	May 22, 1984
<u>AP</u>	HOSPIRA	<u>10MG/ML</u>	<u>A070578</u>	<u>001</u>	Jul 08, 1987
<u>AP</u>		<u>10MG/ML</u>	<u>A075241</u>	<u>001</u>	May 28, 1999
<u>AP</u>		<u>10MG/ML</u>	<u>N018667</u>	<u>001</u>	May 28, 1982
<u>AP</u>	INTL MEDICATION	<u>10MG/ML</u>	<u>N018025</u>	<u>001</u>	
<u>AP</u>	+ LUITPOLD	<u>10MG/ML</u>	<u>N018579</u>	<u>001</u>	Nov 30, 1983

PRESCRIPTION DRUG PRODUCT LIST

3 - 196 (of 424)

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>	WOCKHARDT	<u>10MG/ML</u>	<u>A077941</u> <u>001</u>	Mar 22, 2007
	SOLUTION; ORAL			
		<u>FUROSEMIDE</u>		
<u>AA</u>	+ ROXANE	<u>10MG/ML</u>	<u>A070434</u> <u>001</u>	Apr 22, 1987
<u>AA</u>	WOCKHARDT	<u>10MG/ML</u>	<u>A070655</u> <u>001</u>	Oct 02, 1987
	FUROSEMIDE			
	ROXANE	40MG/5ML	A070433 001	Apr 22, 1987
	TABLET; ORAL			
		<u>FUROSEMIDE</u>		
<u>AB</u>	DAVA PHARMS INC	<u>20MG</u>	<u>N018415</u> <u>001</u>	Jul 27, 1982
<u>AB</u>		<u>40MG</u>	<u>N018415</u> <u>002</u>	Jul 27, 1982
<u>AB</u>		<u>80MG</u>	<u>N018415</u> <u>003</u>	Nov 26, 1984
<u>AB</u>	EXCELLIUM	<u>20MG</u>	<u>A077293</u> <u>001</u>	Nov 09, 2005
<u>AB</u>		<u>40MG</u>	<u>A077293</u> <u>002</u>	Nov 09, 2005
<u>AB</u>		<u>80MG</u>	<u>A077293</u> <u>003</u>	Nov 09, 2005
<u>AB</u>	IPCA LABS LTD	<u>20MG</u>	<u>A078010</u> <u>001</u>	Sep 18, 2006
<u>AB</u>		<u>40MG</u>	<u>A078010</u> <u>002</u>	Sep 18, 2006
<u>AB</u>		<u>80MG</u>	<u>A078010</u> <u>003</u>	Sep 18, 2006
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>20MG</u>	<u>N018413</u> <u>001</u>	Nov 30, 1983
<u>AB</u>		<u>40MG</u>	<u>N018413</u> <u>002</u>	Nov 30, 1983
<u>AB</u>	MYLAN	<u>20MG</u>	<u>N018487</u> <u>001</u>	
<u>AB</u>		<u>40MG</u>	<u>N018487</u> <u>002</u>	
<u>AB</u>		<u>80MG</u>	<u>A070082</u> <u>001</u>	Oct 29, 1986
<u>AB</u>	ROXANE	<u>20MG</u>	<u>N018823</u> <u>001</u>	Nov 10, 1983
<u>AB</u>		<u>40MG</u>	<u>N018823</u> <u>002</u>	Nov 10, 1983
<u>AB</u>		<u>80MG</u>	<u>A070086</u> <u>001</u>	Jan 24, 1986
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>N018569</u> <u>002</u>	
<u>AB</u>		<u>40MG</u>	<u>N018569</u> <u>001</u>	
<u>AB</u>		<u>80MG</u>	<u>N018569</u> <u>005</u>	Aug 14, 1984
<u>AB</u>	VINTAGE PHARMS	<u>20MG</u>	<u>A076796</u> <u>001</u>	Mar 26, 2004
<u>AB</u>		<u>40MG</u>	<u>A076796</u> <u>002</u>	Mar 26, 2004
<u>AB</u>		<u>80MG</u>	<u>A076796</u> <u>003</u>	Mar 26, 2004
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>A070412</u> <u>001</u>	Feb 26, 1986
<u>AB</u>		<u>40MG</u>	<u>A070449</u> <u>001</u>	Nov 22, 1985
<u>AB</u>		<u>80MG</u>	<u>A071379</u> <u>001</u>	Jan 02, 1987
<u>AB</u>		<u>80MG</u>	<u>A070450</u> <u>001</u>	Nov 22, 1985
<u>AB</u>		<u>80MG</u>	<u>A070528</u> <u>001</u>	Jan 07, 1986
<u>AB</u>		<u>80MG</u>	<u>A071594</u> <u>001</u>	Feb 09, 1988
	<u>LASIX</u>			
<u>AB</u>	SANOFI AVENTIS US	<u>20MG</u>	<u>N016273</u> <u>002</u>	
<u>AB</u>		<u>40MG</u>	<u>N016273</u> <u>001</u>	
<u>AB</u>	+	<u>80MG</u>	<u>N016273</u> <u>003</u>	

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A075350</u> <u>001</u>	Sep 12, 2003
<u>AB</u>		<u>300MG</u>	<u>A075350</u> <u>002</u>	Sep 12, 2003
<u>AB</u>		<u>400MG</u>	<u>A075350</u> <u>003</u>	Sep 12, 2003
<u>AB</u>	ALKEM	<u>100MG</u>	<u>A090858</u> <u>001</u>	Dec 17, 2010
<u>AB</u>		<u>300MG</u>	<u>A090858</u> <u>002</u>	Dec 17, 2010
<u>AB</u>		<u>400MG</u>	<u>A090858</u> <u>003</u>	Dec 17, 2010
<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>	<u>A078428</u> <u>001</u>	Jul 25, 2007
<u>AB</u>		<u>300MG</u>	<u>A078428</u> <u>002</u>	Jul 25, 2007
<u>AB</u>		<u>400MG</u>	<u>A078428</u> <u>003</u>	Jul 25, 2007
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A075360</u> <u>001</u>	Apr 06, 2005
<u>AB</u>		<u>300MG</u>	<u>A075360</u> <u>002</u>	Apr 06, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 197 (of 424)

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>	APOTEX INC	<u>400MG</u>	<u>A075360</u>	<u>003</u>	Apr 06, 2005
<u>AB</u>	AUROBINDO PHARM	<u>100MG</u>	<u>A078787</u>	<u>001</u>	Jan 31, 2008
<u>AB</u>		<u>300MG</u>	<u>A078787</u>	<u>002</u>	Jan 31, 2008
<u>AB</u>		<u>400MG</u>	<u>A078787</u>	<u>003</u>	Jan 31, 2008
<u>AB</u>	HIKMA	<u>100MG</u>	<u>A078150</u>	<u>001</u>	Sep 25, 2007
<u>AB</u>		<u>300MG</u>	<u>A078150</u>	<u>002</u>	Sep 25, 2007
<u>AB</u>		<u>400MG</u>	<u>A078150</u>	<u>003</u>	Sep 25, 2007
<u>AB</u>	INVAGEN PHARMS	<u>100MG</u>	<u>A090705</u>	<u>001</u>	Dec 30, 2009
<u>AB</u>		<u>300MG</u>	<u>A090705</u>	<u>002</u>	Dec 30, 2009
<u>AB</u>		<u>400MG</u>	<u>A090705</u>	<u>003</u>	Dec 30, 2009
<u>AB</u>	MARKSANS PHARMA	<u>100MG</u>	<u>A090007</u>	<u>001</u>	Jul 21, 2011
<u>AB</u>		<u>300MG</u>	<u>A090007</u>	<u>002</u>	Jul 21, 2011
<u>AB</u>		<u>400MG</u>	<u>A090007</u>	<u>003</u>	Jul 21, 2011
<u>AB</u>	MYLAN	<u>100MG</u>	<u>A090158</u>	<u>001</u>	Feb 14, 2011
<u>AB</u>		<u>300MG</u>	<u>A090158</u>	<u>002</u>	Feb 14, 2011
<u>AB</u>		<u>400MG</u>	<u>A090158</u>	<u>003</u>	Feb 14, 2011
<u>AB</u>	RANBAXY	<u>100MG</u>	<u>A076606</u>	<u>001</u>	Oct 07, 2005
<u>AB</u>		<u>300MG</u>	<u>A076606</u>	<u>002</u>	Oct 07, 2005
<u>AB</u>		<u>400MG</u>	<u>A076606</u>	<u>003</u>	Oct 07, 2005
<u>AB</u>	SUN PHARM INDS LTD	<u>100MG</u>	<u>A077242</u>	<u>001</u>	Aug 24, 2006
<u>AB</u>		<u>300MG</u>	<u>A077242</u>	<u>002</u>	Aug 24, 2006
<u>AB</u>		<u>400MG</u>	<u>A077242</u>	<u>003</u>	Aug 24, 2006
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>A075435</u>	<u>001</u>	Oct 08, 2004
<u>AB</u>		<u>300MG</u>	<u>A075435</u>	<u>002</u>	Oct 08, 2004
<u>AB</u>		<u>400MG</u>	<u>A075435</u>	<u>003</u>	Oct 08, 2004
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>A075485</u>	<u>003</u>	May 11, 2007
<u>AB</u>		<u>300MG</u>	<u>A075485</u>	<u>002</u>	May 11, 2007
<u>AB</u>		<u>400MG</u>	<u>A075485</u>	<u>001</u>	May 11, 2007
<u>NEURONTIN</u>					
<u>AB</u>	PFIZER PHARMS	<u>100MG</u>	<u>N020235</u>	<u>001</u>	Dec 30, 1993
<u>AB</u>		<u>300MG</u>	<u>N020235</u>	<u>002</u>	Dec 30, 1993
<u>AB</u> +		<u>400MG</u>	<u>N020235</u>	<u>003</u>	Dec 30, 1993

SOLUTION; ORAL

GABAPENTIN

<u>AA</u>	HI TECH PHARMA	<u>250MG/5ML</u>	<u>A078974</u>	<u>001</u>	Feb 18, 2011
<u>AA</u>	+ PARKE DAVIS	<u>250MG/5ML</u>	<u>N021129</u>	<u>001</u>	Mar 02, 2000

TABLET; ORAL

GABAPENTIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>600MG</u>	<u>A075694</u>	<u>001</u>	Oct 21, 2004
<u>AB</u>		<u>800MG</u>	<u>A075694</u>	<u>002</u>	Oct 21, 2004
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077894</u>	<u>001</u>	Oct 10, 2006
<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>	GLENMARK GENERICS	<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>	HIKMA PHARMS	<u>600MG</u>	<u>A078782</u>	<u>001</u>	Jul 21, 2011
<u>AB</u>		<u>800MG</u>	<u>A078782</u>	<u>002</u>	Jul 21, 2011
<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

PREScription DRUG PRODUCT LIST

3 - 198 (of 424)

GABAPENTIN

TABLET; ORAL

GABAPENTIN

<u>AB</u>	MATRIX LABS LTD	<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>	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>	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

NEURONTIN

<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

GLAXO GRP LTD 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	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

OMNITSCAN

+ GE HEALTHCARE 287MG/ML N020123 001 Jan 08, 1993
+ 28.7GM/100ML (287MG/ML) N022066 002 Sep 05, 2007

GADOFOSVESET TRISODIUM

SOLUTION: INTRAVENOUS

SECTION

ABLAVAR

LANTHEUS MEDCL 2440MG/10ML (244MG/ML) N021711 001 Dec 22, 2008
+ 3660MG/15ML (244MG/ML) N021711 002 Dec 22, 2008

GADOPENTETATE DIMEGLUMINE

INJECTABLE; INJECTION

MAGNEVIST

+ BAYER HLTHCARE 469.01MG/ML N019596 001 Jun 02, 1988
+ 469.01MG/ML N021037 001 Mar 10, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 199 (of 424)

GADOTERIDOL

INJECTABLE; INJECTION PROHANCE			
+ BRACCO	279.3MG/ML	N020131 001	Nov 16, 1992
PROHANCE MULTIPACK			
+ BRACCO	279.3MG/ML	N021489 001	Oct 09, 2003

GADOVERSETAMIDE

INJECTABLE; INJECTION OPTIMARK			
+ MALLINCKRODT	1654.5MG/5ML (330.9MG/ML)	N020937 001	Dec 08, 1999
+	3309MG/10ML (330.9MG/ML)	N020937 002	Dec 08, 1999
+	4963.5MG/15ML (330.9MG/ML)	N020937 003	Dec 08, 1999
+	6618MG/20ML (330.9MG/ML)	N020937 004	Dec 08, 1999
+	16.545GM/50ML (330.9MG/ML)	N020975 001	Dec 08, 1999
OPTIMARK IN PLASTIC CONTAINER			
+ MALLINCKRODT	3309MG/10ML (330.9MG/ML)	N020976 002	Dec 08, 1999
+	4963.5MG/15ML (330.9MG/ML)	N020976 003	Dec 08, 1999
+	6618MG/20ML (330.9MG/ML)	N020976 004	Dec 08, 1999
+	9927MG/30ML (330.9MG/ML)	N020976 001	Dec 08, 1999

GADOXETATE DISODIUM

SOLUTION; INTRAVENOUS EOVIST			
+ BAYER HLTHCARE	1.8143GM/10ML (181.43MG/ML)	N022090 001	Jul 03, 2008

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

AB BARR	<u>EQ 8MG BASE</u>	<u>A078189 001</u>	Sep 15, 2008
AB	<u>EQ 16MG BASE</u>	<u>A078189 002</u>	Sep 15, 2008
AB	<u>EQ 24MG BASE</u>	<u>A078189 003</u>	Sep 15, 2008
AB IMPAX LABS	<u>EQ 8MG BASE</u>	<u>A078484 001</u>	May 27, 2009
AB	<u>EQ 16MG BASE</u>	<u>A078484 002</u>	May 27, 2009
AB	<u>EQ 24MG BASE</u>	<u>A078484 003</u>	May 27, 2009
AB MYLAN	<u>EQ 8MG BASE</u>	<u>A090900 001</u>	Jan 24, 2011
AB	<u>EQ 16MG BASE</u>	<u>A090900 002</u>	Jan 24, 2011
AB	<u>EQ 24MG BASE</u>	<u>A090900 003</u>	Jan 24, 2011
AB SUN PHARMA GLOBAL	<u>EQ 8MG BASE</u>	<u>A090178 001</u>	Feb 02, 2011
AB	<u>EQ 16MG BASE</u>	<u>A090178 002</u>	Feb 02, 2011
AB	<u>EQ 24MG BASE</u>	<u>A090178 003</u>	Feb 02, 2011
AB WATSON LABS	<u>EQ 8MG BASE</u>	<u>A079028 001</u>	Dec 15, 2008
AB	<u>EQ 16MG BASE</u>	<u>A079028 002</u>	Dec 15, 2008
AB	<u>EQ 24MG BASE</u>	<u>A079028 003</u>	Dec 15, 2008
<u>RAZADYNE ER</u>			
AB + JANSSEN PHARMS	<u>EQ 8MG BASE</u>	<u>N021615 001</u>	Apr 01, 2005
AB	<u>EQ 16MG BASE</u>	<u>N021615 002</u>	Apr 01, 2005
AB	<u>EQ 24MG BASE</u>	<u>N021615 003</u>	Apr 01, 2005

SOLUTION; ORAL

GALANTAMINE HYDROBROMIDE

AA ROXANE	<u>4MG/ML</u>	<u>A078185 001</u>	Jan 30, 2009
AA RAZADYNE			

AA + JANSSEN PHARMS	<u>4MG/ML</u>	<u>N021224 001</u>	Jun 22, 2001
---------------------	---------------	--------------------	--------------

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

AB APOTEX INC	<u>EQ 4MG BASE</u>	<u>A077781 001</u>	Sep 27, 2011
AB	<u>EQ 8MG BASE</u>	<u>A077781 002</u>	Sep 27, 2011
AB	<u>EQ 12MG BASE</u>	<u>A077781 003</u>	Sep 27, 2011
AB AUROBINDO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A090957 001</u>	Mar 29, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 200 (of 424)

GALANTAMINE HYDROBROMIDE

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 8MG BASE</u>	<u>A090957 002</u>	Mar 29, 2011
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A090957 003</u>	Mar 29, 2011
<u>AB</u>	BARR	<u>EQ 4MG BASE</u>	<u>A077605 001</u>	Aug 28, 2008
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077605 002</u>	Aug 28, 2008
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077605 003</u>	Aug 28, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A077593 001</u>	Sep 11, 2008
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077593 002</u>	Sep 11, 2008
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077593 003</u>	Sep 11, 2008
<u>AB</u>	MYLAN	<u>EQ 4MG BASE</u>	<u>A077590 001</u>	May 29, 2009
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A077603 001</u>	Aug 28, 2008
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077590 002</u>	May 29, 2009
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077603 002</u>	Aug 28, 2008
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077590 003</u>	May 29, 2009
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077603 003</u>	Aug 28, 2008
<u>AB</u>	ROXANE	<u>EQ 4MG BASE</u>	<u>A077608 001</u>	Feb 11, 2009
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077608 002</u>	Feb 11, 2009
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077608 003</u>	Feb 11, 2009
<u>AB</u>	SANDOZ	<u>EQ 4MG BASE</u>	<u>A077589 001</u>	Jun 22, 2009
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077589 002</u>	Jun 22, 2009
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077589 003</u>	Jun 22, 2009
<u>AB</u>	TEVA PHARMS	<u>EQ 4MG BASE</u>	<u>A077587 001</u>	Jul 09, 2009
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077587 002</u>	Jul 09, 2009
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077587 003</u>	Jul 09, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 4MG BASE</u>	<u>A078898 001</u>	Feb 17, 2011
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078898 002</u>	Feb 17, 2011
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A078898 003</u>	Feb 17, 2011
<u>RAZADYNE</u>				
<u>AB</u>	+ JANSSEN PHARMS	<u>EQ 4MG BASE</u>	<u>N021169 001</u>	Feb 28, 2001
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N021169 002</u>	Feb 28, 2001
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>N021169 003</u>	Feb 28, 2001

GALLIUM CITRATE GA-67INJECTABLE; INJECTION
GALLIUM CITRATE GA 67

BS	LANTHEUS MEDCL	2mCi/ML	N017478 001
BS	MALLINCKRODT	2mCi/ML	N018058 001

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

+ GENTA	25MG/ML	N019961 002	Jan 17, 1991
---------	---------	-------------	--------------

GANCICLOVIRCAPSULE; ORAL
GANCICLOVIR

RANBAXY	250MG	A076457 001	Jun 27, 2003
+	500MG	A076457 002	Jun 27, 2003

GEL; OPHTHALMIC
ZIRGAN

+ BAUSCH AND LOMB	0.15%	N022211 001	Sep 15, 2009
-------------------	-------	-------------	--------------

IMPLANT; IMPLANTATION
VITRASERT

+ BAUSCH AND LOMB	4.5MG	N020569 001	Mar 04, 1996
-------------------	-------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 201 (of 424)

GANCICLOVIR SODIUM

INJECTABLE; INJECTION		
CYTOVENE		
<u>AP</u>	+ ROCHE PALO	<u>EQ 500MG BASE/VIAL</u>
	GANCICLOVIR	
<u>AP</u>	APP PHARMS	<u>EQ 500MG BASE/VIAL</u>

GANIRELIX ACETATE

INJECTABLE; INJECTION		
GANIRELIX ACETATE INJECTION		
+ ORGANON USA INC	EQ 250MCG BASE/0.5ML	N021057 001 Jul 29, 1999

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC		
GATIFLOXACIN		
<u>AT</u>	APOTEX CORP	<u>0.3%</u>
	ZYMAR	
<u>AT</u>	+ ALLERGAN	<u>0.3%</u>
	ZYMAXID	
+ ALLERGAN		0.5%

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION		
GEMCITABINE HYDROCHLORIDE		
<u>AP</u>	ACCORD HLTHCARE	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>
<u>AP</u>	ACTAVIS TOTOWA	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>	APP PHARMS	<u>EQ 2GM BASE/VIAL</u>
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>	FRESENIUS KABI ONCOL	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>
<u>AP</u>	HOSPIRA	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>	+ HOSPIRA INC	<u>EQ 2GM BASE/VIAL</u>
<u>AP</u>	ONCO THERAPIES LTD	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>
<u>AP</u>	SUN PHARMA GLOBAL	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>	TEVA PARENTERAL	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
<u>AP</u>	WATSON LABS	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>
	GEMZAR	
<u>AP</u>	+ LILLY	<u>EQ 200MG BASE/VIAL</u>
<u>AP</u>	+ GEMCITABINE	<u>EQ 1GM BASE/VIAL</u>
+ HOSPIRA INC		200MG/5.26ML (38MG/ML)
+		1GM/26.3ML (38MG/ML)
+		2GM/52.6ML (38MG/ML)

GEMFIBROZIL

TABLET; ORAL		
GEMFIBROZIL		
<u>AB</u>	APOTEX	<u>600MG</u>
<u>AB</u>	BLU CARIBE	<u>600MG</u>

PREScription DRUG PRODUCT LIST

3 - 202 (of 424)

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

<u>AB</u>	DAVA PHARMS INC	<u>600MG</u>	<u>A074270</u>	<u>001</u>	Sep 27, 1993
<u>AB</u>	HIKMA PHARMS	<u>600MG</u>	<u>A078599</u>	<u>001</u>	Aug 16, 2010
<u>AB</u>	IMPAKX PHARMS	<u>600MG</u>	<u>A078207</u>	<u>001</u>	Jun 01, 2007
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A077836</u>	<u>001</u>	Jul 27, 2006
<u>AB</u>	NORTHSTAR HLTHCARE	<u>600MG</u>	<u>A079072</u>	<u>001</u>	Sep 13, 2010
<u>AB</u>	SUN PHARM INDs INC	<u>600MG</u>	<u>A079239</u>	<u>001</u>	Dec 29, 2008
<u>AB</u>	TEVA	<u>600MG</u>	<u>A074256</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>	WATSON LABS	<u>600MG</u>	<u>A074442</u>	<u>001</u>	Apr 28, 1995
<u>LOPID</u>					
<u>AB</u> +	PFIZER PHARMS	<u>600MG</u>	<u>N018422</u>	<u>003</u>	Nov 20, 1986

GEMFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

± CORNERSTONE THERAPY

EQ 320MG BASE

N021158 001 Apr 04, 2003

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

AT + FOUGERA EQ 0.1% BASE A062531 001 Jul 05, 1984
AT PERRIGO NEW YORK EQ 0.1% BASE A062307 001
AT TARO EQ 0.1% BASE A062427 001 May 26, 1983

INJECTABLE; INJECTION

GENTAMICIN SULFATE

<u>AP</u>	+	APP PHARMS	<u>EQ 10MG BASE/ML</u>	<u>A062356</u>	<u>001</u>	Mar 04, 1982
<u>AP</u>	+		<u>EQ 40MG BASE/ML</u>	<u>A062356</u>	<u>002</u>	Mar 04, 1982
<u>AP</u>			<u>EQ 40MG BASE/ML</u>	<u>A062366</u>	<u>001</u>	Aug 04, 1983
<u>AP</u>		HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A062420</u>	<u>001</u>	Aug 15, 1983
<u>AP</u>			<u>EQ 10MG BASE/ML</u>	<u>A062612</u>	<u>004</u>	Feb 20, 1986
<u>AP</u>			<u>EQ 40MG BASE/ML</u>	<u>A062420</u>	<u>002</u>	Aug 15, 1983

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

GENTAMICIN SULFATE IN SOLUBLE POWDER 100MG IN PLASTIC CONTAINER	
AP +	B BRAUN
	EQ 0.8MG BASE/ML
	EQ 1.2MG BASE/ML
	EQ 1.4MG BASE/ML
	EQ 1.6MG BASE/ML
	EQ 1.8MG BASE/ML
	EQ 2MG BASE/ML
	EQ 2.4MG BASE/ML
	EQ 40MG BASE/100ML
	EQ 60MG BASE/100ML
	EQ 70MG BASE/100ML
	EQ 80MG BASE/100ML
	EQ 90MG BASE/100ML
	EQ 100MG BASE/100ML
	EQ 120MG BASE/100ML
AP	HOSPIRA
	EQ 1.2MG BASE/ML
	EQ 1.4MG BASE/ML
	EQ 1.6MG BASE/ML
	EQ 1.8MG BASE/ML
	EQ 2MG BASE/ML
	EQ 60MG BASE/100ML
	EQ 70MG BASE/100ML
	EQ 80MG BASE/100ML
	EQ 90MG BASE/100ML
	EQ 100MG BASE/100ML
ISOTONIC GENTAMICIN SULFATE IN PLASTIC CONTAINER	
AP	PAXTER ULTRACARE
	EQ 0.8MG BASE/ML
	EQ 1.2MG BASE/ML
	EQ 1.4MG BASE/ML
	EQ 1.6MG BASE/ML
	EQ 1.8MG BASE/ML
	EQ 2MG BASE/ML
	EQ 60MG BASE/100ML
	EQ 70MG BASE/100ML
	EQ 80MG BASE/100ML
	EQ 90MG BASE/100ML
	EQ 100MG BASE/100ML

PRESCRIPTION DRUG PRODUCT LIST

3 - 203 (of 424)

GENTAMICIN SULFATE

INJECTABLE; INJECTION

ISOTONIC GENTAMICIN SULFATE 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 2MG BASE/ML</u>	<u>A062373 009</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 2.4MG BASE/ML</u>	<u>A062373 010</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 40MG BASE/100ML</u>	<u>A062373 003</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 60MG BASE/100ML</u>	<u>A062373 004</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>		<u>EQ 120MG BASE/100ML</u>	<u>A062373 006</u>	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>	FERA PHARMS	<u>EQ 0.3% BASE</u>	<u>A065024 001</u>	Jul 30, 2004

OINTMENT; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>	FOUGERA	<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
-----------	----------	---------------------	--------------------	--------------

GENTAK

<u>AT</u>	AKORN	<u>EQ 0.3% BASE</u>	<u>A064163 001</u>	Oct 12, 2001
-----------	-------	---------------------	--------------------	--------------

GENTAMICIN SULFATE

<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>	FALCON PHARMS	<u>EQ 0.3% BASE</u>	<u>A062196 001</u>	
<u>AT</u>	FERA PHARMS	<u>EQ 0.3% BASE</u>	<u>A065121 001</u>	Jan 30, 2004

GENTAMICIN SULFATE; PREDNISOLONE ACETATEOINTMENT; OPHTHALMIC
PRED-G

+ ALLERGAN	EQ 0.3% BASE; 0.6%	N050612 001	Dec 01, 1989
------------	--------------------	-------------	--------------

SUSPENSION/DROPS; OPHTHALMIC

+ ALLERGAN	EQ 0.3% BASE; 1%	N050586 001	Jun 10, 1988
------------	------------------	-------------	--------------

GLATIRAMER ACETATE

INJECTABLE; SUBCUTANEOUS

COPAXONE

+ TEVA	20MG/ML	N020622 002	Feb 12, 2002
--------	---------	-------------	--------------

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>	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>	COREPHARMA	<u>1MG</u>	<u>A077274 001</u>	Oct 06, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 204 (of 424)

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

<u>AB</u>	COREPHARMA	<u>2MG</u>	<u>A077274</u> <u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A077274</u> <u>003</u>	Oct 06, 2005
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A077091</u> <u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A077091</u> <u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A077091</u> <u>003</u>	Oct 06, 2005
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A077295</u> <u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A077295</u> <u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A077295</u> <u>003</u>	Oct 06, 2005
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A077486</u> <u>001</u>	Feb 10, 2006
<u>AB</u>		<u>1MG</u>	<u>A077624</u> <u>001</u>	Nov 28, 2005
<u>AB</u>		<u>2MG</u>	<u>A077486</u> <u>002</u>	Feb 10, 2006
<u>AB</u>		<u>2MG</u>	<u>A077624</u> <u>002</u>	Nov 28, 2005
<u>AB</u>		<u>4MG</u>	<u>A077486</u> <u>003</u>	Feb 10, 2006
<u>AB</u>		<u>4MG</u>	<u>A077624</u> <u>003</u>	Nov 28, 2005
<u>AB</u>	RANBAXY	<u>1MG</u>	<u>A076875</u> <u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A076875</u> <u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A076875</u> <u>003</u>	Oct 06, 2005
<u>AB</u>		<u>8MG</u>	<u>A076875</u> <u>004</u>	Oct 06, 2005
<u>AB</u>	TEVA	<u>1MG</u>	<u>A076802</u> <u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A076802</u> <u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A076802</u> <u>003</u>	Oct 06, 2005
<u>AB</u>	VINTAGE	<u>1MG</u>	<u>A077370</u> <u>001</u>	Dec 23, 2005
<u>AB</u>		<u>2MG</u>	<u>A077370</u> <u>002</u>	Dec 23, 2005
<u>AB</u>		<u>4MG</u>	<u>A077370</u> <u>003</u>	Dec 23, 2005
<u>AB</u>		<u>8MG</u>	<u>A077370</u> <u>004</u>	Dec 23, 2005
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>A077280</u> <u>001</u>	Feb 03, 2006
<u>AB</u>		<u>2MG</u>	<u>A077280</u> <u>002</u>	Feb 03, 2006
<u>AB</u>		<u>4MG</u>	<u>A077280</u> <u>003</u>	Feb 03, 2006
<u>AB</u>	WATSON LABS FLORIDA	<u>1MG</u>	<u>A076995</u> <u>001</u>	Apr 27, 2010
<u>AB</u>		<u>2MG</u>	<u>A076995</u> <u>002</u>	Apr 27, 2010
<u>AB</u>		<u>4MG</u>	<u>A076995</u> <u>003</u>	Apr 27, 2010

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

+ TAKEDA GLOBAL

2MG; 30MG

N021925 001 Jul 28, 2006

4MG; 30MG

N021925 002 Jul 28, 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

<u>AB</u>	ACCORD HLTHCARE INC	<u>5MG</u>	<u>A074550</u> <u>001</u>	Sep 11, 1997
<u>AB</u>		<u>10MG</u>	<u>A074550</u> <u>002</u>	Sep 11, 1997
<u>AB</u>	APOTEX	<u>5MG</u>	<u>A075795</u> <u>001</u>	Jun 13, 2001
<u>AB</u>		<u>10MG</u>	<u>A075795</u> <u>002</u>	Jun 13, 2001
<u>AB</u>	CARACO	<u>5MG</u>	<u>A077820</u> <u>001</u>	Jul 11, 2006
<u>AB</u>		<u>10MG</u>	<u>A077820</u> <u>002</u>	Jul 11, 2006
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>5MG</u>	<u>A074497</u> <u>001</u>	Aug 31, 1995

PREScription DRUG PRODUCT LIST

3 - 205 (of 424)

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

<u>AB</u>	IVAX	SUB	TEVA PHARMS	<u>10MG</u>	<u>A074497</u>	<u>002</u>	Aug 31, 1995
<u>AB</u>	MYLAN			<u>5MG</u>	<u>A074226</u>	<u>001</u>	May 10, 1994
<u>AB</u>				<u>5MG</u>	<u>A074438</u>	<u>001</u>	Jun 20, 1995
<u>AB</u>				<u>10MG</u>	<u>A074226</u>	<u>002</u>	May 10, 1994
<u>AB</u>				<u>10MG</u>	<u>A074438</u>	<u>002</u>	Jun 20, 1995
<u>AB</u>	SANDOZ			<u>5MG</u>	<u>A074305</u>	<u>001</u>	Apr 07, 1995
<u>AB</u>				<u>10MG</u>	<u>A074305</u>	<u>002</u>	Apr 07, 1995
<u>AB</u>	WATSON LABS			<u>5MG</u>	<u>A074223</u>	<u>001</u>	Feb 27, 1995
<u>AB</u>				<u>5MG</u>	<u>A074370</u>	<u>001</u>	Nov 22, 1994
<u>AB</u>				<u>10MG</u>	<u>A074223</u>	<u>002</u>	Feb 27, 1995
<u>AB</u>				<u>10MG</u>	<u>A074370</u>	<u>002</u>	Nov 22, 1994
<u>GLUCOTROL</u>							
<u>AB</u>	PFIZER			<u>5MG</u>	<u>N017783</u>	<u>001</u>	May 08, 1984
<u>AB</u>	+			<u>10MG</u>	<u>N017783</u>	<u>002</u>	May 08, 1984

TABLET, EXTENDED RELEASE; OBAT-

GLIPIZIDE

<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A076467</u>	<u>003</u>	Mar 27, 2006
<u>AB</u>		<u>5MG</u>	<u>A076467</u>	<u>001</u>	Sep 08, 2003
<u>AB</u>		<u>10MG</u>	<u>A076467</u>	<u>002</u>	Nov 07, 2003
 <u>GLUCOTROL XL</u>					
<u>AB</u>	PFIZER	<u>2.5MG</u>	<u>N020329</u>	<u>003</u>	Aug 10, 1999
<u>AB</u>		<u>5MG</u>	<u>N020329</u>	<u>001</u>	Apr 26, 1994
<u>AB</u> +		<u>10MG</u>	<u>N020329</u>	<u>002</u>	Apr 26, 1994

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	CARACO	<u>2.5MG;250MG</u>	<u>A077620</u>	<u>001</u>	Jan 11, 2008
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077620</u>	<u>002</u>	Jan 11, 2008
<u>AB</u>		<u>5MG;500MG</u>	<u>A077620</u>	<u>003</u>	Jan 11, 2008
<u>AB</u>	COREPHARMA	<u>2.5MG;250MG</u>	<u>A077507</u>	<u>001</u>	Oct 27, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077507</u>	<u>002</u>	Oct 27, 2005
<u>AB</u>		<u>5MG;500MG</u>	<u>A077507</u>	<u>003</u>	Oct 27, 2005
<u>AB</u>	HERITAGE PHARMS INC	<u>2.5MG;250MG</u>	<u>A078728</u>	<u>001</u>	Jun 23, 2010
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078728</u>	<u>002</u>	Jun 23, 2010
<u>AB</u>		<u>5MG;500MG</u>	<u>A078728</u>	<u>003</u>	Jun 23, 2010
<u>AB</u>	MYLAN	<u>2.5MG;250MG</u>	<u>A078083</u>	<u>001</u>	Apr 12, 2007
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078083</u>	<u>002</u>	Apr 12, 2007
<u>AB</u>		<u>5MG;500MG</u>	<u>A078083</u>	<u>003</u>	Apr 12, 2007
<u>AB</u>	TEVA PHARMS	<u>2.5MG;250MG</u>	<u>A077270</u>	<u>001</u>	Oct 28, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077270</u>	<u>002</u>	Oct 28, 2005
<u>AB</u>	+	<u>5MG;500MG</u>	<u>A077270</u>	<u>003</u>	Oct 28, 2005
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2.5MG;250MG</u>	<u>A078905</u>	<u>001</u>	Jan 31, 2011
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078905</u>	<u>002</u>	Jan 31, 2011
<u>AB</u>		<u>5MG;500MG</u>	<u>A078905</u>	<u>003</u>	Jan 31, 2011

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

PREScription DRUG PRODUCT LIST

3 - 206 (of 424)

GLUTAMINE

FOR SOLUTION; ORAL
NUTRESTORE
+ EMMAUS MEDCL 5GM / PACKET

N021667 001 Jun 10, 2004

GLYBURIDE

TABLET; ORAL

GLYBURIDE

<u>AB</u>	AUROBINDO PHARMA	<u>1.25MG</u>	<u>A077537</u>	<u>001</u>	Oct 18, 2007
<u>AB</u>		<u>2.5MG</u>	<u>A077537</u>	<u>002</u>	Oct 18, 2007
<u>AB</u>		<u>5MG</u>	<u>A077537</u>	<u>003</u>	Oct 18, 2007
<u>AB</u>	COREPHARMA	<u>1.25MG</u>	<u>A076257</u>	<u>001</u>	Jun 27, 2002
<u>AB</u>		<u>2.5MG</u>	<u>A076257</u>	<u>002</u>	Jun 27, 2002
<u>AB</u>		<u>5MG</u>	<u>A076257</u>	<u>003</u>	Jun 27, 2002
<u>AB</u>	HERITAGE PHARMS INC	<u>1.25MG</u>	<u>A090937</u>	<u>001</u>	Feb 28, 2011
<u>AB</u>		<u>2.5MG</u>	<u>A090937</u>	<u>002</u>	Feb 28, 2011
<u>AB</u>		<u>5MG</u>	<u>A090937</u>	<u>003</u>	Feb 28, 2011
<u>AB</u>	TEVA	<u>1.25MG</u>	<u>A074388</u>	<u>001</u>	Aug 29, 1995
<u>AB</u>		<u>2.5MG</u>	<u>A074388</u>	<u>002</u>	Aug 29, 1995
<u>AB</u>	+	<u>5MG</u>	<u>A074388</u>	<u>003</u>	Aug 29, 1995
<u>GLYBURIDE (MICRONIZED)</u>					
<u>AB</u>	DAVA PHARMS INC	<u>1.5MG</u>	<u>A074591</u>	<u>001</u>	Dec 22, 1997
<u>AB</u>		<u>3MG</u>	<u>A074591</u>	<u>002</u>	Dec 22, 1997
<u>AB</u>		<u>4.5MG</u>	<u>A074591</u>	<u>003</u>	Dec 22, 1997
<u>AB</u>		<u>6MG</u>	<u>A074591</u>	<u>004</u>	Dec 22, 1997
<u>AB</u>	HIKMA	<u>1.5MG</u>	<u>A075890</u>	<u>001</u>	Jul 31, 2003
<u>AB</u>		<u>3MG</u>	<u>A075890</u>	<u>002</u>	Jul 31, 2003
<u>AB</u>		<u>6MG</u>	<u>A075890</u>	<u>003</u>	Jul 31, 2003
<u>AB</u>	MYLAN	<u>1.5MG</u>	<u>A074792</u>	<u>001</u>	Jun 26, 1998
<u>AB</u>		<u>3MG</u>	<u>A074792</u>	<u>002</u>	Jun 26, 1998
<u>AB</u>		<u>6MG</u>	<u>A074792</u>	<u>003</u>	Aug 17, 1999
<u>AB</u>	TEVA	<u>1.5MG</u>	<u>A074686</u>	<u>001</u>	Apr 20, 1999
<u>AB</u>		<u>3MG</u>	<u>A074686</u>	<u>002</u>	Apr 20, 1999
<u>AB</u>		<u>4.5MG</u>	<u>A074686</u>	<u>003</u>	Apr 20, 1999
<u>AB</u>		<u>6MG</u>	<u>A074686</u>	<u>004</u>	Apr 20, 1999
<u>GLYNASE</u>					
<u>AB</u>	PHARMACIA AND UPJOHN	<u>1.5MG</u>	<u>N020051</u>	<u>001</u>	Mar 04, 1992
<u>AB</u>		<u>3MG</u>	<u>N020051</u>	<u>002</u>	Mar 04, 1992
<u>AB</u>	+	<u>6MG</u>	<u>N020051</u>	<u>004</u>	Sep 24, 1993
<u>DIABETA</u>					
<u>BX</u>	SANOFI AVENTIS US	<u>1.25MG</u>	<u>N017532</u>	<u>001</u>	May 01, 1984
<u>BX</u>		<u>2.5MG</u>	<u>N017532</u>	<u>002</u>	May 01, 1984
<u>BX</u>	+	<u>5MG</u>	<u>N017532</u>	<u>003</u>	May 01, 1984

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>1.25MG;250MG</u>	<u>N021178</u>	<u>001</u>	Jul 31, 2000
<u>AB</u>	<u>+</u>	<u>2.5MG;500MG</u>	<u>N021178</u>	<u>002</u>	Jul 31, 2000
<u>AB</u>		<u>5MG;500MG</u>	<u>N021178</u>	<u>003</u>	Jul 31, 2000
GLYBURIDE AND METFORMIN HYDROCHLORIDE					
<u>AB</u>	ACTAVIS ELIZABETH	<u>1.25MG;250MG</u>	<u>A076716</u>	<u>001</u>	Jun 28, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A076716</u>	<u>002</u>	Jun 28, 2005
<u>AB</u>		<u>5MG;500MG</u>	<u>A076716</u>	<u>003</u>	Jun 28, 2005
<u>AB</u>	AUROBINDO PHARMA	<u>1.25MG;250MG</u>	<u>A077870</u>	<u>001</u>	Nov 14, 2007
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077870</u>	<u>002</u>	Nov 14, 2007
<u>AB</u>		<u>5MG;500MG</u>	<u>A077870</u>	<u>003</u>	Nov 14, 2007
<u>AB</u>	COREPHARMA	<u>1.25MG;250MG</u>	<u>A076731</u>	<u>001</u>	Nov 19, 2004
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A076731</u>	<u>002</u>	Nov 19, 2004
<u>AB</u>		<u>5MG;500MG</u>	<u>A076731</u>	<u>003</u>	Nov 19, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 207 (of 424)

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS INC	<u>1.25MG;250MG</u>	<u>A079009</u> <u>001</u>	Jun 03, 2009
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A079009</u> <u>002</u>	Jun 03, 2009
<u>AB</u>		<u>5MG;500MG</u>	<u>A079009</u> <u>003</u>	Jun 03, 2009
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>1.25MG;250MG</u>	<u>A076345</u> <u>001</u>	Feb 18, 2004
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A076345</u> <u>002</u>	Feb 18, 2004
<u>AB</u>		<u>5MG;500MG</u>	<u>A076345</u> <u>003</u>	Feb 18, 2004

GLYCINE

SOLUTION; IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

<u>AT</u>	BAXTER HLTHCARE	<u>1.5GM/100ML</u>	<u>N017865</u> <u>001</u>
		<u>GLYCINE 1.5% IN PLASTIC CONTAINER</u>	
<u>AT</u>	B BRAUN	<u>1.5GM/100ML</u>	<u>N016784</u> <u>001</u>

GLCOPYRROLATE

INJECTABLE; INJECTION

GLCOPYRROLATE

<u>AP</u>	HIKMA FARMACEUTICA	<u>0.2MG/ML</u>	<u>A090963</u> <u>001</u>	Sep 21, 2011
<u>AP</u>	LUITPOLD	<u>0.2MG/ML</u>	<u>A089335</u> <u>001</u>	Jul 23, 1986
		<u>ROBINUL</u>		
<u>AP</u>	+ BAXTER HLTHCARE	<u>0.2MG/ML</u>	<u>N017558</u> <u>001</u>	

SOLUTION; ORAL

CUVPOSA

+ SHIONOGI INC 1MG/5ML

N022571 001 Jul 28, 2010

TABLET; ORAL

GLCOPYRROLATE

<u>AA</u>	BOCA PHARMA	<u>1MG</u>	<u>A090020</u> <u>001</u>	Oct 19, 2011
<u>AA</u>		<u>2MG</u>	<u>A090020</u> <u>002</u>	Oct 19, 2011
<u>AA</u>	COREPHARMA	<u>1MG</u>	<u>A040568</u> <u>001</u>	Dec 22, 2004
<u>AA</u>		<u>2MG</u>	<u>A040568</u> <u>002</u>	Dec 22, 2004
<u>AA</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A040847</u> <u>001</u>	Mar 21, 2008
<u>AA</u>		<u>2MG</u>	<u>A040847</u> <u>002</u>	Mar 21, 2008
<u>AA</u>	PAR PHARM	<u>1MG</u>	<u>A040653</u> <u>001</u>	Aug 31, 2006
<u>AA</u>		<u>2MG</u>	<u>A040653</u> <u>002</u>	Aug 31, 2006
<u>AA</u>	RANBAXY	<u>1MG</u>	<u>A040844</u> <u>001</u>	Aug 18, 2009
<u>AA</u>		<u>2MG</u>	<u>A040844</u> <u>002</u>	Aug 18, 2009
<u>AA</u>	VINTAGE	<u>1MG</u>	<u>A040821</u> <u>001</u>	Dec 29, 2008
<u>AA</u>		<u>2MG</u>	<u>A040821</u> <u>002</u>	Dec 29, 2008
<u>AA</u>	WEST WARD	<u>1MG</u>	<u>A040836</u> <u>001</u>	Mar 05, 2009
<u>AA</u>		<u>2MG</u>	<u>A040836</u> <u>002</u>	Mar 05, 2009
		<u>ROBINUL</u>		
<u>AA</u>	+ SHIONOGI INC	<u>1MG</u>	<u>N012827</u> <u>001</u>	
		<u>ROBINUL FORTE</u>		
<u>AA</u>	+ SHIONOGI INC	<u>2MG</u>	<u>N012827</u> <u>002</u>	

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

CHORIONIC GONADOTROPIN

<u>AP</u>	+ APP PHARMS	<u>10,000 UNITS/VIAL</u>	<u>N017067</u> <u>002</u>
<u>AP</u>	+ FERRING	<u>10,000 UNITS/VIAL</u>	<u>N017016</u> <u>007</u>
		<u>PREGNYL</u>	
<u>AP</u>	+ ORGANON USA INC	<u>10,000 UNITS/VIAL</u>	<u>N017692</u> <u>001</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 208 (of 424)

GOSERELIN ACETATE

IMPLANT; IMPLANTATION

ZOLADEX

+ ASTRazeneca	EQ 3.6MG BASE	N019726 001	Dec 29, 1989
+	EQ 10.8MG BASE	N020578 001	Jan 11, 1996

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

AT + BAUSCH AND LOMB	<u>0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML</u>	<u>A064047 001</u>	Jan 31, 1996
AT LUITPOLD	<u>0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML</u>	<u>A065187 001</u>	Oct 28, 2005
	<u>NEOSPORIN</u>		
AT + 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

+ PROSTRAKAN INC	3.1MG/24HR	N022198 001	Sep 12, 2008
------------------	------------	-------------	--------------

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AP AKORN INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A079119 001</u>	Sep 10, 2009
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A079078 002</u>	Sep 14, 2009
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A079078 001</u>	Sep 14, 2009
AP APP PHARMS	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078522 001</u>	Dec 31, 2007
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078090 001</u>	Jun 30, 2008
AP BAXTER HLTHCARE	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077177 001</u>	Dec 31, 2007
AP BEDFORD LABS	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A077913 001</u>	Jun 26, 2008
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077186 001</u>	Jun 30, 2008
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077187 001</u>	Jun 30, 2008
AP CLARIS LIFESCIENCES	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078197 001</u>	Dec 31, 2007
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078198 001</u>	Jun 30, 2008
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078198 002</u>	Jun 30, 2008
AP DR REDDYS LABS INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078863 001</u>	Jun 30, 2008
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078880 001</u>	Jun 30, 2008
AP EBEWE PHARMA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078808 001</u>	Apr 29, 2008
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078835 001</u>	Jun 30, 2008
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078835 002</u>	Jun 30, 2008
AP LUITPOLD	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A091274 001</u>	Sep 22, 2010
AP SAGENT STRIDES	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A091136 001</u>	Apr 09, 2010
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A091136 002</u>	Apr 09, 2010
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A091137 002</u>	Apr 09, 2010
AP SANDOZ	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078534 001</u>	Apr 30, 2009
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078531 001</u>	Apr 30, 2009
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078531 002</u>	Apr 30, 2009
AP + TEVA PARENTERAL	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078392 001</u>	Dec 31, 2007
AP +	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077963 001</u>	Jan 03, 2008
AP +	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077297 001</u>	Jun 30, 2008
AP WATSON LABS	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078262 001</u>	Dec 31, 2007
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078258 001</u>	Jun 30, 2008
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078258 002</u>	Jun 30, 2008
AP WOCKHARDT USA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078566 001</u>	Feb 29, 2008
AP	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078564 001</u>	Jun 30, 2008
AP	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078565 001</u>	Jun 30, 2008
	<u>GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE</u>		
AP APP PHARMS	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078096 001</u>	Jun 30, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 209 (of 424)

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	DR REDDYS LABS INC	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078863 002</u>	Jun 30, 2008
<u>AP</u>	+ TEVA PARENTERAL	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077165 001</u>	Dec 31, 2007
<u>GRANISTERON HYDROCHLORIDE</u>				
<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

SOLUTION; ORAL

GRANISETRON HYDROCHLORIDE

+ PEDIATRX EQ 2MG BASE/10ML

A078334 001 Feb 28, 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>	COREPHARMA	<u>EQ 1MG BASE</u>	<u>A078260 001</u>	Dec 31, 2007
<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>	ROXANE	<u>EQ 1MG BASE</u>	<u>A077842 001</u>	Dec 31, 2007
<u>AB</u>	TARO	<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

GRISEOFULVIN

SUSPENSION; ORAL

GRISEOFULVIN

<u>AB</u>	VINTAGE	<u>125MG/5ML</u>	<u>A065438 001</u>	Oct 08, 2010
-----------	---------	------------------	--------------------	--------------

GRISEOFULVIN, MICROCRYSTALLINE

SUSPENSION; ORAL

GRIFULVIN V

<u>AB</u>	+ ORTHONEUTROGENA	<u>125MG/5ML</u>	<u>A062483 001</u>	Jan 26, 1984
<u>GRISEOFULVIN</u>				
<u>AB</u>	ACTAVIS MID ATLANTIC	<u>125MG/5ML</u>	<u>A065394 001</u>	Jul 06, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>125MG/5ML</u>	<u>A065354 001</u>	Sep 10, 2007
<u>AB</u>	PERRIGO CO TENNESSEE	<u>125MG/5ML</u>	<u>A065200 001</u>	Mar 02, 2005

TABLET; ORAL

GRIFULVIN V

+ ORTHONEUTROGENA 500MG

A062279 003

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL

GRIS-PEG

PEDINOL 125MG

N050475 001

+ 250MG

N050475 002

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

IVAX SUB TEVA PHARMS EQ 4MG BASE

A074149 001 Apr 07, 1995

+ EQ 8MG BASE

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	<u>EQ 1MG BASE</u>	<u>A074673 001</u>	Feb 28, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 210 (of 424)

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA	<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
	<u>TENEX</u>			
<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

INTUNIV

SHIRE	EQ 1MG BASE	N022037 001	Sep 02, 2009
	EQ 2MG BASE	N022037 002	Sep 02, 2009
	EQ 3MG BASE	N022037 003	Sep 02, 2009
+	EQ 4MG BASE	N022037 004	Sep 02, 2009

GUANIDINE HYDROCHLORIDE

TABLET; ORAL

GUANIDINE HYDROCHLORIDE

SCHERING	125MG	N001546 001
----------	-------	-------------

HALCINONIDE

CREAM; TOPICAL

HALOG

+	RANBAXY	0.1%	N017556 001
---	---------	------	-------------

OINTMENT; TOPICAL

HALOG

+	RANBAXY	0.1%	N017824 001
---	---------	------	-------------

HALOBETASOL PROPIONATE

CREAM; TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A077001 001</u>	Dec 16, 2004
<u>AB</u>	G AND W LABS	<u>0.05%</u>	<u>A078162 001</u>	Apr 24, 2007
<u>AB</u>	PERRIGO ISRAEL	<u>0.05%</u>	<u>A077123 001</u>	Dec 16, 2004
<u>AB</u>	TARO	<u>0.05%</u>	<u>A077227 001</u>	Aug 04, 2005
	<u>ULTRAVATE</u>			
<u>AB</u> +	RANBAXY	<u>0.05%</u>	<u>N019967 001</u>	Dec 27, 1990

OINTMENT; TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>A076903 001</u>	Dec 16, 2004
<u>AB</u>	G AND W LABS	<u>0.05%</u>	<u>A077721 001</u>	Sep 07, 2006
<u>AB</u>	PERRIGO	<u>0.05%</u>	<u>A076872 001</u>	Dec 16, 2004
<u>AB</u>	TARO	<u>0.05%</u>	<u>A076994 001</u>	Dec 16, 2004
	<u>ULTRAVATE</u>			
<u>AB</u> +	RANBAXY	<u>0.05%</u>	<u>N019968 001</u>	Dec 17, 1990

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

<u>AB</u>	MYLAN	<u>0.5MG</u>	<u>A070278 006</u>	Jun 10, 1986
<u>AB</u>		<u>1MG</u>	<u>A070278 004</u>	Jun 10, 1986
<u>AB</u>		<u>2MG</u>	<u>A070278 001</u>	Jun 10, 1986
<u>AB</u>		<u>5MG</u>	<u>A070278 005</u>	Jun 10, 1986
<u>AB</u>		<u>10MG</u>	<u>A070278 002</u>	Jul 16, 2009
<u>AB</u>		<u>20MG</u>	<u>A070278 003</u>	Jul 16, 2009
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>A071206 001</u>	Nov 17, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 211 (of 424)

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

<u>AB</u>	SANDOZ	<u>1MG</u>	<u>A071207</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>	+	<u>2MG</u>	<u>A071208</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>		<u>5MG</u>	<u>A071209</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>		<u>10MG</u>	<u>A071210</u>	<u>001</u>	Mar 11, 1988
<u>AB</u>		<u>20MG</u>	<u>A071211</u>	<u>001</u>	Mar 11, 1988
<u>AB</u>	ZYDUS PHARMS USA	<u>5MG</u>	<u>A077580</u>	<u>003</u>	Nov 29, 2007
<u>AB</u>		<u>10MG</u>	<u>A077580</u>	<u>004</u>	Nov 29, 2007
<u>AB</u>		<u>20MG</u>	<u>A077580</u>	<u>005</u>	Nov 29, 2007

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL

<u>AO</u>	+	JANSEN PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N018701</u>	<u>001</u>	Jan 14, 1986
<u>AO</u>	+		<u>EQ 100MG BASE/ML</u>	<u>N018701</u>	<u>002</u>	Jan 31, 1997
<u>AO</u>		<u>HALOPERIDOL DECANOATE</u>				
<u>AO</u>		APP PHARMS	<u>EQ 50MG BASE/ML</u>	<u>A074893</u>	<u>001</u>	Dec 19, 1997
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A074893</u>	<u>002</u>	Dec 19, 1997
<u>AO</u>		BEDFORD	<u>EQ 50MG BASE/ML</u>	<u>A074811</u>	<u>001</u>	Jan 30, 1998
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A075305</u>	<u>001</u>	Sep 28, 1998
<u>AO</u>		CLARIS LIFESCIENCES	<u>EQ 50MG BASE/ML</u>	<u>A075440</u>	<u>001</u>	Feb 28, 2000
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A075440</u>	<u>002</u>	Feb 28, 2000
<u>AO</u>		TEVA PARENTERAL	<u>EQ 50MG BASE/ML</u>	<u>A075393</u>	<u>001</u>	May 11, 1999
<u>AO</u>			<u>EQ 100MG BASE/ML</u>	<u>A075393</u>	<u>002</u>	May 11, 1999

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALOPERIDOL

<u>AA</u>	PHARM ASSOC	<u>EQ 2MG BASE/ML</u>	<u>A073037</u>	<u>001</u>	Feb 26, 1993	
<u>AA</u>	SILARX	<u>EQ 2MG BASE/ML</u>	<u>A073364</u>	<u>001</u>	Sep 28, 1993	
<u>AA</u>	+	TEVA PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A071617</u>	<u>001</u>	Dec 01, 1988

INJECTABLE; INJECTION

HALDOL

<u>AP</u>	+	JANSEN PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N015923</u>	<u>001</u>
<u>AP</u>		<u>HALOPERIDOL</u>			
<u>AP</u>		APP PHARMS	<u>EQ 5MG BASE/ML</u>	<u>A075689</u>	<u>001</u>
<u>AP</u>		BEDFORD	<u>EQ 5MG BASE/ML</u>	<u>A075858</u>	<u>001</u>
<u>AP</u>		CLARIS LIFESCIENCES	<u>EQ 5MG BASE/ML</u>	<u>A076791</u>	<u>001</u>
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A076828</u>	<u>001</u>
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A076774</u>	<u>001</u>
<u>AP</u>		PFIZER	<u>EQ 5MG BASE/ML</u>	<u>A078347</u>	<u>001</u>
<u>AP</u>		SAGENT PHARMS	<u>EQ 5MG BASE/ML</u>	<u>A091637</u>	<u>001</u>
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A200742</u>	<u>001</u>
<u>AP</u>		TEVA PARENTERAL	<u>EQ 5MG BASE/ML</u>	<u>A076035</u>	<u>001</u>

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>	+	APP PHARMS	<u>1,000 UNITS/ML</u>	<u>N017029</u>	<u>001</u>
<u>AP</u>	+		<u>5,000 UNITS/ML</u>	<u>N017651</u>	<u>006</u>
<u>AP</u>	+		<u>10,000 UNITS/ML</u>	<u>N017029</u>	<u>003</u>
<u>AP</u>	+		<u>20,000 UNITS/ML</u>	<u>N017029</u>	<u>004</u>
<u>AP</u>	+	BAXTER HLTHCARE	<u>1,000 UNITS/ML</u>	<u>N017037</u>	<u>001</u>
<u>AP</u>	+		<u>5,000 UNITS/ML</u>	<u>N017037</u>	<u>002</u>
<u>AP</u>	+		<u>10,000 UNITS/ML</u>	<u>N017037</u>	<u>003</u>
<u>AP</u>		HOSPIRA	<u>5,000 UNITS/ML</u>	<u>A088100</u>	<u>001</u>
<u>AP</u>		HOSPIRA INC	<u>1,000 UNITS/ML</u>	<u>A090571</u>	<u>001</u>
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A090571</u>	<u>002</u>

Apr 28, 1983
Aug 31, 2009
Aug 31, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 212 (of 424)

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>	HOSPIRA INC	<u>10,000 UNITS/ML</u>	<u>A090571</u> <u>003</u>	Aug 31, 2009
<u>AP</u>	SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090808</u> <u>001</u>	Jun 30, 2010
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A090808</u> <u>002</u>	Jun 30, 2010
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A090808</u> <u>003</u>	Jun 30, 2010
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>A090809</u> <u>001</u>	Jun 30, 2010
<u>AP</u>	SANDOZ	<u>1,000 UNITS/ML</u>	<u>A091682</u> <u>001</u>	Jun 08, 2011
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A091659</u> <u>001</u>	Jun 08, 2011
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>A091682</u> <u>002</u>	Jun 08, 2011
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>A201002</u> <u>001</u>	Jun 08, 2011
	<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</u> <u>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</u> <u>001</u>	Jul 20, 1992
<u>AP</u>	HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916</u> <u>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</u> <u>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</u> <u>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</u> <u>011</u>	Jun 23, 1989
	<u>HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	BAXTER HLTHCARE	<u>4,000 UNITS/100ML</u>	<u>N018814</u> <u>001</u>	Oct 31, 1983
	<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</u> <u>001</u>	Jul 20, 1992
<u>AP</u>	HOSPIRA	<u>4,000 UNITS/100ML</u>	<u>N019805</u> <u>001</u>	Jan 25, 1989
	<u>HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	BAXTER HLTHCARE	<u>5,000 UNITS/100ML</u>	<u>N018814</u> <u>003</u>	Jul 09, 1985
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N018814</u> <u>004</u>	Jul 02, 1987
	<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</u> <u>004</u>	Jul 20, 1992
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N019952</u> <u>005</u>	Jul 20, 1992
<u>AP</u>	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N019339</u> <u>004</u>	Mar 27, 1985
<u>AP</u>		<u>5,000 UNITS/100ML</u>	<u>N019805</u> <u>002</u>	Jan 25, 1989
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N019339</u> <u>002</u>	Mar 27, 1985
	<u>HEPARIN SODIUM IN PLASTIC CONTAINER</u>			
<u>AP</u>	+ APP PHARMS	<u>1,000 UNITS/ML</u>	<u>N017029</u> <u>013</u>	Dec 05, 1985
<u>AP</u>	+	<u>5,000 UNITS/ML</u>	<u>N017029</u> <u>014</u>	Dec 05, 1985
<u>AP</u>	+	<u>10,000 UNITS/ML</u>	<u>N017029</u> <u>015</u>	Dec 05, 1985
<u>AP</u>	+	<u>20,000 UNITS/ML</u>	<u>N017029</u> <u>016</u>	Dec 05, 1985
	<u>HEPARIN SODIUM PRESERVATIVE FREE</u>			
<u>AP</u>	+ APP PHARMS	<u>1,000 UNITS/ML</u>	<u>N017029</u> <u>010</u>	Apr 28, 1986
<u>AP</u>	+ HOSPIRA	<u>10,000 UNITS/ML</u>	<u>A089522</u> <u>001</u>	May 04, 1987
<u>AP</u>	SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090810</u> <u>001</u>	Jun 30, 2010
	<u>HEPARIN SODIUM</u>			
	PFIZER	<u>1,000 UNITS/ML</u>	<u>N201370</u> <u>001</u>	Jul 21, 2011
		<u>5,000 UNITS/ML</u>	<u>N201370</u> <u>002</u>	Jul 21, 2011
		<u>10,000 UNITS/ML</u>	<u>N201370</u> <u>003</u>	Jul 21, 2011
	<u>HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N019339</u> <u>001</u>	Mar 27, 1985
	<u>HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N018916</u> <u>006</u>	Jan 31, 1984
	<u>HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N018916</u> <u>007</u>	Jan 31, 1984
		<u>10,000 UNITS/100ML</u>	<u>N018916</u> <u>008</u>	Jan 31, 1984
	<u>HEPARIN SODIUM PRESERVATIVE FREE</u>			
	PFIZER	<u>1,000 UNITS/ML</u>	<u>N201370</u> <u>004</u>	Jul 21, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 213 (of 424)

HEXACHLOROPHENONE

EMULSION; TOPICAL PHISOHEX			
+ SANOFI AVENTIS US	3%	N006882	001
SPONGE; TOPICAL <u>PRE-OP</u>			
<u>AT</u> + DAVIS AND GECK	<u>480MG</u>	<u>N017433</u>	<u>001</u>
<u>PRE-OP II</u>			
<u>AT</u> DAVIS AND GECK	<u>480MG</u>	<u>N017433</u>	<u>002</u>

HEXAMINOLEVULINATE HYDROCHLORIDE

FOR SOLUTION; INTRAVESICAL CYSVIEW KIT			
+ GE HEALTHCARE	100MG/VIAL	N022555	001 May 28, 2010

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS SUPPRELIN LA			
+ ENDO PHARM	50MG	N022058	001 May 03, 2007
VANTAS			
+ ENDO PHARM	50MG	N021732	001 Oct 12, 2004

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL <u>HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE</u>			
<u>AA</u> ACTAVIS MID ATLANTIC	<u>1.5MG/5ML;5MG/5ML</u>	<u>A088017</u>	<u>001 Jul 05, 1983</u>
<u>AA</u> + HI TECH PHARMA	<u>1.5MG/5ML;5MG/5ML</u>	<u>A040613</u>	<u>001 Feb 08, 2008</u>
<u>AA</u> WOCKHARDT	<u>1.5MG/5ML;5MG/5ML</u>	<u>A088008</u>	<u>001 Mar 03, 1983</u>
TABLET; ORAL <u>HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE</u>			
<u>AB</u> NOVEL LABS INC	<u>1.5MG;5MG</u>	<u>A091528</u>	<u>001 Apr 20, 2011</u>
<u>TUSSIGON</u>			
<u>AB</u> + KING PHARMS	<u>1.5MG;5MG</u>	<u>A088508</u>	<u>001 Jul 30, 1985</u>

HYALURONIDASE

INJECTABLE; INJECTION AMPHADASE			
+ AMPHASTAR PHARM	150 UNITS/ML	N021665	001 Oct 26, 2004
HYDASE			
+ AKORN INC	150 UNITS/ML	N021716	001 Oct 25, 2005
VITRASE			
+ ISTA PHARMS	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

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION <u>HYDRALAZINE HYDROCHLORIDE</u>			
<u>AP</u> AKORN	<u>20MG/ML</u>	<u>A040730</u>	<u>001 Apr 21, 2009</u>
<u>AP</u> APP PHARMS	<u>20MG/ML</u>	<u>A040388</u>	<u>001 Mar 13, 2001</u>
<u>AP</u> + LUITPOLD	<u>20MG/ML</u>	<u>A040136</u>	<u>001 Jun 30, 1997</u>
TABLET; ORAL <u>HYDRALAZINE HYDROCHLORIDE</u>			
<u>AA</u> GLENMARK PHARMS LTD	<u>10MG</u>	<u>A090527</u>	<u>001 May 27, 2009</u>
	<u>25MG</u>	<u>A090527</u>	<u>002 May 27, 2009</u>
	<u>50MG</u>	<u>A090527</u>	<u>003 May 27, 2009</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 214 (of 424)

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

<u>AA</u>	GLENMARK PHARMS LTD	<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 UNIT 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>	MYLAN	<u>10MG</u>	<u>A090413</u>	<u>001</u>	Dec 08, 2010
<u>AA</u>		<u>25MG</u>	<u>A090413</u>	<u>002</u>	Dec 08, 2010
<u>AA</u>		<u>50MG</u>	<u>A090413</u>	<u>003</u>	Dec 08, 2010
<u>AA</u>		<u>100MG</u>	<u>A090413</u>	<u>004</u>	Dec 08, 2010
<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>	WATSON LABS	<u>25MG</u>	<u>A084504</u>	<u>001</u>	
<u>AA</u>		<u>50MG</u>	<u>A084503</u>	<u>001</u>	
<u>AA</u>	ZYDUS PHARMS USA	<u>10MG</u>	<u>A040858</u>	<u>001</u>	Feb 26, 2010
<u>AA</u>		<u>25MG</u>	<u>A040858</u>	<u>002</u>	Feb 26, 2010
<u>AA</u>		<u>50MG</u>	<u>A040858</u>	<u>003</u>	Feb 26, 2010
<u>AA</u>		<u>100MG</u>	<u>A040858</u>	<u>004</u>	Feb 26, 2010

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

PAR PHARM	25MG; 25MG	
+	50MG; 50MG	

A088957 001 Oct 21, 1985
A088946 001 Oct 21, 1985

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

+ ARBOR PHARMS INC	37.5MG; 20MG	
--------------------	--------------	--

N020727 001 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>	UNICHEM	<u>12.5MG</u>	<u>A090510</u>	<u>001</u>	Jan 19, 2010
<u>AB</u>	VINTAGE PHARMS	<u>12.5MG</u>	<u>A075907</u>	<u>001</u>	Sep 17, 2002
<u>AB</u>	WEST WARD	<u>12.5MG</u>	<u>A077885</u>	<u>001</u>	Nov 26, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 215 (of 424)

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

MICROZIDE

<u>AB</u>	+ WATSON LABS	<u>12.5MG</u>	<u>N020504</u>	<u>001</u>	Dec 27, 1996
TABLET; ORAL					
<u>HYDROCHLOROTHIAZIDE</u>					
<u>AB</u>	ACTAVIS ELIZABETH	<u>12.5MG</u>	<u>A040707</u>	<u>001</u>	Feb 27, 2007
<u>AB</u>	APOTEX	<u>25MG</u>	<u>A040774</u>	<u>001</u>	Oct 03, 2007
<u>AB</u>		<u>50MG</u>	<u>A040774</u>	<u>002</u>	Oct 03, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A040780</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040780</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>	CARACO	<u>12.5MG</u>	<u>A040857</u>	<u>001</u>	May 30, 2008
<u>AB</u>		<u>25MG</u>	<u>A040810</u>	<u>001</u>	Mar 27, 2007
<u>AB</u>		<u>50MG</u>	<u>A040810</u>	<u>002</u>	Mar 27, 2007
<u>AB</u>	DAVA PHARMS INC	<u>25MG</u>	<u>A087059</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A087068</u>	<u>001</u>	
<u>AB</u>	EXCELLIUM	<u>25MG</u>	<u>A040702</u>	<u>001</u>	Mar 16, 2007
<u>AB</u>		<u>50MG</u>	<u>A040702</u>	<u>002</u>	Mar 16, 2007
<u>AB</u>	HERITAGE PHARMS INC	<u>25MG</u>	<u>A085182</u>	<u>002</u>	
<u>AB</u>		<u>50MG</u>	<u>A085182</u>	<u>001</u>	
<u>AB</u>	IPCA LABS LTD	<u>12.5MG</u>	<u>A040807</u>	<u>001</u>	Jul 20, 2007
<u>AB</u>		<u>25MG</u>	<u>A040807</u>	<u>002</u>	Jul 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040807</u>	<u>003</u>	Jul 20, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A083177</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A083177</u>	<u>002</u>	
<u>AB</u>	+ JUBILANT CADISTA	<u>25MG</u>	<u>A040809</u>	<u>001</u>	Sep 04, 2007
<u>AB</u>		<u>50MG</u>	<u>A040809</u>	<u>002</u>	Sep 04, 2007
<u>AB</u>	LANNETT	<u>25MG</u>	<u>A084325</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A084324</u>	<u>001</u>	
<u>AB</u>	MYLAN	<u>12.5MG</u>	<u>A040770</u>	<u>001</u>	Jan 23, 2007
<u>AB</u>	MYLAN PHARMS INC	<u>25MG</u>	<u>A040735</u>	<u>002</u>	Jan 23, 2007
<u>AB</u>		<u>50MG</u>	<u>A040735</u>	<u>003</u>	Jan 23, 2007
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A040907</u>	<u>001</u>	Aug 15, 2008
<u>AB</u>		<u>50MG</u>	<u>A040907</u>	<u>002</u>	Aug 15, 2008
<u>AB</u>	VINTAGE PHARMS	<u>25MG</u>	<u>A040412</u>	<u>001</u>	Mar 29, 2002
<u>AB</u>		<u>50MG</u>	<u>A040412</u>	<u>002</u>	Mar 29, 2002
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A081189</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>	WEST WARD	<u>25MG</u>	<u>A084878</u>	<u>002</u>	Jul 12, 2006
<u>AB</u>		<u>50MG</u>	<u>A084878</u>	<u>001</u>	
<u>ORETIC</u>					
<u>AB</u>	ABBOTT	<u>50MG</u>	<u>N011971</u>	<u>002</u>	

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

+ SANOFI AVENTIS	12.5MG;150MG	<u>N020758</u>	<u>002</u>	Sep 30, 1997
	12.5MG;300MG	<u>N020758</u>	<u>003</u>	Aug 31, 1998

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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 216 (of 424)

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET; ORAL

LISINOPRIL AND HYDROCHLOROTHIAZIDE

<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>	RANBAXY	<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>	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>	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
<u>AB</u>	WEST WARD	<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>PRINZIDE</u>				
<u>AB</u>	MERCK	<u>12.5MG;10MG</u>	<u>N019778</u>	<u>003</u>	Nov 18, 1993
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N019778</u>	<u>001</u>	Feb 16, 1989
	<u>ZESTORETIC</u>				
<u>AB</u>	ASTRAZENECA	<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	<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>	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>	LUPIN LTD	<u>12.5MG;50MG</u>	<u>A078245</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A078245</u>	<u>002</u>	May 21, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078245</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	MYLAN	<u>12.5MG;50MG</u>	<u>A091652</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091652</u>	<u>002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A091652</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	ROXANE	<u>12.5MG;50MG</u>	<u>A077732</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077732</u>	<u>001</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077732</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	SANDOZ	<u>12.5MG;50MG</u>	<u>A077948</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077948</u>	<u>003</u>	Aug 19, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077948</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>	TEVA PHARMS	<u>12.5MG;50MG</u>	<u>A077157</u>	<u>001</u>	Apr 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077157</u>	<u>002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077157</u>	<u>003</u>	Apr 06, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 217 (of 424)

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	TORRENT PHARMS	<u>12.5MG;50MG</u>	<u>A090528</u> <u>001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A090528</u> <u>003</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A090528</u> <u>002</u>	Oct 06, 2010
<u>AB</u>	WATSON LABS	<u>12.5MG;50MG</u>	<u>A200180</u> <u>001</u>	Jan 12, 2011
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A200180</u> <u>002</u>	Jan 12, 2011
<u>AB</u>		<u>25MG;100MG</u>	<u>A200180</u> <u>003</u>	Jan 12, 2011
<u>AB</u>	ZYDUS PHARMS USA INC	<u>12.5MG;50MG</u>	<u>A078385</u> <u>001</u>	Oct 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078385</u> <u>002</u>	Oct 06, 2010

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

MYLAN	15MG;250MG	A070264	001	Jan 23, 1986
+	25MG;250MG	A070265	001	Jan 23, 1986

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

<u>AB</u>	NOVARTIS	<u>25MG;50MG</u>	<u>N018303</u> <u>001</u>	Dec 31, 1984
<u>AB</u>	+	<u>25MG;100MG</u>	<u>N018303</u> <u>002</u>	Dec 31, 1984
		<u>METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE</u>		
<u>AB</u>	MYLAN	<u>25MG;50MG</u>	<u>A076792</u> <u>001</u>	Aug 20, 2004
<u>AB</u>		<u>25MG;100MG</u>	<u>A076792</u> <u>002</u>	Aug 20, 2004
		<u>METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE</u>		
	MYLAN	<u>50MG;100MG</u>	<u>A076792</u> <u>003</u>	Aug 20, 2004

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	GLENMARK PHARMS	<u>12.5MG;7.5MG</u>	<u>A090718</u> <u>001</u>	Mar 17, 2010
<u>AB</u>		<u>12.5MG;15MG</u>	<u>A090718</u> <u>002</u>	Mar 17, 2010
<u>AB</u>		<u>25MG;15MG</u>	<u>A090718</u> <u>003</u>	Mar 17, 2010
<u>AB</u>	PADDOCK LLC	<u>12.5MG;7.5MG</u>	<u>A090096</u> <u>001</u>	Sep 25, 2008
<u>AB</u>		<u>12.5MG;15MG</u>	<u>A090096</u> <u>002</u>	Sep 25, 2008
<u>AB</u>		<u>25MG;15MG</u>	<u>A090096</u> <u>003</u>	Sep 25, 2008
<u>AB</u>	TEVA	<u>12.5MG;7.5MG</u>	<u>A076980</u> <u>001</u>	Mar 07, 2007
<u>AB</u>		<u>12.5MG;15MG</u>	<u>A076980</u> <u>003</u>	Mar 07, 2007
<u>AB</u>		<u>25MG;15MG</u>	<u>A076980</u> <u>002</u>	Mar 07, 2007
		<u>UNIRETIC</u>		
<u>AB</u>	UCB INC	<u>12.5MG;7.5MG</u>	<u>N020729</u> <u>001</u>	Jun 27, 1997
<u>AB</u>		<u>12.5MG;15MG</u>	<u>N020729</u> <u>003</u>	Feb 14, 2002
<u>AB</u>	+	<u>25MG;15MG</u>	<u>N020729</u> <u>002</u>	Jun 27, 1997

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

	DAIICHI SANKYO	12.5MG;20MG	N021532	002	Jun 05, 2003
		12.5MG;40MG	N021532	003	Jun 05, 2003
+		25MG;40MG	N021532	005	Jun 05, 2003

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

INDERIDE-40/25

<u>AB</u>	+ AKRIMAX PHARMS	<u>25MG;40MG</u>	<u>N018031</u> <u>001</u>	
		<u>PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE</u>		
<u>AB</u>	MYLAN	<u>25MG;40MG</u>	<u>A070946</u> <u>001</u>	Mar 04, 1987
<u>AB</u>	+	<u>25MG;80MG</u>	<u>A070947</u> <u>001</u>	Apr 01, 1987

PREScription DRUG PRODUCT LIST

3 - 218 (of 424)

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB	PLIVA	25MG;40MG	A072042	001	Mar 14, 1988
AB		25MG;80MG	A072043	001	Mar 14, 1988
AB	WATSON LABS	25MG;40MG	A070301	001	Apr 18, 1986
AB		25MG;80MG	A070305	001	Apr 18, 1986

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

<u>AB</u>	PFIZER PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>N020125</u>	<u>001</u>	Dec 28, 1999
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>N020125</u>	<u>002</u>	Dec 28, 1999
<u>AB</u>	<u>+</u>	<u>25MG;EQ 20MG BASE</u>	<u>N020125</u>	<u>003</u>	Dec 28, 1999
		<u>QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE</u>			
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG;EQ 10MG BASE</u>	<u>A078450</u>	<u>001</u>	Aug 24, 2007
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>A078450</u>	<u>002</u>	Aug 24, 2007
<u>AB</u>		<u>25MG;EQ 20MG BASE</u>	<u>A078450</u>	<u>003</u>	Aug 24, 2007
<u>AB</u>	INVAGEN PHARMS	<u>12.5MG;10MG</u>	<u>A201356</u>	<u>001</u>	Apr 20, 2011
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A201356</u>	<u>002</u>	Apr 20, 2011
<u>AB</u>		<u>25MG;20MG</u>	<u>A201356</u>	<u>003</u>	Apr 20, 2011
<u>AB</u>	MYLAN	<u>12.5MG;EQ 10MG BASE</u>	<u>A077093</u>	<u>001</u>	Mar 28, 2005
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>A077093</u>	<u>002</u>	Mar 28, 2005
<u>AB</u>		<u>25MG;EQ 20MG BASE</u>	<u>A077093</u>	<u>003</u>	Mar 28, 2005
<u>AB</u>	RANBAXY	<u>12.5MG;EQ 10MG BASE</u>	<u>A078211</u>	<u>001</u>	Mar 04, 2009
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>A078211</u>	<u>002</u>	Mar 04, 2009
<u>AB</u>		<u>25MG;EQ 20MG BASE</u>	<u>A078211</u>	<u>003</u>	Mar 04, 2009
		<u>QUINARETIC</u>			
<u>AB</u>	GAVIS PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>A076374</u>	<u>001</u>	Mar 31, 2004
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>A076374</u>	<u>002</u>	Mar 31, 2004
<u>AB</u>		<u>25MG;EO 20MG BASE</u>	<u>A076374</u>	<u>003</u>	Mar 31, 2004

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

ALDACTAZIDE

<u>AB</u>	GD SEARLE LLC	<u>25MG; 25MG</u>	<u>N012616</u>	<u>004</u>	Dec 30, 1982
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE					
<u>AB</u>	MUTUAL PHARM	<u>25MG; 25MG</u>	<u>A089534</u>	<u>001</u>	Jul 02, 1987
<u>AB</u>	MYLAN	<u>25MG; 25MG</u>	<u>A086513</u>	<u>001</u>	
<u>AB</u>	WATSON LABS	<u>25MG; 25MG</u>	<u>A087398</u>	<u>001</u>	
ALDACTAZIDE					
+ GD SEARLE LLC	50MG; 50MG	N012616	005	Dec 30, 1982	

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

MICARDIS HCT

BOEHRINGER INGELHEIM 12.5MG;40MG N021162 001 Nov 17, 2000
12.5MG;80MG N021162 002 Nov 17, 2000
+ 25MG;80MG N021162 003 Apr 19, 2004

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

<u>AB</u>	<u>+ GLAXOSMITHKLINE LLC</u>	<u>25MG;37.5MG</u>	<u>N016042</u>	<u>003</u>	Mar 03, 1994
	<u>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	DURAMED PHARMS BARR	<u>25MG;37.5MG</u>	<u>A075052</u>	<u>001</u>	Jun 18, 1999
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>25MG;50MG</u>	<u>A074259</u>	<u>001</u>	Mar 30, 1995
<u>AB</u>	LANNETT HOLDINGS INC	<u>25MG;37.5MG</u>	<u>A201407</u>	<u>001</u>	Dec 09, 2011
<u>AB</u>	MYLAN	<u>25MG;37.5MG</u>	<u>A074701</u>	<u>001</u>	Jun 07, 1996
<u>AB</u>	SANDOZ	<u>25MG;37.5MG</u>	<u>A074821</u>	<u>001</u>	Jun 05, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 219 (of 424)

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u> + SANDOZ	<u>25MG;50MG</u>	<u>A073191</u> <u>001</u>	Jul 31, 1991
TABLET; ORAL			
	<u>MAXZIDE</u>		
<u>AB</u> + MYLAN BERTEK	<u>50MG;75MG</u>	<u>N019129</u> <u>001</u>	Oct 22, 1984
	<u>MAXZIDE-25</u>		
<u>AB</u> MYLAN BERTEK	<u>25MG;37.5MG</u>	<u>N019129</u> <u>003</u>	May 13, 1988
<u>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</u>			
<u>AB</u> APOTEX INC	<u>25MG;37.5MG</u>	<u>A071251</u> <u>002</u>	May 05, 1998
<u>AB</u>	<u>50MG;75MG</u>	<u>A071251</u> <u>001</u>	Apr 17, 1988
<u>AB</u> PLIVA	<u>25MG;37.5MG</u>	<u>A074026</u> <u>001</u>	Apr 26, 1996
<u>AB</u>	<u>50MG;75MG</u>	<u>A073467</u> <u>001</u>	Jan 31, 1996
<u>AB</u> SANDOZ	<u>25MG;37.5MG</u>	<u>A073281</u> <u>001</u>	Apr 30, 1992
<u>AB</u>	<u>50MG;75MG</u>	<u>A072011</u> <u>001</u>	Jun 17, 1988
<u>AB</u> WATSON LABS	<u>25MG;37.5MG</u>	<u>A073449</u> <u>001</u>	Sep 23, 1993
<u>AB</u>	<u>50MG;75MG</u>	<u>A071851</u> <u>001</u>	Nov 30, 1988
<u>AB</u>	<u>50MG;75MG</u>	<u>A071969</u> <u>001</u>	Apr 17, 1988

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

NOVARTIS

12.5MG;80MG	N020818	001	Mar 06, 1998
12.5MG;160MG	N020818	002	Mar 06, 1998
12.5MG;320MG	N020818	004	Apr 28, 2006
25MG;160MG	N020818	003	Jan 17, 2002
+ 25MG;320MG	N020818	005	Apr 28, 2006

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

<u>AB</u> AMNEAL PHARMS NY	<u>5MG;200MG</u>	<u>A076642</u> <u>002</u>	Mar 18, 2004
	<u>7.5MG;200MG</u>	<u>A076642</u> <u>001</u>	Oct 12, 2004
<u>AB</u> TEVA	<u>7.5MG;200MG</u>	<u>A076023</u> <u>001</u>	Apr 11, 2003
<u>AB</u> VINTAGE PHARMS	<u>5MG;200MG</u>	<u>A077727</u> <u>001</u>	Nov 06, 2006
<u>AB</u>	<u>7.5MG;200MG</u>	<u>A077723</u> <u>001</u>	Nov 06, 2006
<u>AB</u>	<u>10MG;200MG</u>	<u>A077723</u> <u>002</u>	Nov 06, 2006
<u>AB</u> WATSON LABS FLORIDA	<u>5MG;200MG</u>	<u>A077454</u> <u>001</u>	Jun 23, 2010
<u>AB</u>	<u>7.5MG;200MG</u>	<u>A076604</u> <u>001</u>	Dec 31, 2003
<u>VICOPROFEN</u>			
<u>AB</u> + ABBOTT	<u>7.5MG;200MG</u>	<u>N020716</u> <u>001</u>	Sep 23, 1997
REPREXAIN			
AMNEAL PHARMS NY	2.5MG;200MG	A076642	Oct 19, 2007
	10MG;200MG	A076642	Oct 19, 2007

HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

REZIRA

+ CYPRESS PHARM	5MG/5ML;60MG/5ML	N022442	001	Jun 08, 2011
-----------------	------------------	---------	-----	--------------

HYDROCORTISONE

CREAM; TOPICAL

ALA-CORT

<u>AT</u> CROWN LABS	<u>1%</u>	<u>A080706</u> <u>006</u>	
<u>ANUSOL HC</u>			
<u>AT</u> SALIX PHARMS	<u>2.5%</u>	<u>A088250</u> <u>001</u>	Jun 06, 1984
<u>HYDROCORTISONE</u>			
<u>AT</u> ACTAVIS MID ATLANTIC	<u>1%</u>	<u>A087795</u> <u>001</u>	May 03, 1983
<u>AT</u>	<u>2.5%</u>	<u>A089682</u> <u>001</u>	Mar 10, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 220 (of 424)

HYDROCORTISONE

CREAM; TOPICAL			
<u>HYDROCORTISONE</u>			
AT	+ FOUGERA	<u>1%</u>	A080693 003
AT	+	<u>2.5%</u>	A089414 001
AT	LYNE	<u>2.5%</u>	A040879 001
AT	PERRIGO NEW YORK	<u>2.5%</u>	A085025 001
AT	TARO	<u>1%</u>	A086155 001
AT		<u>2.5%</u>	A088799 001
AT	VINTAGE PHARMS	<u>2.5%</u>	A040503 001
<u>SYNACORT</u>			
AT	MEDICIS	<u>1%</u>	A087458 001
AT		<u>2.5%</u>	A087457 001
ENEMA; RECTAL			
<u>COLOCORT</u>			
AB	PADDOCK LLC	<u>100MG/60ML</u>	A075172 001
<u>CORTENEMA</u>			
AB	+ ANI PHARMS	<u>100MG/60ML</u>	N016199 001
<u>HYDROCORTISONE</u>			
AB	TEVA PHARMS	<u>100MG/60ML</u>	A074171 001
LOTION; TOPICAL			
<u>ALA-CORT</u>			
AT	CROWN LABS	<u>1%</u>	A083201 001
<u>HYDROCORTISONE</u>			
AT	+ ALTANA	<u>2.5%</u>	A040351 001
AT	TARO	<u>2.5%</u>	A040247 001
AT	VINTAGE PHARMS	<u>2.5%</u>	A040417 001
<u>NUTRACORT</u>			
AT	CORIA	<u>1%</u>	A080443 003
AT		<u>2.5%</u>	A087644 001
<u>STIE-CORT</u>			
AT	PERRIGO	<u>1%</u>	A089066 001
AT		<u>2.5%</u>	A089074 001
ALA-SCALP			
	CROWN LABS	<u>2%</u>	A083231 001
OINTMENT; TOPICAL			
<u>HYDROCORTISONE</u>			
AT	ACTAVIS MID ATLANTIC	<u>1%</u>	A087796 001
AT	+ ALTANA	<u>1%</u>	A080692 001
AT	+ FOUGERA	<u>2.5%</u>	A081203 001
AT	PERRIGO NEW YORK	<u>2.5%</u>	A085027 001
AT	TARO	<u>1%</u>	A086257 001
AT		<u>2.5%</u>	A040310 001
<u>HYDROCORTISONE IN ABSORB BASE</u>			
AT	CAROLINA MEDCL	<u>1%</u>	A088138 001
POWDER; FOR RX COMPOUNDING			
HYDRO-RX			
+ X GEN PHARMS		100%	A085982 001
SOLUTION; TOPICAL			
TEXACORT			
+ MISSION PHARMA		2.5%	A081271 001
Apr 17, 1992			
TABLET; ORAL			
<u>CORTEF</u>			
AB	PHARMACIA AND UPJOHN	<u>5MG</u>	N008697 003
AB		<u>10MG</u>	N008697 001
AB	+	<u>20MG</u>	N008697 002
<u>HYDROCORTISONE</u>			
AB	COREPHARMA	<u>5MG</u>	A040646 001
AB		<u>10MG</u>	A040646 002
Mar 30, 2007			
Mar 30, 2007			

PRESCRIPTION DRUG PRODUCT LIST

3 - 221 (of 424)

HYDROCORTISONE

TABLET; ORAL

HYDROCORTISONE

<u>AB</u>	COREPHARMA	<u>20MG</u>	<u>A040646</u>	<u>003</u>	Mar 30, 2007
<u>AB</u>	VINTAGE	<u>5MG</u>	<u>A040761</u>	<u>001</u>	Jul 16, 2007
<u>AB</u>		<u>10MG</u>	<u>A040761</u>	<u>002</u>	Jul 16, 2007
<u>AB</u>		<u>20MG</u>	<u>A040761</u>	<u>003</u>	Jul 16, 2007
	HYDROCORTISONE				
BP	WEST WARD	20MG		A083365	001

HYDROCORTISONE ACETATEAEROSOL, METERED; RECTAL
CORTIFOAM

+ MEDA PHARMS 10%

N017351 001 Feb 10, 1982

CREAM; TOPICAL

HYDROCORTISONE ACETATE

+ FERNDALE LABS 2.5%

A040259 001 Jul 29, 1999

MICORT-HC

+ FERNDALE LABS 2%

A040398 001 Mar 29, 2002

2.5%

A040396 001 Feb 27, 2001

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; 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 MEDA PHARMS 1%;1% A086457 001

PROCTOFOAM HC

BX MEDA PHARMS 1%;1% A086195 001

CREAM; TOPICAL

PRAMOSONE

FERNDALE LABS 0.5%;1%

A083778 001

1%;1%

A085368 001

LOTION; TOPICAL

PRAMOSONE

FERNDALE LABS 1%;1%

A085980 001

2.5%;1%

A085979 001

HYDROCORTISONE ACETATE; UREA

CREAM; TOPICAL

CARMOL HC

AT NYCOMED US 1%;10% A080505 001

U-CORT

AT TARO 1%;10% A089472 001 Jun 13, 1988

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATE

AB TARO PHARM IND 0.1% A076654 001 Aug 03, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 222 (of 424)

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL <u>LOCOID</u>	<u>0.1%</u>	<u>N018514 001</u>	Mar 31, 1982
AB + TRIAX PHARMS LLC			
LOCOID LIPOCREAM			
+ TRIAX PHARMS LLC	<u>0.1%</u>	<u>N020769 001</u>	Sep 08, 1997
LOTION; TOPICAL LOCOID			
+ TRIAX PHARMS LLC	<u>0.1%</u>	<u>N022076 001</u>	May 18, 2007
OINTMENT; TOPICAL <u>HYDROCORTISONE BUTYRATE</u>	<u>0.1%</u>	<u>A076842 001</u>	Dec 27, 2004
AB TARO			
<u>LOCOID</u>			
AB + TRIAX PHARMS LLC	<u>0.1%</u>	<u>N018652 001</u>	Oct 29, 1982
SOLUTION; TOPICAL <u>HYDROCORTISONE BUTYRATE</u>	<u>0.1%</u>	<u>A076364 001</u>	Jan 14, 2004
AT TARO PHARM INDS			
<u>LOCOID</u>			
AT + TRIAX PHARMS LLC	<u>0.1%</u>	<u>N019116 001</u>	Feb 25, 1987

HYDROCORTISONE PROBUTATE

CREAM; TOPICAL PANDEL	<u>0.1%</u>	<u>N020453 001</u>	Feb 28, 1997
+ FOUGERA PHARMS			

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION <u>A-HYDROCORT</u>			
AP HOSPIRA	<u>EQ 100MG BASE/VIAL</u>	<u>A040666 001</u>	Apr 06, 2006
<u>SOLU-CORTEF</u>			
AP + PHARMACIA AND UPJOHN	<u>EQ 100MG BASE/VIAL</u>	<u>N009866 001</u>	
AP +	<u>EQ 250MG BASE/VIAL</u>	<u>N009866 002</u>	
AP +	<u>EQ 500MG BASE/VIAL</u>	<u>N009866 003</u>	
AP +	<u>EQ 1GM BASE/VIAL</u>	<u>N009866 004</u>	

HYDROCORTISONE VALERATE

CREAM; TOPICAL <u>HYDROCORTISONE VALERATE</u>	<u>0.2%</u>	<u>A075666 001</u>	May 24, 2000
AB PERRIGO NEW YORK			
AB TARO	<u>0.2%</u>	<u>A075042 001</u>	Aug 25, 1998
<u>WESTCORT</u>			
AB + RANBAXY	<u>0.2%</u>	<u>N017950 001</u>	
OINTMENT; TOPICAL <u>HYDROCORTISONE VALERATE</u>			
AB ALTANA	<u>0.2%</u>	<u>A075085 001</u>	Jul 31, 2001
AB TARO	<u>0.2%</u>	<u>A075043 001</u>	Aug 25, 1998
<u>WESTCORT</u>			
AB + RANBAXY	<u>0.2%</u>	<u>N018726 001</u>	Aug 08, 1983

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC <u>CORTISPORIN</u>			
AT + MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N050479 001</u>	
<u>NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE</u>			
AT ALCON	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062423 001</u>	Aug 25, 1983
AT BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064053 001</u>	Dec 29, 1995
AT LUITPOLD	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A065216 001</u>	Oct 31, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 223 (of 424)

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SUSPENSION/DROPS; OPHTHALMIC

+ ALCON UNIVERSAL	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062874 001	May 11, 1988
SUSPENSION/DROPS; OTIC			
<u>CORTISPORIN</u>			
AT + MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A060613 001</u>	
<u>NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE</u>			
AT ALCON UNIVERSAL	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062488 001</u>	Nov 06, 1985
AT LUITPOLD	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A065219 001</u>	May 01, 2006
<u>OTICAIR</u>			
AT BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064065 001</u>	Aug 28, 1996
<u>PEDIOTIC</u>			
AT MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062822 001</u>	Sep 29, 1987

HYDROFLUMETHIAZIDE

TABLET; ORAL

SALURON

AB + SHIRE	<u>50MG</u>	<u>N011949 001</u>
------------	-------------	--------------------

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDID

AP + PURDUE PHARM PRODS	<u>1MGM/ML</u>	<u>N019034 003</u>	Apr 30, 2009
AP +	<u>2MGM/ML</u>	<u>N019034 004</u>	Apr 30, 2009
AP +	<u>4MGM/ML</u>	<u>N019034 005</u>	Apr 30, 2009
<u>DILAUDID-HP</u>			
AP + PURDUE PHARM PRODS	<u>10MGM/ML</u>	<u>N019034 001</u>	Jan 11, 1984
<u>HYDROMORPHONE HYDROCHLORIDE</u>			
AP AKORN	<u>10MG/ML</u>	<u>A078228 001</u>	Apr 14, 2010
AP	<u>10MG/ML</u>	<u>A078261 001</u>	Apr 14, 2010
AP BARR	<u>10MG/ML</u>	<u>A076444 001</u>	Apr 25, 2003
AP HOSPIRA	<u>10MG/ML</u>	<u>A074598 001</u>	Jun 19, 1997
AP HOSPIRA INC	<u>1MGM/ML</u>	<u>N200403 001</u>	Dec 01, 2011
AP	<u>2MGM/ML</u>	<u>N200403 002</u>	Dec 01, 2011
AP	<u>4MGM/ML</u>	<u>N200403 003</u>	Dec 01, 2011
AP	<u>10MG/ML</u>	<u>A078591 001</u>	Jun 17, 2008
DILAUDID-HP			
+ PURDUE PHARM PRODS	250MG/VIAL	N019034 002	Aug 04, 1994

SOLUTION; ORAL

DILAUDID

AA + PURDUE PHARM PRODS	<u>5MGM/5ML</u>	<u>N019891 001</u>	Dec 07, 1992
<u>HYDROMORPHONE HYDROCHLORIDE</u>			
AA ROXANE	<u>5MGM/5ML</u>	<u>A074653 001</u>	Jul 29, 1998

TABLET; ORAL

DILAUDID

AB PURDUE PHARM PRODS	<u>2MG</u>	<u>N019892 003</u>	Nov 09, 2007
AB	<u>4MG</u>	<u>N019892 002</u>	Nov 09, 2007
AB +	<u>8MG</u>	<u>N019892 001</u>	Dec 07, 1992
<u>HYDROMORPHONE HYDROCHLORIDE</u>			
AB ELITE LABS	<u>8MG</u>	<u>A076723 001</u>	Oct 18, 2005
AB LANNETT	<u>2MG</u>	<u>A078439 001</u>	Dec 09, 2009
AB	<u>4MG</u>	<u>A078439 002</u>	Dec 09, 2009
AB	<u>8MG</u>	<u>A077471 001</u>	Dec 09, 2009
AB MALLINCKRODT	<u>8MG</u>	<u>A076855 001</u>	Dec 23, 2004
AB MALLINCKRODT INC	<u>2MG</u>	<u>A078273 001</u>	Sep 19, 2007
AB	<u>4MG</u>	<u>A078273 002</u>	Sep 19, 2007
AB NESHER PHARMS	<u>2MG</u>	<u>A077311 001</u>	Nov 09, 2005
AB	<u>4MG</u>	<u>A077311 002</u>	Nov 09, 2005
AB	<u>8MG</u>	<u>A077311 003</u>	Nov 09, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 224 (of 424)

HYDROMORPHONE HYDROCHLORIDE

TABLET; ORAL

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>	ROXANE	<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

MALLINCKRODT INC	8MG	N021217 001	Mar 01, 2010
	12MG	N021217 002	Mar 01, 2010
+	16MG	N021217 003	Mar 01, 2010

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+ MERCK SANTE SAS	5GM/VIAL (5GM/KIT)	N022041 001	Apr 08, 2011
HYDROXOCOBALAMIN			
+ WATSON LABS	1MG/ML	A085998 001	

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

+ AKORN	1%;0.25%	N019261 001	Jan 30, 1992
---------	----------	-------------	--------------

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<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>		<u>200MG</u>	<u>A040150 001</u>	Jan 27, 1996
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>A040081 001</u>	Sep 30, 1994
<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>A040133 001</u>	Nov 30, 1995
<u>AB</u>	WEST WARD	<u>200MG</u>	<u>A040760 001</u>	Aug 15, 2007
<u>AB</u>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A040657 001</u>	Sep 21, 2007
<u>PLAQUENIL</u>				
<u>AB</u>	+ SANOFI AVENTIS US	<u>200MG</u>	<u>N009768 001</u>	

HYDROXYPROGESTERONE CAPROATE

SOLUTION; INTRAMUSCULAR

MAKENA

+ KV PHARM	1250MG/5ML (250MG/ML)	N021945 001	Feb 03, 2011
------------	-----------------------	-------------	--------------

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>
<u>AB</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

PRESCRIPTION DRUG PRODUCT LIST

3 - 225 (of 424)

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

<u>AP</u> + APP PHARMS	<u>25MG/ML</u>	<u>A087329</u> <u>001</u>
<u>AP</u> +	<u>50MG/ML</u>	<u>A087329</u> <u>002</u>
<u>AP</u> LUITPOLD	<u>25MG/ML</u>	<u>A087408</u> <u>001</u>
<u>AP</u>	<u>50MG/ML</u>	<u>A087408</u> <u>002</u>

SYRUP; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AA</u> + HI TECH PHARMA	<u>10MG/5ML</u>	<u>A040010</u> <u>001</u>	Oct 28, 1994
<u>AA</u> + VINTAGE PHARMS	<u>10MG/5ML</u>	<u>A040391</u> <u>001</u>	Apr 10, 2002
<u>AA</u> + WOCKHARDT	<u>10MG/5ML</u>	<u>A087294</u> <u>001</u>	Apr 12, 1982

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u> AMNEAL PHARM	<u>10MG</u>	<u>A040808</u> <u>001</u>	Sep 24, 2008
<u>AB</u>	<u>25MG</u>	<u>A040808</u> <u>002</u>	Sep 24, 2008
<u>AB</u>	<u>50MG</u>	<u>A040808</u> <u>003</u>	Sep 24, 2008
<u>AB</u> HERITAGE PHARMS INC	<u>10MG</u>	<u>A040804</u> <u>001</u>	Jun 30, 2008
<u>AB</u>	<u>25MG</u>	<u>A040804</u> <u>002</u>	Jun 30, 2008
<u>AB</u>	<u>50MG</u>	<u>A040804</u> <u>003</u>	Jun 30, 2008
<u>AB</u> HETERO LABS UNIT III	<u>10MG</u>	<u>A040805</u> <u>001</u>	May 29, 2008
<u>AB</u>	<u>25MG</u>	<u>A040805</u> <u>002</u>	May 29, 2008
<u>AB</u>	<u>50MG</u>	<u>A040805</u> <u>003</u>	May 29, 2008
<u>AB</u> INVAGEN PHARMS	<u>10MG</u>	<u>A040812</u> <u>001</u>	Mar 12, 2008
<u>AB</u>	<u>25MG</u>	<u>A040812</u> <u>002</u>	Mar 12, 2008
<u>AB</u>	<u>50MG</u>	<u>A040812</u> <u>003</u>	Mar 12, 2008
<u>AB</u> KVK TECH	<u>10MG</u>	<u>A040786</u> <u>001</u>	Mar 20, 2007
<u>AB</u>	<u>25MG</u>	<u>A040787</u> <u>001</u>	Mar 20, 2007
<u>AB</u>	<u>50MG</u>	<u>A040788</u> <u>001</u>	Mar 20, 2007
<u>AB</u> MUTUAL PHARM	<u>10MG</u>	<u>A089381</u> <u>001</u>	May 19, 1986
<u>AB</u>	<u>25MG</u>	<u>A089382</u> <u>001</u>	May 19, 1986
<u>AB</u>	<u>50MG</u>	<u>A089383</u> <u>001</u>	May 19, 1986
<u>AB</u> MYLAN	<u>10MG</u>	<u>A091176</u> <u>001</u>	Jun 07, 2010
<u>AB</u>	<u>25MG</u>	<u>A091176</u> <u>002</u>	Jun 07, 2010
<u>AB</u>	<u>50MG</u>	<u>A091176</u> <u>003</u>	Jun 07, 2010
<u>AB</u> NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A040841</u> <u>001</u>	Mar 31, 2008
<u>AB</u>	<u>25MG</u>	<u>A040842</u> <u>001</u>	Mar 31, 2008
<u>AB</u>	<u>50MG</u>	<u>A040840</u> <u>001</u>	Mar 31, 2008
<u>AB</u> + PLIVA	<u>10MG</u>	<u>A088617</u> <u>001</u>	Jan 10, 1986
<u>AB</u> +	<u>25MG</u>	<u>A088618</u> <u>001</u>	Jan 10, 1986
<u>AB</u> +	<u>50MG</u>	<u>A088619</u> <u>001</u>	Jan 10, 1986
<u>AB</u> SUN PHARM INDs INC	<u>10MG</u>	<u>A040899</u> <u>001</u>	Jun 10, 2008
<u>AB</u>	<u>25MG</u>	<u>A040899</u> <u>002</u>	Jun 10, 2008
<u>AB</u>	<u>50MG</u>	<u>A040899</u> <u>003</u>	Jun 10, 2008
<u>AB</u> VINTAGE PHARMS	<u>10MG</u>	<u>A040579</u> <u>001</u>	May 27, 2005
<u>AB</u>	<u>25MG</u>	<u>A040574</u> <u>001</u>	May 27, 2005
<u>AB</u>	<u>50MG</u>	<u>A040580</u> <u>001</u>	May 27, 2005
<u>AB</u> WATSON LABS	<u>10MG</u>	<u>A081149</u> <u>001</u>	Mar 18, 1994
<u>AB</u>	<u>10MG</u>	<u>A088348</u> <u>001</u>	Sep 15, 1983
<u>AB</u>	<u>25MG</u>	<u>A081150</u> <u>001</u>	Mar 18, 1994
<u>AB</u>	<u>25MG</u>	<u>A088349</u> <u>001</u>	Sep 15, 1983
<u>AB</u>	<u>50MG</u>	<u>A081151</u> <u>001</u>	Mar 18, 1994
<u>AB</u>	<u>50MG</u>	<u>A088350</u> <u>001</u>	Sep 15, 1983

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u> BARR	<u>EQ 25MG HCL</u>	<u>A088496</u> <u>001</u>	Jun 15, 1984
<u>AB</u>	<u>EQ 50MG HCL</u>	<u>A088487</u> <u>001</u>	Jun 15, 1984
<u>AB</u> SANDOZ	<u>EQ 25MG HCL</u>	<u>A087479</u> <u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 226 (of 424)

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u>	SANDOZ	<u>EQ 50MG HCL</u>	<u>A086183 001</u>	
<u>AB</u>	WATSON LABS	<u>EQ 25MG HCL</u>	<u>A040156 001</u>	Jul 15, 1996
<u>AB</u>		<u>EQ 25MG HCL</u>	<u>A081165 001</u>	Jul 31, 1991
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>A040156 002</u>	Jul 15, 1996
	<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	EQ 100MG HCL	A088488 001	Jun 15, 1984
	SUSPENSION; ORAL			
	VISTARIL			
	+ PFIZER	EQ 25MG HCL/5ML	N011795 001	

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

+ ROCHE

EQ 3MG BASE/3ML

N021858 001 Jan 06, 2006

TABLET; ORAL

BONIVA

+ ROCHE

EQ 150MG BASE

N021455 002 Mar 24, 2005

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

CUMBERLAND PHARMS

400MG/4ML (100MG/ML)

N022348 001 Jun 11, 2009

+

800MG/8ML (100MG/ML)

N022348 002 Jun 11, 2009

SUSPENSION; ORAL

IBUPROFEN

<u>AB</u>	+	ACTAVIS MID ATLANTIC	<u>100MG/5ML</u>	<u>A074978 001</u>	Mar 25, 1998
<u>AB</u>		PERRIGO R AND D	<u>100MG/5ML</u>	<u>A076925 001</u>	Sep 23, 2004

TABLET; ORAL

IBUPROFEN

<u>AB</u>	AMNEAL PHARMS NY	<u>400MG</u>	<u>A071334 001</u>	Nov 25, 1986
<u>AB</u>		<u>400MG</u>	<u>A078558 001</u>	Jun 18, 2007
<u>AB</u>		<u>600MG</u>	<u>A071335 001</u>	Nov 25, 1986
<u>AB</u>		<u>600MG</u>	<u>A078558 002</u>	Jun 18, 2007
<u>AB</u>		<u>800MG</u>	<u>A071935 001</u>	Oct 13, 1987
<u>AB</u>		<u>800MG</u>	<u>A078558 003</u>	Jun 18, 2007
<u>AB</u>	CONTRACT PHARMACAL	<u>400MG</u>	<u>A071267 001</u>	Oct 15, 1986
<u>AB</u>		<u>600MG</u>	<u>A071268 001</u>	Oct 15, 1986
<u>AB</u>		<u>800MG</u>	<u>A072300 001</u>	Jul 01, 1988
<u>AB</u>	DR REDDYS LA	<u>400MG</u>	<u>A075682 001</u>	Nov 14, 2001
<u>AB</u>		<u>600MG</u>	<u>A075682 002</u>	Nov 14, 2001
<u>AB</u>	+	<u>800MG</u>	<u>A075682 003</u>	Nov 14, 2001
<u>AB</u>	DR REDDYS LABS INC	<u>400MG</u>	<u>A076112 001</u>	Oct 31, 2001
<u>AB</u>		<u>600MG</u>	<u>A076112 002</u>	Oct 31, 2001
<u>AB</u>		<u>800MG</u>	<u>A076112 003</u>	Oct 31, 2001
<u>AB</u>	MARKSANS PHARMA	<u>400MG</u>	<u>A090796 001</u>	Dec 21, 2010
<u>AB</u>		<u>600MG</u>	<u>A090796 002</u>	Dec 21, 2010
<u>AB</u>		<u>800MG</u>	<u>A090796 003</u>	Dec 21, 2010
<u>AB</u>	NORTHSTAR HLTHCARE	<u>400MG</u>	<u>A078132 001</u>	Sep 10, 2007
<u>AB</u>		<u>600MG</u>	<u>A078132 002</u>	Sep 10, 2007
<u>AB</u>		<u>800MG</u>	<u>A078132 003</u>	Sep 10, 2007
<u>AB</u>	OHM LABS	<u>400MG</u>	<u>A070818 001</u>	Dec 26, 1985
<u>AB</u>	PERRIGO R AND D	<u>400MG</u>	<u>A077114 001</u>	Jul 18, 2005
<u>AB</u>		<u>600MG</u>	<u>A077114 002</u>	Jul 18, 2005

PREScription DRUG PRODUCT LIST

3 - 227 (of 424)

IBUPROFEN

TABLET; ORAL

IBUPROFEN

<u>AB</u>	PERRIGO R AND D	<u>800MG</u>	<u>A077114</u>	<u>003</u>	Jul 18, 2005
<u>AB</u>	SHASUN USA	<u>400MG</u>	<u>A078329</u>	<u>001</u>	Feb 05, 2009
<u>AB</u>		<u>600MG</u>	<u>A078329</u>	<u>002</u>	Feb 05, 2009
<u>AB</u>		<u>800MG</u>	<u>A078329</u>	<u>003</u>	Feb 05, 2009
<u>AB</u>	VINTAGE PHARMS	<u>300MG</u>	<u>A071230</u>	<u>001</u>	Oct 22, 1986
<u>AB</u>		<u>400MG</u>	<u>A071231</u>	<u>001</u>	Oct 22, 1986
<u>AB</u>		<u>400MG</u>	<u>A071644</u>	<u>001</u>	Feb 01, 1988
<u>AB</u>		<u>600MG</u>	<u>A071232</u>	<u>001</u>	Oct 22, 1986
<u>AB</u>		<u>800MG</u>	<u>A072004</u>	<u>001</u>	Nov 18, 1987
<u>AB</u>	WATSON LABS	<u>400MG</u>	<u>A070436</u>	<u>001</u>	Aug 21, 1985
<u>AB</u>		<u>600MG</u>	<u>A070437</u>	<u>001</u>	Aug 21, 1985
<u>IBU-TAB</u>					
<u>AB</u>	ALRA	<u>400MG</u>	<u>A071058</u>	<u>001</u>	Aug 11, 1988
<u>AB</u>		<u>600MG</u>	<u>A071059</u>	<u>001</u>	Aug 11, 1988

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

NEOPROFEN

+ LUNDBECK INC EQ 20MG BASE/2ML (EQ 10MG BASE/ML) N021903 001 Apr 13, 2006

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

AB	ACTAVIS ELIZABETH	<u>400MG ; 5MG</u>	A078769	001	Jan 04, 2008
AB	+ BARR LABS INC	<u>400MG ; 5MG</u>	A078316	001	Nov 29, 2007
AB	WATSON LABS	<u>400MG ; 5MG</u>	A078394	001	Nov 26, 2007

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

CORVERT

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>0.1MG/ML</u>	<u>N020491</u>	<u>001</u>	Dec 28, 1995
<u>IBUTILIDE FUMARATE</u>					
<u>AP</u>	BIONICHE PHARMA USA	<u>0.1MG/ML</u>	<u>A090643</u>	<u>001</u>	Jan 11, 2010
<u>AP</u>		<u>0.1MG/ML</u>	<u>A090924</u>	<u>001</u>	Jan 11, 2010
<u>AP</u>	LUITPOLD	<u>0.1MG/ML</u>	<u>A090240</u>	<u>001</u>	Jan 11, 2010

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

N021321 001 Dec 20, 2002

TDABUBCTIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>1MG/ML</u>	<u>N050734</u>	<u>001</u>	Feb 17, 1997
	<u>IDARUBICIN HYDROCHLORIDE</u>				
<u>AP</u>	APP PHARMS	<u>1MG/ML</u>	<u>A065440</u>	<u>001</u>	Aug 04, 2009
<u>AP</u>	BEDFORD LABS	<u>1MG/ML</u>	<u>A065275</u>	<u>001</u>	Dec 14, 2006
<u>AP</u>		<u>1MG/ML</u>	<u>A065288</u>	<u>001</u>	May 15, 2007
<u>AP</u>	SANDOZ	<u>1MG/ML</u>	<u>A091293</u>	<u>001</u>	Mar 29, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 228 (of 424)

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

<u>IDARUBICIN HYDROCHLORIDE PFS</u>				
<u>AP</u>	TEVA PARENTERAL	<u>1MG/ML</u>	<u>A065036 001</u>	May 01, 2002
	IDARUBICIN HYDROCHLORIDE			
+	TEVA PARENTERAL	5MG/VIAL	A065037 003	May 01, 2002
+		10MG/VIAL	A065037 002	May 01, 2002
+		20MG/VIAL	A065037 001	May 01, 2002

IDOXURIDINE

SOLUTION/DROPS; OPHTHALMIC

<u>DENDRID</u>				
<u>AT</u>	+	ALCON	<u>0.1%</u>	<u>N014169 001</u>
<u>HERPLEX</u>				
<u>AT</u>	+	ALLERGAN	<u>0.1%</u>	<u>N013935 002</u>

IFOSFAMIDE

INJECTABLE; INJECTION

<u>IFEX</u>				
<u>AP</u>	BAXTER HLTHCARE	<u>1GM/VIAL</u>	<u>N019763 001</u>	Dec 30, 1988
<u>AP</u>		<u>3GM/VIAL</u>	<u>N019763 002</u>	Dec 30, 1988
<u>IFOSFAMIDE</u>				
<u>AP</u>	+	APP PHARMS	<u>1GM/VIAL</u>	<u>A076078 001</u>
<u>AP</u>			<u>1GM/20ML (50MG/ML)</u>	<u>A090181 001</u>
<u>AP</u>	+		<u>3GM/VIAL</u>	<u>A076078 002</u>
<u>AP</u>			<u>3GM/60ML (50MG/ML)</u>	<u>A090181 002</u>
<u>AP</u>	BEDFORD LABS	<u>1GM/20ML(50MG/ML)</u>	<u>A076619 001</u>	Jun 29, 2011
<u>AP</u>		<u>3GM/60ML(50MG/ML)</u>	<u>A076619 002</u>	Jun 29, 2011
<u>AP</u>	+	TEVA PARENTERAL	<u>1GM/20ML (50MG/ML)</u>	<u>A076657 001</u>
<u>AP</u>	+		<u>3GM/60ML (50MG/ML)</u>	<u>A076657 002</u>

IFOSFAMIDE; MESNA

INJECTABLE; INTRAVENOUS

IFOSFAMIDE/MESNA KIT

+	TEVA PARENTERAL	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

ILOPERIDONETABLET; ORAL
FANAPT

+	NOVARTIS	1MG	N022192 001	May 06, 2009
		2MG	N022192 002	May 06, 2009
		4MG	N022192 003	May 06, 2009
		6MG	N022192 004	May 06, 2009
		8MG	N022192 005	May 06, 2009
		10MG	N022192 006	May 06, 2009
		12MG	N022192 007	May 06, 2009

ILOPROST

SOLUTION; INHALATION

VENTAVIS

+	ACTELION PHARMS LTD	10MCG/ML (10MCG/ML)	N021779 002	Dec 08, 2005
+		20MCG/ML (20MCG/ML)	N021779 003	Aug 07, 2009

IMATINIB MESYLATETABLET; ORAL
GLEEVEC
NOVARTIS

+		EQ 100MG BASE	N021588 001	Apr 18, 2003
+		EQ 400MG BASE	N021588 002	Apr 18, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 229 (of 424)

IMIGLUCERASE

INJECTABLE; INJECTION

CEREZYME

GENZYME

200 UNITS/VIAL

+

400 UNITS/VIAL

N020367 001 May 23, 1994

N020367 002 Sep 22, 1999

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

<u>AB</u>	LUPIN LTD	<u>10MG</u>	<u>A090443</u> <u>001</u>	Mar 11, 2010
<u>AB</u>		<u>25MG</u>	<u>A090442</u> <u>001</u>	Mar 11, 2010
<u>AB</u>		<u>50MG</u>	<u>A090441</u> <u>001</u>	Mar 11, 2010
<u>AB</u>	MUTUAL PHARM	<u>10MG</u>	<u>A081048</u> <u>001</u>	Jun 05, 1990
<u>AB</u>		<u>25MG</u>	<u>A081049</u> <u>001</u>	Jun 05, 1990
<u>AB</u>		<u>50MG</u>	<u>A081050</u> <u>001</u>	Jun 05, 1990
<u>AB</u>	PAR PHARM	<u>10MG</u>	<u>A088292</u> <u>001</u>	Oct 21, 1983
<u>AB</u>		<u>10MG</u>	<u>A089422</u> <u>001</u>	Jul 14, 1987
<u>AB</u>		<u>25MG</u>	<u>A088262</u> <u>001</u>	Oct 21, 1983
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A084936</u> <u>002</u>	
<u>AB</u>		<u>25MG</u>	<u>A083745</u> <u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A084937</u> <u>001</u>	
<u>TOFRANIL</u>				
<u>AB</u>	MALLINCKRODT INC	<u>10MG</u>	<u>A087844</u> <u>001</u>	May 22, 1984
<u>AB</u>		<u>25MG</u>	<u>A087845</u> <u>001</u>	May 22, 1984
<u>AB</u> +		<u>50MG</u>	<u>A087846</u> <u>001</u>	May 22, 1984

IMIPRAMINE PAMOATE

CAPSULE; ORAL

IMIPRAMINE PAMOATE

<u>AB</u>	LUPIN LTD	<u>EQ 75MG HCL</u>	<u>A090444</u> <u>001</u>	Apr 16, 2010
<u>AB</u>		<u>EQ 100MG HCL</u>	<u>A090444</u> <u>002</u>	Apr 16, 2010
<u>AB</u>		<u>EQ 125MG HCL</u>	<u>A090444</u> <u>003</u>	Apr 16, 2010
<u>AB</u>		<u>EQ 150MG HCL</u>	<u>A090444</u> <u>004</u>	Apr 16, 2010
<u>AB</u>	ROXANE	<u>EQ 75MG HCL</u>	<u>A091099</u> <u>001</u>	Apr 16, 2010
<u>AB</u>		<u>EQ 100MG HCL</u>	<u>A091099</u> <u>002</u>	Apr 16, 2010
<u>AB</u>		<u>EQ 125MG HCL</u>	<u>A091099</u> <u>003</u>	Apr 16, 2010
<u>AB</u>		<u>EQ 150MG HCL</u>	<u>A091099</u> <u>004</u>	Apr 16, 2010
<u>TOFRANIL-PM</u>				
<u>AB</u> +	MALLINCKRODT INC	<u>EQ 75MG HCL</u>	<u>N017090</u> <u>001</u>	
<u>AB</u>		<u>EQ 100MG HCL</u>	<u>N017090</u> <u>004</u>	
<u>AB</u>		<u>EQ 125MG HCL</u>	<u>N017090</u> <u>003</u>	
<u>AB</u>		<u>EQ 150MG HCL</u>	<u>N017090</u> <u>002</u>	

IMIQUIMOD

CREAM; TOPICAL

ALDARA

<u>AB</u> +	MEDICIS	<u>5%</u>	<u>N020723</u> <u>001</u>	Feb 27, 1997
<u>IMIQUIMOD</u>				
<u>AB</u>	NYCOMED US	<u>5%</u>	<u>A078548</u> <u>001</u>	Feb 25, 2010
<u>AB</u>	PERRIGO ISRAEL	<u>5%</u>	<u>A078837</u> <u>001</u>	Sep 07, 2010
<u>AB</u>	TARO	<u>5%</u>	<u>A200173</u> <u>001</u>	Apr 15, 2011
<u>AB</u>	TEVA PHARMS USA	<u>5%</u>	<u>A200481</u> <u>001</u>	Apr 18, 2011
<u>AB</u>	TOLMAR	<u>5%</u>	<u>A091044</u> <u>001</u>	Feb 28, 2011
ZYCLARA				
+ MEDICIS		2.5%	N022483 002	Jul 15, 2011
+		3.75%	N022483 001	Mar 25, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 230 (of 424)

INAMRINONE LACTATE

INJECTABLE; INJECTION
 AMRINONE LACTATE
 + BEDFORD EQ 5MG BASE/ML A075513 001 May 09, 2000

INDACATEROL MALEATE

POWDER; INHALATION
 ARCAPTA NEOHALER
 + NOVARTIS EQ 75MCG BASE N022383 001 Jul 01, 2011

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>1.25MG</u>	<u>A074722</u> <u>001</u>	Jun 17, 1996
<u>AB</u>		<u>2.5MG</u>	<u>A074722</u> <u>002</u>	Jun 17, 1996
<u>AB</u>	ALPHAPHARM	<u>1.25MG</u>	<u>A075105</u> <u>001</u>	Jul 23, 1998
<u>AB</u>		<u>2.5MG</u>	<u>A075105</u> <u>002</u>	Jul 23, 1998
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>1.25MG</u>	<u>A074299</u> <u>002</u>	Apr 29, 1996
<u>AB</u>		<u>2.5MG</u>	<u>A074299</u> <u>001</u>	Jul 27, 1995
<u>AB</u>	MYLAN	<u>1.25MG</u>	<u>A074461</u> <u>002</u>	Mar 26, 1997
<u>AB</u> +		<u>2.5MG</u>	<u>A074461</u> <u>001</u>	Mar 27, 1996
<u>AB</u>	SANDOZ	<u>1.25MG</u>	<u>A074594</u> <u>001</u>	May 23, 1996
<u>AB</u>		<u>2.5MG</u>	<u>A074594</u> <u>002</u>	May 23, 1996
<u>AB</u>	WATSON LABS	<u>1.25MG</u>	<u>A074585</u> <u>001</u>	Sep 26, 1996
<u>AB</u>		<u>2.5MG</u>	<u>A074585</u> <u>002</u>	Sep 26, 1996

INDINAVIR SULFATE

CAPSULE; ORAL
 CRIXIVAN
 MERCK SHARP DOHME EQ 100MG BASE N020685 006 Apr 19, 2000
 EQ 200MG BASE N020685 003 Mar 13, 1996
 + EQ 400MG BASE N020685 001 Mar 13, 1996

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION
 INDICLOR
 + GE HEALTHCARE 2mCi/0.2ML N019862 001 Dec 29, 1992
 INDIUM IN 111 CHLORIDE
 + MALLINCKRODT 5mCi/0.5ML 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
 + MALLINCKRODT 3mCi/ML N020314 001 Jun 02, 1994

INDOCYANINE GREEN

INJECTABLE; INJECTION
IC-GREEN
AP + AKORN 25MG/VIAL N011525 001

PRESCRIPTION DRUG PRODUCT LIST

3 - 231 (of 424)

INDOCYANINE GREEN

INJECTABLE; INJECTION

INDOCYANINE GREENAP PULSION MEDCL 25MG/VIALA040811 001 Nov 21, 2007INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

AB	GLENMARK GENERICS	<u>25MG</u>	<u>A091276 001</u>	Dec 22, 2010
AB		<u>50MG</u>	<u>A091276 002</u>	Dec 22, 2010
AB	HERITAGE PHARMS INC	<u>25MG</u>	<u>N018851 001</u>	May 18, 1984
AB		<u>50MG</u>	<u>N018851 002</u>	May 18, 1984
AB	HETERO LABS UNIT III	<u>25MG</u>	<u>A091240 001</u>	Apr 12, 2011
AB		<u>50MG</u>	<u>A091240 002</u>	Apr 12, 2011
AB	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A070719 001</u>	Feb 12, 1986
AB		<u>50MG</u>	<u>A070756 001</u>	Feb 12, 1986
AB	MYLAN	<u>25MG</u>	<u>N018858 001</u>	Apr 20, 1984
AB	+	<u>50MG</u>	<u>A070624 001</u>	Sep 04, 1985
AB	SANDOZ	<u>25MG</u>	<u>A070673 001</u>	Apr 29, 1987
AB		<u>50MG</u>	<u>A070674 001</u>	Apr 29, 1987
AB	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A090403 001</u>	Nov 15, 2010
AB		<u>50MG</u>	<u>A090403 002</u>	Nov 15, 2010

CAPSULE, EXTENDED RELEASE; ORAL

INDOMETHACIN

AB	AMNEAL PHARMS	<u>75MG</u>	<u>A091549 001</u>	Dec 01, 2010
AB	AVANTHI INC	<u>75MG</u>	<u>A079175 001</u>	Mar 06, 2009
AB	PADDOCK LLC	<u>75MG</u>	<u>A200529 001</u>	Nov 30, 2010
AB	+	<u>SANDOZ</u>	<u>A074464 001</u>	May 28, 1998

INJECTABLE; INJECTION

INDOMETHACIN

+ APP PHARMS 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

INDOCINAP + LUNDBECK INC EQ 1MG BASE/VIALN018878 001 Jan 30, 1985AP INDOMETHACIN SODIUM EQ 1MG BASE/VIALA078713 001 Jul 16, 2008INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG MIX 70/30

+ NOVO NORDISK INC 70 UNITS/ML;30 UNITS/ML

N021172 001 Nov 01, 2001

INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS

NOVOLOG

+ NOVO NORDISK INC 100 UNITS/ML

N020986 001 Jun 07, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 232 (of 424)

INSULIN DETEMIR RECOMBINANT

INJECTABLE; SUBCUTANEOUS
 LEVEMIR
 + NOVO NORDISK INC 100 UNITS/ML N021536 001 Jun 16, 2005

INSULIN GLARGINE RECOMBINANT

INJECTABLE; INJECTION
 LANTUS
 + SANOFI AVENTIS US 100 UNITS/ML N021081 001 Apr 20, 2000

INSULIN GLULISINE RECOMBINANT

INJECTABLE; IV (INFUSION)-SC
 + 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

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 50/50 PEN
 + LILLY 50 UNITS/ML;50 UNITS/ML N021018 003 Dec 22, 1999
 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
 HUMALOG MIX 75/25 PEN
 + LILLY 75 UNITS/ML;25 UNITS/ML N021017 003 Dec 22, 1999

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
 HUMALOG PEN
 + LILLY 100 UNITS/ML N020563 002 Aug 06, 1998

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION
 HUMULIN R
 + LILLY 500 UNITS/ML N018780 004 Mar 31, 1994

IOBENGUANE SULFATE I-123

SOLUTION; INTRAVENOUS
 ADREVIEW
 + GE HEALTHCARE 10MCI/5ML (2MCI/ML) N022290 001 Sep 19, 2008

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION
 CHOLOGRAFIN MEGLUMINE
 + BRACCO 52% N009321 003

PRESCRIPTION DRUG PRODUCT LIST

3 - 233 (of 424)

IODIXANOL

INJECTABLE; INJECTION				
VISIPAQUE 270				
+ GE HEALTHCARE	55%		N020351 001	Mar 22, 1996
	55%		N020808 001	Aug 29, 1997
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

IOHEXOL

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
OMNIPAQ 240				
+ GE HEALTHCARE	51.8%		N018956 002	Dec 26, 1985
	51.8%		N020608 001	Oct 24, 1995
OMNIPAQ 300				
+ GE HEALTHCARE	64.7%		N018956 003	Dec 26, 1985
	64.7%		N020608 002	Oct 24, 1995

IOPAMIDOL

INJECTABLE; INJECTION				
<u>IOPAMIDOL-250</u>				
<u>AP APP PHARMS</u>	<u>51%</u>		<u>A074679 001</u>	Apr 02, 1997
<u>IOPAMIDOL-300</u>				
<u>AP APP PHARMS</u>	<u>61%</u>		<u>A074679 002</u>	Apr 02, 1997
<u>IOPAMIDOL-370</u>				
<u>AP APP PHARMS</u>	<u>76%</u>		<u>A074679 003</u>	Apr 02, 1997
<u>ISOVUE-200</u>				
<u>AP + BRACCO</u>	<u>41%</u>		<u>N018735 006</u>	Jul 07, 1987
<u>ISOVUE-250</u>				
<u>AP + BRACCO</u>	<u>51%</u>		<u>N018735 007</u>	Jul 06, 1992
<u>AP +</u>	<u>51%</u>		<u>N020327 002</u>	Oct 12, 1994
<u>ISOVUE-300</u>				
<u>AP + BRACCO</u>	<u>61%</u>		<u>N018735 002</u>	Dec 31, 1985
<u>AP +</u>	<u>61%</u>		<u>N020327 003</u>	Oct 12, 1994
<u>ISOVUE-370</u>				
<u>AP + BRACCO</u>	<u>76%</u>		<u>N018735 003</u>	Dec 31, 1985
<u>AP +</u>	<u>76%</u>		<u>N020327 004</u>	Oct 12, 1994
<u>SCANLUX-300</u>				
<u>AP SANOCHEMIA CORP USA</u>	<u>61%</u>		<u>A090394 001</u>	Jun 18, 2010
<u>SCANLUX-370</u>				
<u>AP SANOCHEMIA CORP USA</u>	<u>76%</u>		<u>A090394 002</u>	Jun 18, 2010
ISOVUE-M 200				
+ BRACCO	41%		N018735 001	Dec 31, 1985
ISOVUE-M 300				
+ BRACCO	61%		N018735 004	Dec 31, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 234 (of 424)

IOPROMIDE

INJECTABLE; INJECTION			
ULTRAVIST (PHARMACY BULK)			
+ BAYER HLTHCARE	49.9%	N021425 003	Mar 12, 2004
+	62.3%	N021425 001	Sep 20, 2002
+	76.9%	N021425 002	Sep 20, 2002
ULTRAVIST 150			
+ BAYER HLTHCARE	31.2%	N020220 004	May 10, 1995
ULTRAVIST 240			
+ BAYER HLTHCARE	49.9%	N020220 003	May 10, 1995
ULTRAVIST 300			
+ BAYER HLTHCARE	62.3%	N020220 002	May 10, 1995
ULTRAVIST 300 IN PLASTIC CONTAINER			
+ BAYER HLTHCARE	62.3%	N020220 005	Nov 18, 2008
ULTRAVIST 370			
+ BAYER HLTHCARE	76.9%	N020220 001	May 10, 1995

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION			
CONRAY			
+ MALLINCKRODT	60%	N013295 001	
CONRAY 30			
+ MALLINCKRODT	30%	N016983 001	
CONRAY 43			
+ MALLINCKRODT	43%	N013295 002	
SOLUTION; INTRAVESICAL			
CYSTO-CONRAY II			
MALLINCKRODT	17.2%	N017057 002	

LOVERSOL

INJECTABLE; INJECTION			
OPTIRAY 240			
+ MALLINCKRODT	51%	N019710 002	Dec 30, 1988
OPTIRAY 300			
+ MALLINCKRODT	64%	N019710 004	Jan 22, 1992
+	64%	N020923 004	May 13, 1999
OPTIRAY 320			
+ MALLINCKRODT	68%	N019710 001	Dec 30, 1988
OPTIRAY 350			
+ MALLINCKRODT	74%	N019710 005	Jan 22, 1992
+	74%	N020923 003	May 28, 1998

IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM

INJECTABLE; INJECTION			
HEXBABRIX			
+ 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

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION			
ATROVENT HFA			
+ BOEHRINGER INGELHEIM	0.021MG/INH	N021527 001	Nov 27, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 235 (of 424)

IPRATROPIUM BROMIDE

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

<u>AN</u>	BAUSCH AND LOMB	<u>0.02%</u>	<u>A075835</u>	<u>001</u>	Oct 15, 2001
<u>AN</u>	+ DEY	<u>0.02%</u>	<u>A074755</u>	<u>001</u>	Jan 10, 1997
<u>AN</u>	LANDELA PHARM	<u>0.02%</u>	<u>A077072</u>	<u>001</u>	Jul 19, 2005
<u>AN</u>	NEPHRON	<u>0.02%</u>	<u>A075562</u>	<u>001</u>	Sep 27, 2001
<u>AN</u>	NOVEX	<u>0.02%</u>	<u>A075441</u>	<u>001</u>	Mar 28, 2001
<u>AN</u>	RITEDOSE CORP	<u>0.02%</u>	<u>A075693</u>	<u>001</u>	Jan 26, 2001
<u>AN</u>	TEVA PARENTERAL	<u>0.02%</u>	<u>A075313</u>	<u>001</u>	Feb 07, 2000
<u>AN</u>	WATSON LABS	<u>0.02%</u>	<u>A076291</u>	<u>001</u>	May 09, 2005

SPRAY, METERED; NASAL

ATROVENT

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>0.021MG/SPRAY</u>	<u>N020393</u>	<u>001</u>	Oct 20, 1995
<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>0.042MG/SPRAY</u>	<u>N020394</u>	<u>001</u>	Oct 20, 1995

IPRATROPIUM BROMIDE

<u>AB</u>	BAUSCH AND LOMB	<u>0.021MG/SPRAY</u>	<u>A076025</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A076103</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>	DEY	<u>0.021MG/SPRAY</u>	<u>A075552</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A075553</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>	NOVEX	<u>0.021MG/SPRAY</u>	<u>A076156</u>	<u>001</u>	Apr 18, 2003
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A076155</u>	<u>001</u>	Apr 18, 2003
<u>AB</u>	ROXANE	<u>0.021MG/SPRAY</u>	<u>A076664</u>	<u>001</u>	Nov 05, 2003
<u>AB</u>		<u>0.042MG/SPRAY</u>	<u>A076598</u>	<u>001</u>	Nov 05, 2003

IRBESARTAN

TABLET; ORAL

AVAPRO

	SANOFI AVENTIS US	75MG	N020757	001	Sep 30, 1997
		150MG	N020757	002	Sep 30, 1997
+		300MG	N020757	003	Sep 30, 1997

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>	+ PFIZER INC	<u>100MG/5ML (20MG/ML)</u>	<u>N020571</u>	<u>002</u>	Jun 14, 1996

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>	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>	APP PHARMS	<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>	BEDFORD LABS	<u>40MG/2ML (20MG/ML)</u>	<u>A078753</u>	<u>001</u>	Dec 24, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078753</u>	<u>002</u>	Dec 24, 2008
<u>AP</u>	BIONICHE PHARMA	<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>	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>	EBEWE PHARMA	<u>40MG/2ML (20MG/ML)</u>	<u>A090137</u>	<u>001</u>	Nov 12, 2009
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090137</u>	<u>002</u>	Nov 12, 2009
<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>	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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 236 (of 424)

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

<u>AP</u>	+ HOSPIRA	<u>500MG/25ML (20MG/ML)</u>	<u>A078796 001</u>	Feb 27, 2008
<u>AP</u>	JIANGSU HENGRUI MED	<u>40MG/2ML (20MG/ML)</u>	<u>A090675 002</u>	Dec 16, 2011
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090675 001</u>	Dec 16, 2011
<u>AP</u>	PLIVA LACHEMA	<u>40MG/2ML (20MG/ML)</u>	<u>A078122 001</u>	Oct 31, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078122 002</u>	Oct 31, 2008
<u>AP</u>	SUN PHARMA GLOBAL	<u>40MG/2ML (20MG/ML)</u>	<u>A078805 001</u>	Apr 21, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078805 002</u>	Apr 21, 2008
<u>AP</u>	TEVA PARENTERAL	<u>40MG/2ML (20MG/ML)</u>	<u>A077260 001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077260 002</u>	Feb 27, 2008
<u>AP</u>		<u>500MG/25ML (20MG/ML)</u>	<u>A090101 001</u>	Nov 26, 2008
<u>AP</u>	WATSON LABS	<u>40MG/2ML (20MG/ML)</u>	<u>A077219 001</u>	Feb 20, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077219 002</u>	Feb 20, 2008
<u>AP</u>	X-GEN PHARMS	<u>40MG/2ML (20MG/ML)</u>	<u>A090016 001</u>	Jan 28, 2009
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090016 002</u>	Jan 28, 2009
	CAMPTOSAR			
+ PFIZER INC		<u>300MG/15ML (20MG/ML)</u>	<u>N020571 003</u>	Aug 05, 2010

IRON DEXTRAN

INJECTABLE; INJECTION

DEXFERRUM

<u>BP</u>	LUITPOLD	EQ 50MG IRON/ML	<u>N040024 001</u>	Feb 23, 1996
	INFED			
<u>BP</u>	+ WATSON LABS (UTAH)	EQ 50MG IRON/ML	<u>N017441 001</u>	
	PROFERDEX			
<u>BP</u>	NEW RIVER	EQ 50MG IRON/ML	<u>N017807 001</u>	

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

<u>LUITPOLD</u>	EQ 50MG BASE/2.5ML (EQ 20MG BASE/ML)	<u>N021135 002</u>	Mar 20, 2005
<u>+</u>	EQ 100MG BASE/5ML (EQ 20MG BASE/ML)	<u>N021135 001</u>	Nov 06, 2000
	EQ 200MG BASE/10ML (EQ 20MG BASE/ML)	<u>N021135 004</u>	Feb 09, 2007

ISOCARBOAZID

TABLET; ORAL

MARPLAN

<u>+</u>	VALIDUS PHARMS INC	10MG	<u>N011961 001</u>
----------	--------------------	------	--------------------

ISOFLURANE

LIQUID; INHALATION

FORANE

<u>AN</u>	+ BAXTER HLTHCARE CORP	<u>99.9%</u>	<u>N017624 001</u>
	<u>ISOFLURANE</u>		
<u>AN</u>	HALOCARBON PRODS	<u>99.9%</u>	<u>A075225 001</u>
<u>AN</u>	HOSPIRA	<u>99.9%</u>	<u>A074097 001</u>
<u>AN</u>	PIRAMAL CRITICAL	<u>99.9%</u>	<u>A074416 001</u>
<u>AN</u>	RHODIA	<u>99.9%</u>	<u>A074502 001</u>

ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

<u>+</u>	SANDOZ	100MG/ML	<u>A040648 001</u>	Jul 05, 2005
----------	--------	----------	--------------------	--------------

SYRUP; ORAL

ISONIAZID

<u>+</u>	CAROLINA MEDCL	50MG/5ML	<u>A088235 001</u>	Nov 10, 1983
----------	----------------	----------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 237 (of 424)

ISONIAZID

TABLET; ORAL

ISONIAZID

<u>AA</u>	BARR	<u>100MG</u>	<u>A080936</u>	<u>001</u>	
<u>AA</u>		<u>300MG</u>	<u>A080937</u>	<u>002</u>	
<u>AA</u>	MIKART	<u>100MG</u>	<u>A040090</u>	<u>001</u>	Jun 26, 1997
<u>AA</u>		<u>300MG</u>	<u>A040090</u>	<u>002</u>	Jun 26, 1997
<u>AA</u>	+ SANDOZ	<u>100MG</u>	<u>N008678</u>	<u>002</u>	
<u>AA</u>	+	<u>300MG</u>	<u>N008678</u>	<u>003</u>	
<u>AA</u>	WATSON LABS	<u>300MG</u>	<u>A080521</u>	<u>001</u>	
<u>AA</u>	WEST WARD	<u>100MG</u>	<u>A080212</u>	<u>001</u>	
<u>AA</u>		<u>300MG</u>	<u>A087425</u>	<u>001</u>	

ISONIAZID; PYRAZINAMIDE; RIFAMPIN

TABLET; ORAL

RIFATER

+ SANOFI AVENTIS US 50MG;300MG;120MG N050705 001 May 31, 1994

ISONIAZID; RIFAMPIN

CAPSULE; ORAL

RIFAMATE

<u>AB</u>	+ SANOFI AVENTIS US	<u>150MG;300MG</u>	<u>A061884</u>	<u>001</u>	
<u>AB</u>	<u>RIFAMPIN AND ISONIAZID</u>	<u>150MG;300MG</u>	<u>A065221</u>	<u>001</u>	Jul 29, 2005

ISOPROTERENOL HYDROCHLORIDE

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

<u>AP</u>	INTL MEDICATION	<u>0.2MG/ML</u>	<u>A083724</u>	<u>001</u>	
<u>AP</u>	+ HOSPIRA	<u>0.2MG/ML</u>	<u>N010515</u>	<u>001</u>	

ISOSORBIDE DINITRATECAPSULE, EXTENDED RELEASE; ORAL
DILATRATE-SR

+ SCHWARZ PHARMA 40MG N019790 001 Sep 02, 1988

TABLET; ORAL

ISORDIL

<u>AB</u>	VALEANT INTL	<u>5MG</u>	<u>N012093</u>	<u>007</u>	Jul 29, 1988
<u>AB</u>		<u>10MG</u>	<u>N012093</u>	<u>002</u>	Jul 29, 1988
<u>AB</u>		<u>20MG</u>	<u>N012093</u>	<u>006</u>	Jul 29, 1988
<u>AB</u>	+	<u>30MG</u>	<u>N012093</u>	<u>005</u>	Jul 29, 1988

ISOSORBIDE DINITRATE

<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>A086923</u>	<u>001</u>	Mar 12, 1987
<u>AB</u>		<u>10MG</u>	<u>A086925</u>	<u>001</u>	Mar 12, 1987
<u>AB</u>		<u>20MG</u>	<u>A087537</u>	<u>001</u>	Oct 02, 1987
<u>AB</u>		<u>30MG</u>	<u>A087946</u>	<u>001</u>	Jan 12, 1988
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A086221</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>		<u>10MG</u>	<u>A086223</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>		<u>20MG</u>	<u>A089367</u>	<u>001</u>	Apr 07, 1988
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>A086034</u>	<u>001</u>	Jan 06, 1988
<u>AB</u>		<u>10MG</u>	<u>A086032</u>	<u>001</u>	Jan 07, 1988
<u>AB</u>	WEST WARD	<u>5MG</u>	<u>A086067</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>		<u>10MG</u>	<u>A086066</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>		<u>20MG</u>	<u>A088088</u>	<u>001</u>	Nov 02, 1987
<u>AB</u>		<u>30MG</u>	<u>A040591</u>	<u>001</u>	Jan 10, 2007

ISORDIL

+ VALEANT INTL 40MG N012093 001 Jul 29, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 238 (of 424)

ISOSORBIDE DINITRATE

TABLET; SUBLINGUAL

ISOSORBIDE DINITRATE

<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A086033</u>	<u>001</u>	Feb 26, 1988
<u>AB</u>	+	<u>5MG</u>	<u>A086031</u>	<u>001</u>	Sep 29, 1987
<u>AB</u>	WEST WARD	<u>2.5MG</u>	<u>A086054</u>	<u>001</u>	Oct 29, 1987
<u>AB</u>		<u>5MG</u>	<u>A086055</u>	<u>001</u>	Nov 02, 1987

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE DINITRATE

<u>AB</u>	+	<u>CARACO</u>	<u>40MG</u>	<u>A040009</u>	<u>001</u>	Dec 30, 1998
<u>AB</u>		<u>COREPHARMA</u>	<u>40MG</u>	<u>A040723</u>	<u>001</u>	Mar 17, 2008

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075037</u>	<u>002</u>	Oct 30, 1998
<u>AB</u>		<u>20MG</u>	<u>A075037</u>	<u>001</u>	Oct 30, 1998
<u>AB</u>	TEVA	<u>20MG</u>	<u>A075147</u>	<u>001</u>	Nov 27, 1998
<u>AB</u>	WEST WARD	<u>20MG</u>	<u>A075361</u>	<u>001</u>	Oct 05, 2000
	<u>MONOKET</u>				
<u>AB</u>	UCB INC	<u>10MG</u>	<u>N020215</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>	+	<u>20MG</u>	<u>N020215</u>	<u>001</u>	Jun 30, 1993

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>30MG</u>	<u>A075306</u>	<u>001</u>	Dec 31, 1998
<u>AB</u>		<u>60MG</u>	<u>A075306</u>	<u>002</u>	Dec 31, 1998
<u>AB</u>	BRIGHTSTONE	<u>60MG</u>	<u>A075166</u>	<u>001</u>	Oct 07, 1999
<u>AB</u>	DEXCEL LTD	<u>60MG</u>	<u>A075522</u>	<u>001</u>	Apr 17, 2000
<u>AB</u>	ELAN PHARM	<u>60MG</u>	<u>A075041</u>	<u>001</u>	Sep 22, 1998
<u>AB</u>	KREMERS URBAN PHARMS	<u>30MG</u>	<u>A075155</u>	<u>002</u>	Jan 13, 2000
<u>AB</u>		<u>60MG</u>	<u>A075155</u>	<u>001</u>	Oct 30, 1998
<u>AB</u>	+	<u>120MG</u>	<u>A075155</u>	<u>003</u>	Aug 04, 2000
<u>AB</u>	NESHER PHARMS	<u>30MG</u>	<u>A075395</u>	<u>001</u>	Mar 16, 2000
<u>AB</u>		<u>60MG</u>	<u>A075395</u>	<u>002</u>	Mar 16, 2000
<u>AB</u>		<u>120MG</u>	<u>A075395</u>	<u>003</u>	Mar 16, 2000
<u>AB</u>	TORRENT PHARMS	<u>30MG</u>	<u>A200270</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>		<u>60MG</u>	<u>A200495</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>		<u>120MG</u>	<u>A200495</u>	<u>002</u>	Jun 03, 2011
<u>AB</u>	VINTAGE PHARMS	<u>30MG</u>	<u>A090598</u>	<u>001</u>	Aug 11, 2010
<u>AB</u>		<u>60MG</u>	<u>A090598</u>	<u>002</u>	Aug 11, 2010
<u>AB</u>		<u>120MG</u>	<u>A090598</u>	<u>003</u>	Aug 11, 2010
<u>AB</u>	WEST WARD	<u>30MG</u>	<u>A076813</u>	<u>002</u>	Mar 30, 2006
<u>AB</u>		<u>60MG</u>	<u>A076813</u>	<u>001</u>	Jan 07, 2005

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

<u>AP</u>	SYNERX	<u>1%</u>	<u>A090874</u>	<u>001</u>	Jul 20, 2010
<u>AP</u>	+	<u>LYMPHAZURIN</u>			
<u>AP</u>	COVIDIEN	<u>1%</u>	<u>N018310</u>	<u>001</u>	

ISOTRETINOIN

CAPSULE; ORAL

AMNESTEEM

<u>AB</u>	MYLAN	<u>10MG</u>	<u>A075945</u>	<u>001</u>	Nov 08, 2002
<u>AB</u>	+	<u>20MG</u>	<u>A075945</u>	<u>002</u>	Nov 08, 2002
<u>AB</u>	+	<u>40MG</u>	<u>A075945</u>	<u>003</u>	Nov 08, 2002

CLARAVIS

<u>AB</u>	BARR	<u>10MG</u>	<u>A076356</u>	<u>001</u>	Apr 11, 2003
-----------	------	-------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 239 (of 424)

ISOTRETINOIN

CAPSULE; ORAL				
CLARAVIS				
AB BARR	<u>20MG</u>		<u>A076135 002</u>	Apr 11, 2003
AB	<u>30MG</u>		<u>A076135 003</u>	May 11, 2006
AB	<u>40MG</u>		<u>A076135 001</u>	Apr 11, 2003
SOTRET				
AB RANBAXY	<u>10MG</u>		<u>A076041 001</u>	Dec 24, 2002
AB	<u>20MG</u>		<u>A076041 002</u>	Dec 24, 2002
AB	<u>30MG</u>		<u>A076503 001</u>	Jun 20, 2003
AB	<u>40MG</u>		<u>A076041 003</u>	Dec 24, 2002

ISRADIPINE

CAPSULE; ORAL				
ISRADIPINE				
AB MIKAH PHARMA	<u>2.5MG</u>		<u>A077169 001</u>	Apr 24, 2006
AB	<u>5MG</u>		<u>A077169 002</u>	Apr 24, 2006
AB WATSON LABS	<u>2.5MG</u>		<u>A077317 001</u>	Jan 05, 2006
AB +	<u>5MG</u>		<u>A077317 002</u>	Jan 05, 2006
TABLET, EXTENDED RELEASE; ORAL				
DYNACIRC CR				
GLAXOSMITHKLINE LLC	5MG		N020336 001	Jun 01, 1994
+ 10MG			N020336 002	Jun 01, 1994

ITRACONAZOLE

CAPSULE; ORAL				
ITRACONAZOLE				
AB SANDOZ	<u>100MG</u>		<u>A076104 001</u>	May 28, 2004
SPORANOX				
AB + JANSSEN PHARMS	<u>100MG</u>		<u>N020083 001</u>	Sep 11, 1992
SOLUTION; ORAL				
SPORANOX				
+ JANSSEN PHARMS	10MG/ML		N020657 001	Feb 21, 1997
TABLET; ORAL				
ONMEL				
+ STIEFEL LABS INC	200MG		N022484 001	Apr 29, 2010

IVERMECTIN

TABLET; ORAL				
STROMECTOL				
+ MERCK	3MG		N050742 002	Oct 08, 1998

IXABEPILONE

INJECTABLE; IV (INFUSION)				
+ BRISTOL MYERS SQUIBB	15MG/VIAL		N022065 001	Oct 16, 2007
+ 45MG/VIAL			N022065 002	Oct 16, 2007

KANAMYCIN SULFATE

INJECTABLE; INJECTION				
KANAMYCIN SULFATE				
APP PHARMS	EQ 500MG BASE/2ML		A065111 001	Dec 17, 2002
+ EQ 1GM BASE/3ML			A065111 002	Dec 17, 2002

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION				
KETALAR				
AP + JHP PHARMS	<u>EQ 10MG BASE/ML</u>		<u>N016812 001</u>	
AP +	<u>EQ 50MG BASE/ML</u>		<u>N016812 002</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 240 (of 424)

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALAR

<u>AP</u>	+ JHP PHARMS	<u>EQ 100MG BASE/ML</u>	<u>N016812 003</u>	
	<u>KETAMINE HYDROCHLORIDE</u>			
<u>AP</u>	BEDFORD	<u>EQ 50MG BASE/ML</u>	<u>A074524 001</u>	Mar 22, 1996
<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>A074524 002</u>	Mar 22, 1996
<u>AP</u>	BIONICHE PHARMA	<u>EQ 10MG BASE/ML</u>	<u>A076092 001</u>	Sep 30, 2008
<u>AP</u>		<u>EQ 50MG BASE/ML</u>	<u>A076092 002</u>	Dec 28, 2001
<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>A076092 003</u>	Oct 25, 2002
<u>AP</u>	HOSPIRA	<u>EQ 50MG BASE/ML</u>	<u>A074549 001</u>	Jun 27, 1996
<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>A074549 002</u>	Jun 27, 1996

KETOCONAZOLE

AEROSOL, FOAM; TOPICAL

EXTINA

<u>AT</u>	+ STIEFEL LABS INC	<u>2%</u>	<u>N021738 001</u>	Jun 12, 2007
	<u>KETOCONAZOLE</u>			

<u>AT</u>	PERRIGO ISRAEL	<u>2%</u>	<u>A091550 001</u>	Aug 25, 2011
	<u>KETOCONAZOLE</u>			

CREAM; TOPICAL

KETOCONAZOLE

<u>AB</u>	ALTANA	<u>2%</u>	<u>A076294 001</u>	Apr 28, 2004
	<u>KETOZOLE</u>			

<u>AB</u>	TARO	<u>2%</u>	<u>A075638 001</u>	Dec 18, 2002
-----------	------	-----------	--------------------	--------------

GEL; TOPICAL

XOLEGEL

+ AQUA PHARMS	2%	N021946 001	Jul 28, 2006
---------------	----	-------------	--------------

SHAMPOO; TOPICAL

KETOCONAZOLE

<u>AB</u>	PERRIGO NEW YORK	<u>2%</u>	<u>A076419 001</u>	Jan 07, 2004
	<u>NIZORAL</u>			

<u>AB</u>	+ JANSSEN PHARMS	<u>2%</u>	<u>N019927 001</u>	Aug 31, 1990
-----------	------------------	-----------	--------------------	--------------

TABLET; ORAL

KETOCONAZOLE

<u>AB</u>	APOTEX	<u>200MG</u>	<u>A075912 001</u>	Jan 10, 2002
<u>AB</u>	MUTUAL PHARMA	<u>200MG</u>	<u>A075314 001</u>	Jun 15, 1999
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A075597 001</u>	Dec 23, 1999
<u>AB</u>	PLIVA	<u>200MG</u>	<u>A075362 001</u>	Jun 15, 1999
<u>AB</u>	TARO	<u>200MG</u>	<u>A075319 001</u>	Jun 15, 1999
<u>AB</u>	TEVA	<u>200MG</u>	<u>A075273 001</u>	Jun 15, 1999

NIZORAL

<u>AB</u>	+ ORTHO MCNEIL JANSSEN	<u>200MG</u>	<u>N018533 001</u>	
-----------	------------------------	--------------	--------------------	--

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

<u>AB</u>	HERITAGE PHARMS INC	<u>50MG</u>	<u>A074014 002</u>	Jan 29, 1993
<u>AB</u>		<u>75MG</u>	<u>A074014 003</u>	Jan 29, 1993
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A074035 002</u>	Dec 31, 1996
<u>AB</u>		<u>75MG</u>	<u>A074035 003</u>	Dec 31, 1996
<u>AB</u>	TEVA	<u>50MG</u>	<u>A073516 001</u>	Dec 22, 1992
<u>AB</u>	+ ELAN PHARM	<u>75MG</u>	<u>A073517 001</u>	Dec 22, 1992

KETOPROFEN

HERITAGE PHARMS INC 25MG

A074014 001 Jan 29, 1993

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

<u>AB</u>	ELAN PHARM	<u>200MG</u>	<u>A074879 001</u>	Dec 10, 1997
-----------	------------	--------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 241 (of 424)

KETOPROFEN

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

<u>AB</u>	MYLAN	<u>100MG</u>	<u>A075679</u>	<u>003</u>	Feb 20, 2002
<u>AB</u>		<u>150MG</u>	<u>A075679</u>	<u>002</u>	Feb 20, 2002
<u>AB</u>	+	<u>200MG</u>	<u>A075679</u>	<u>001</u>	Feb 20, 2002
<u>AB</u>	WATSON LABS FLORIDA	<u>100MG</u>	<u>A075270</u>	<u>002</u>	Mar 24, 1999
<u>AB</u>		<u>150MG</u>	<u>A075270</u>	<u>003</u>	Mar 24, 1999
<u>AB</u>		<u>200MG</u>	<u>A075270</u>	<u>001</u>	Mar 24, 1999

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

<u>AP</u>	APP PHARMS	<u>15MG/ML</u>	<u>A075784</u>	<u>001</u>	Jan 11, 2002
<u>AP</u>		<u>30MG/ML</u>	<u>A075784</u>	<u>002</u>	Jan 11, 2002
<u>AP</u>	BAXTER HLTHCARE	<u>15MG/ML</u>	<u>A075772</u>	<u>001</u>	Jul 21, 2004
<u>AP</u>		<u>30MG/ML</u>	<u>A075772</u>	<u>002</u>	Jul 21, 2004
<u>AP</u>	BAXTER HLTHCARE CORP	<u>15MG/ML</u>	<u>A075299</u>	<u>001</u>	Nov 03, 1999
<u>AP</u>		<u>30MG/ML</u>	<u>A075299</u>	<u>002</u>	Nov 03, 1999
<u>AP</u>	+	<u>BEDFORD</u>	<u>A075222</u>	<u>001</u>	Apr 26, 1999
<u>AP</u>	+		<u>A075222</u>	<u>002</u>	Apr 26, 1999
<u>AP</u>	+		<u>A075228</u>	<u>001</u>	Apr 26, 1999
<u>AP</u>	CLARIS LIFESCIENCES	<u>15MG/ML</u>	<u>A075631</u>	<u>002</u>	Jun 29, 2001
<u>AP</u>		<u>30MG/ML</u>	<u>A075631</u>	<u>001</u>	Jun 29, 2001
<u>AP</u>	HOSPIRA	<u>15MG/ML</u>	<u>A074802</u>	<u>001</u>	Jun 05, 1997
<u>AP</u>		<u>15MG/ML</u>	<u>A074993</u>	<u>001</u>	Jan 27, 1999
<u>AP</u>		<u>30MG/ML</u>	<u>A074802</u>	<u>002</u>	Jun 05, 1997
<u>AP</u>		<u>30MG/ML</u>	<u>A074993</u>	<u>002</u>	Jan 27, 1999
<u>AP</u>	LUITPOLD	<u>15MG/ML</u>	<u>A078145</u>	<u>001</u>	Jan 14, 2008
<u>AP</u>		<u>30MG/ML</u>	<u>A078145</u>	<u>002</u>	Jan 14, 2008
<u>AP</u>	PFIZER	<u>15MG/ML</u>	<u>A078299</u>	<u>001</u>	Jul 16, 2007
<u>AP</u>		<u>30MG/ML</u>	<u>A078299</u>	<u>002</u>	Jul 16, 2007
<u>AP</u>	SANDOZ	<u>15MG/ML</u>	<u>A076271</u>	<u>001</u>	Oct 06, 2004
<u>AP</u>		<u>30MG/ML</u>	<u>A076271</u>	<u>002</u>	Oct 06, 2004
<u>AP</u>	SUN PHARMA GLOBAL	<u>15MG/ML</u>	<u>A078737</u>	<u>001</u>	Oct 06, 2008
<u>AP</u>		<u>30MG/ML</u>	<u>A078737</u>	<u>002</u>	Oct 06, 2008
<u>AP</u>	WOCKHARDT	<u>15MG/ML</u>	<u>A077942</u>	<u>001</u>	Mar 27, 2007
<u>AP</u>		<u>30MG/ML</u>	<u>A077942</u>	<u>002</u>	Mar 27, 2007
<u>AP</u>		<u>30MG/ML</u>	<u>A077943</u>	<u>001</u>	Mar 27, 2007

SOLUTION/DROPS; OPHTHALMIC

ACULAR

<u>AT</u>	+	ALLERGAN	<u>0.5%</u>	<u>N019700</u>	<u>001</u>	Nov 09, 1992
<u>AT</u>	+	ALLERGAN	<u>0.4%</u>	<u>N021528</u>	<u>001</u>	May 30, 2003

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>		<u>0.5%</u>	<u>A076583</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>	SUN PHARMA GLOBAL	<u>0.5%</u>	<u>A090017</u>	<u>001</u>	Nov 05, 2009

ACULAR PRESERVATIVE FREE

+ ALLERGAN	0.5%	N020811	001	Nov 03, 1997
ACUVAIL				
+ ALLERGAN	0.45%	N022427	001	Jul 22, 2009

SPRAY, METERED; NASAL

SPRIX

+ LUITPOLD	15.75MG/SPRAY	N022382	001	May 14, 2010
------------	---------------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 242 (of 424)

KETOROLAC TROMETHAMINE

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

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

<u>AP</u>	BEDFORD	<u>5MG/ML</u>	<u>A075303</u>	<u>001</u>	May 28, 1999
<u>AP</u>	CLARIS LIFESCIENCES	<u>5MG/ML</u>	<u>A076051</u>	<u>001</u>	Jul 05, 2002
<u>AP</u>	+ HOSPIRA	<u>5MG/ML</u>	<u>A075239</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>	+ TAYLOR	<u>5MG/ML</u>	<u>A075240</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>	TAYLOR	<u>5MG/ML</u>	<u>A075431</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>	SAGENT STRIDES	<u>5MG/ML</u>	<u>A075524</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>	<u>LABETALOL HYDROCHLORIDE</u>	<u>5MG/ML</u>	<u>A079134</u>	<u>001</u>	Feb 03, 2010

TABLET; ORAL

LABETALOL HYDROCHLORIDE

<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>	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

TRANDATE

<u>AB</u>	PROMETHEUS LABS	<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

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

+	INALCO	10GM/PACKET	A074712	001	Dec 10, 1997
+		20GM/PACKET	A074712	002	Dec 10, 1997

SOLUTION; ORAL

CONSTILAC

<u>AA</u>	ALRA	<u>10GM/15ML</u>	<u>A071054</u>	<u>001</u>	Jul 26, 1988
-----------	------	------------------	----------------	------------	--------------

CONSTULOSE

<u>AA</u>	+ ACTAVIS MID ATLANTIC	<u>10GM/15ML</u>	<u>A070288</u>	<u>001</u>	Aug 15, 1988
-----------	------------------------	------------------	----------------	------------	--------------

LACTULOSE

<u>AA</u>	ANI PHARMS	<u>10GM/15ML</u>	<u>A078430</u>	<u>001</u>	Nov 28, 2007
-----------	------------	------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 243 (of 424)

LACTULOSE

SOLUTION; ORAL

LACTULOSE

<u>AA</u>	HI TECH PHARMA	<u>10GM/15ML</u>	<u>A074076</u> <u>001</u>	Jul 03, 1995
<u>AA</u>	MORTON GROVE	<u>10GM/15ML</u>	<u>A074602</u> <u>001</u>	Nov 14, 1996
<u>AA</u>	NOVEX	<u>10GM/15ML</u>	<u>A075911</u> <u>001</u>	Feb 21, 2002
<u>AA</u>	PHARM ASSOC	<u>10GM/15ML</u>	<u>A074623</u> <u>001</u>	Jul 30, 1996
<u>AA</u>	ROXANE	<u>10GM/15ML</u>	<u>A073591</u> <u>001</u>	May 29, 1992
<u>AA</u>	VINTAGE PHARMS	<u>10GM/15ML</u>	<u>A075993</u> <u>001</u>	Jul 26, 2001
<u>AA</u>	VISTAPHARM	<u>10GM/15ML</u>	<u>A074138</u> <u>001</u>	Sep 30, 1992

SOLUTION; ORAL, RECTAL

CHOLAC

<u>AA</u>	ALRA	<u>10GM/15ML</u>	<u>A071331</u> <u>001</u>	Jul 26, 1988
<u>AA</u>	<u>ENULOSE</u>			
<u>AA</u>	+ ACTAVIS MID ATLANTIC	<u>10GM/15ML</u>	<u>A071548</u> <u>001</u>	Aug 15, 1988
<u>AA</u>	<u>GENERLAC</u>			
<u>AA</u>	MORTON GROVE PHARMS	<u>10GM/15ML</u>	<u>A074603</u> <u>001</u>	Oct 31, 1996
<u>AA</u>	<u>LACTULOSE</u>			
<u>AA</u>	ANI PHARMS	<u>10GM/15ML</u>	<u>A090426</u> <u>001</u>	Nov 21, 2008
<u>AA</u>	HI TECH PHARMA	<u>10GM/15ML</u>	<u>A074077</u> <u>001</u>	Jul 03, 1995
<u>AA</u>	NOVEX	<u>10GM/15ML</u>	<u>A076645</u> <u>001</u>	Jul 28, 2003

LAMIVUDINE

SOLUTION; ORAL

EPIVIR

+ VIIIV HLTHCARE	10MG/ML	N020596	001	Nov 17, 1995
EPIVIR-HBV				
+ GLAXOSMITHKLINE	5MG/ML	N021004	001	Dec 08, 1998

TABLET; ORAL

EPIVIR

<u>AB</u>	VIIIV HLTHCARE	<u>150MG</u>	<u>N020564</u> <u>001</u>	Nov 17, 1995
<u>AB</u>	+	<u>300MG</u>	<u>N020564</u> <u>003</u>	Jun 24, 2002
<u>AB</u>	<u>LAMIVUDINE</u>			
<u>AB</u>	APOTEX	<u>150MG</u>	<u>A091606</u> <u>001</u>	Dec 02, 2011
<u>AB</u>		<u>300MG</u>	<u>A091606</u> <u>002</u>	Dec 02, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>150MG</u>	<u>A202032</u> <u>001</u>	Nov 17, 2011
<u>AB</u>		<u>300MG</u>	<u>A202032</u> <u>002</u>	Nov 17, 2011
EPIVIR-HBV				
+ GLAXOSMITHKLINE	100MG	N021003	001	Dec 08, 1998

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

COMBIVIR

<u>AB</u>	+ VIIIV HLTHCARE	<u>150MG;300MG</u>	<u>N020857</u> <u>001</u>	Sep 26, 1997
<u>AB</u>	<u>LAMIVUDINE AND ZIDOVUDINE</u>			
<u>AB</u>	TEVA PHARMS	<u>150MG;300MG</u>	<u>A079081</u> <u>001</u>	May 25, 2011

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

<u>AB</u>	+ GLAXOSMITHKLINE	<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>	ACTAVIS TOTOWA	<u>25MG</u>	<u>A078669</u> <u>001</u>	Apr 08, 2011
<u>AB</u>		<u>100MG</u>	<u>A078669</u> <u>002</u>	Apr 08, 2011
<u>AB</u>		<u>150MG</u>	<u>A078669</u> <u>003</u>	Apr 08, 2011
<u>AB</u>		<u>200MG</u>	<u>A078669</u> <u>004</u>	Apr 08, 2011
<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090607</u> <u>001</u>	Jan 13, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 244 (of 424)

LAMOTRIGINE

TABLET; ORAL

LAMOTRIGINE

AB	ALEMBIC PHARMS LTD	<u>100MG</u>	<u>A090607</u>	<u>002</u>	Jan 13, 2011
AB		<u>150MG</u>	<u>A090607</u>	<u>003</u>	Jan 13, 2011
AB		<u>200MG</u>	<u>A090607</u>	<u>004</u>	Jan 13, 2011
AB	APOTEX INC	<u>25MG</u>	<u>A078625</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A078625</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A078625</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A078625</u>	<u>004</u>	Jan 27, 2009
AB	AUROBINDO PHARMA	<u>25MG</u>	<u>A078956</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A078956</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A078956</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A078956</u>	<u>004</u>	Jan 27, 2009
AB	CADISTA PHARMS	<u>25MG</u>	<u>A079132</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A079132</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A079132</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A079132</u>	<u>004</u>	Jan 27, 2009
AB	DR REDDYS LABS LTD	<u>25MG</u>	<u>A076708</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A076708</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A076708</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A076708</u>	<u>004</u>	Jan 27, 2009
AB	HIKMA PHARMS	<u>25MG</u>	<u>A078134</u>	<u>001</u>	Apr 19, 2011
AB		<u>100MG</u>	<u>A078134</u>	<u>002</u>	Apr 19, 2011
AB		<u>150MG</u>	<u>A078134</u>	<u>003</u>	Apr 19, 2011
AB		<u>200MG</u>	<u>A078134</u>	<u>004</u>	Apr 19, 2011
AB	LUPIN LTD	<u>25MG</u>	<u>A078691</u>	<u>001</u>	Jun 01, 2010
AB		<u>100MG</u>	<u>A078691</u>	<u>002</u>	Jun 01, 2010
AB		<u>150MG</u>	<u>A078691</u>	<u>003</u>	Jun 01, 2010
AB		<u>200MG</u>	<u>A078691</u>	<u>004</u>	Jun 01, 2010
AB	MYLAN	<u>25MG</u>	<u>A077420</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A077420</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A077420</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A077420</u>	<u>004</u>	Jan 27, 2009
AB	SANDOZ	<u>25MG</u>	<u>A078645</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A078645</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A078645</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A078645</u>	<u>004</u>	Jan 27, 2009
AB	TARO PHARM INDs	<u>25MG</u>	<u>A078525</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A078525</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A078525</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A078525</u>	<u>004</u>	Jan 27, 2009
AB	TEVA	<u>25MG</u>	<u>A076388</u>	<u>001</u>	Aug 30, 2006
AB		<u>100MG</u>	<u>A076388</u>	<u>002</u>	Aug 30, 2006
AB		<u>150MG</u>	<u>A076388</u>	<u>003</u>	Aug 30, 2006
AB		<u>200MG</u>	<u>A076388</u>	<u>004</u>	Aug 30, 2006
AB	TORRENT PHARMS	<u>25MG</u>	<u>A078947</u>	<u>001</u>	Jan 27, 2009
AB		<u>100MG</u>	<u>A078947</u>	<u>002</u>	Jan 27, 2009
AB		<u>150MG</u>	<u>A078947</u>	<u>003</u>	Jan 27, 2009
AB		<u>200MG</u>	<u>A078947</u>	<u>004</u>	Jan 27, 2009
AB	UNICHEM LABS LTD	<u>25MG</u>	<u>A090170</u>	<u>001</u>	Oct 06, 2011
AB		<u>100MG</u>	<u>A090170</u>	<u>002</u>	Oct 06, 2011
AB		<u>150MG</u>	<u>A090170</u>	<u>003</u>	Oct 06, 2011
AB		<u>200MG</u>	<u>A090170</u>	<u>004</u>	Oct 06, 2011
AB	UPSHER SMITH	<u>25MG</u>	<u>A078310</u>	<u>001</u>	Feb 04, 2009
AB		<u>100MG</u>	<u>A078310</u>	<u>002</u>	Feb 04, 2009
AB		<u>150MG</u>	<u>A078310</u>	<u>003</u>	Feb 04, 2009
AB		<u>200MG</u>	<u>A078310</u>	<u>004</u>	Feb 04, 2009
AB	WATSON LABS	<u>25MG</u>	<u>A077783</u>	<u>001</u>	Nov 01, 2010
AB		<u>100MG</u>	<u>A077783</u>	<u>002</u>	Nov 01, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 245 (of 424)

LAMOTRIGINE

TABLET; ORAL

LAMOTRIGINE

<u>AB</u>	WATSON LABS	<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>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077633</u> <u>001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A077633</u> <u>003</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A077633</u> <u>004</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A077633</u> <u>005</u>	Jan 27, 2009
	LAMOTRIGINE			
	ZYDUS PHARMS USA	50MG	A077633 002	Jan 27, 2009
		250MG	A077633 006	Jan 27, 2009

TABLET, CHEWABLE; ORAL

LAMICTAL CD

<u>AB</u>	GLAXOSMITHKLINE	<u>2MG</u>	<u>N020764</u> <u>004</u>	Sep 08, 2000
<u>AB</u>		<u>5MG</u>	<u>N020764</u> <u>001</u>	Aug 24, 1998
<u>AB</u>	+	<u>25MG</u>	<u>N020764</u> <u>002</u>	Aug 24, 1998
	<u>LAMOTRIGINE</u>			
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A090401</u> <u>002</u>	Nov 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A090401</u> <u>003</u>	Nov 04, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076701</u> <u>001</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076701</u> <u>002</u>	Jan 22, 2009
<u>AB</u>	GLENMARK GENERICS	<u>5MG</u>	<u>A079099</u> <u>001</u>	Feb 19, 2009
<u>AB</u>		<u>25MG</u>	<u>A079099</u> <u>002</u>	Feb 19, 2009
<u>AB</u>	JUBILANT LIFE	<u>5MG</u>	<u>A200220</u> <u>001</u>	Feb 28, 2011
<u>AB</u>		<u>25MG</u>	<u>A200220</u> <u>002</u>	Feb 28, 2011
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076630</u> <u>001</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076630</u> <u>002</u>	Jan 22, 2009
<u>AB</u>	TARO	<u>5MG</u>	<u>A079204</u> <u>001</u>	Feb 04, 2009
<u>AB</u>		<u>25MG</u>	<u>A079204</u> <u>002</u>	Feb 04, 2009
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076420</u> <u>001</u>	Jun 21, 2006
<u>AB</u>		<u>25MG</u>	<u>A076420</u> <u>002</u>	Jun 21, 2006
<u>AB</u>	WATSON LABS	<u>2MG</u>	<u>A076928</u> <u>001</u>	Jan 22, 2009
<u>AB</u>		<u>5MG</u>	<u>A076928</u> <u>002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A076928</u> <u>003</u>	Jan 22, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A078009</u> <u>002</u>	Jan 22, 2009
<u>AB</u>		<u>25MG</u>	<u>A078009</u> <u>003</u>	Jan 22, 2009

TABLET, EXTENDED RELEASE; ORAL

LAMICTAL XR

	SMITHKLINE BEECHAM	25MG	N022115 001	May 29, 2009
+		50MG	N022115 002	May 29, 2009
		100MG	N022115 003	May 29, 2009
		200MG	N022115 004	May 29, 2009
		250MG	N022115 006	Jun 21, 2011
		300MG	N022115 005	Apr 14, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

LAMICTAL ODT

	SMITHKLINE BEECHAM	25MG	N022251 001	May 08, 2009
+		50MG	N022251 002	May 08, 2009
		100MG	N022251 003	May 08, 2009
		200MG	N022251 004	May 08, 2009

LANREOTIDE ACETATEINJECTABLE; SUBCUTANEOUS
SOMATULINE DEPOT

+	IPSEN PHARMS	EQ 60MG BASE	N022074 001	Aug 30, 2007
+		EQ 90MG BASE	N022074 002	Aug 30, 2007
+		EQ 120MG BASE	N022074 003	Aug 30, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 246 (of 424)

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

LANSOPRAZOLE

<u>AB</u>	DR REDDYS LABS LTD	<u>15MG</u>	<u>A091269</u> <u>001</u>	Oct 15, 2010
<u>AB</u>		<u>30MG</u>	<u>A091269</u> <u>002</u>	Oct 15, 2010
<u>AB</u>	MATRIX LABS LTD	<u>15MG</u>	<u>A090763</u> <u>001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A090763</u> <u>002</u>	Nov 10, 2009
<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>	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>PREVACID</u>		
<u>AB</u>	TAKEDA PHARMS NA	<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, DELAYED RELEASE, ORALLY DISINTEGRATING; ORAL

LANSOPRAZOLE

<u>AB</u>	TEVA PHARMS	<u>15MG</u>	<u>A078730</u> <u>001</u>	Oct 15, 2010
<u>AB</u>		<u>30MG</u>	<u>A078730</u> <u>002</u>	Oct 15, 2010
		<u>PREVACID</u>		
<u>AB</u>	TAKEDA PHARMS NA	<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

TABLET, CHEWABLE; ORAL

FOSRENOL

	SHIRE	EQ 500MG BASE	<u>N021468</u> <u>002</u>	Oct 26, 2004
		EQ 750MG BASE	<u>N021468</u> <u>003</u>	Nov 23, 2005
	+	EQ 1GM BASE	<u>N021468</u> <u>004</u>	Nov 23, 2005

LAPATINIB DITOSYLATE

TABLET; ORAL

TYKERB

	+ SMITHKLINE BEECHAM	EQ 250MG BASE	<u>N022059</u> <u>001</u>	Mar 13, 2007
--	----------------------	---------------	---------------------------	--------------

LATANOPROST

SOLUTION/DROPS; OPHTHALMIC

LATANOPROST

<u>AT</u>	ALCON RES	<u>0.005%</u>	<u>A091449</u> <u>001</u>	Mar 22, 2011
<u>AT</u>	APOTEX	<u>0.005%</u>	<u>A077697</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>	LUITPOLD	<u>0.005%</u>	<u>A200925</u> <u>001</u>	Mar 22, 2011
<u>AT</u>	MYLAN	<u>0.005%</u>	<u>A201786</u> <u>001</u>	Mar 22, 2011
<u>AT</u>	PADDOCK LLC	<u>0.005%</u>	<u>A090887</u> <u>001</u>	Jul 19, 2011
		<u>XALATAN</u>		
<u>AT</u>	+ PHARMACIA AND UPJOHN	<u>0.005%</u>	<u>N020597</u> <u>001</u>	Jun 05, 1996

LEFLUNOMIDE

TABLET; ORAL

ARAVA

<u>AB</u>	SANOFI AVENTIS US	<u>10MG</u>	<u>N020905</u> <u>001</u>	Sep 10, 1998
<u>AB</u>	+	<u>20MG</u>	<u>N020905</u> <u>002</u>	Sep 10, 1998

LEFLUNOMIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>10MG</u>	<u>A091369</u> <u>001</u>	Nov 21, 2011
<u>AB</u>		<u>20MG</u>	<u>A091369</u> <u>002</u>	Nov 21, 2011
<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A077090</u> <u>001</u>	Sep 13, 2005
<u>AB</u>		<u>20MG</u>	<u>A077090</u> <u>002</u>	Sep 13, 2005
<u>AB</u>	BARR	<u>10MG</u>	<u>A077083</u> <u>001</u>	Sep 13, 2005
<u>AB</u>		<u>20MG</u>	<u>A077083</u> <u>002</u>	Sep 13, 2005
<u>AB</u>	HERITAGE PHARMS INC	<u>10MG</u>	<u>A077086</u> <u>001</u>	Sep 13, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 247 (of 424)

LEFLUNOMIDE

TABLET; ORAL

LEFLUNOMIDE

<u>AB</u>	HERITAGE PHARMS INC	<u>20MG</u>	<u>A077086</u>	<u>002</u>	Sep 13, 2005
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A077087</u>	<u>001</u>	Sep 13, 2005
<u>AB</u>		<u>20MG</u>	<u>A077087</u>	<u>002</u>	Sep 13, 2005
<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>A077084</u>	<u>001</u>	Sep 13, 2005
<u>AB</u>		<u>20MG</u>	<u>A077084</u>	<u>002</u>	Sep 13, 2005
	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
+		25MG	N021880	004	Jun 29, 2006

LEPIRUDIN RECOMBINANT

INJECTABLE; INJECTION

REFLUDAN

+	BAYER HLTHCARE	50MG/VIAL	N020807	001	Mar 06, 1998
---	----------------	-----------	---------	-----	--------------

LETROZOLE

TABLET; ORAL

FEMARA

<u>AB</u>	+ NOVARTIS PHARMS	<u>2.5MG</u>	<u>N020726</u>	<u>001</u>	Jul 25, 1997
	<u>LETROZOLE</u>				
<u>AB</u>	ACCORD HLTHCARE	<u>2.5MG</u>	<u>A090934</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	ACTAVIS TOTOWA	<u>2.5MG</u>	<u>A090292</u>	<u>001</u>	Jul 13, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>2.5MG</u>	<u>A091191</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	ENDO PHARMS	<u>2.5MG</u>	<u>A090789</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	FRESENIUS KABI ONCOL	<u>2.5MG</u>	<u>A090491</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	IMPAX LABS	<u>2.5MG</u>	<u>A091638</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	INDICUS PHARMA	<u>2.5MG</u>	<u>A201804</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	KUDCO IRELAND	<u>2.5MG</u>	<u>A091098</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A078190</u>	<u>001</u>	Dec 24, 2008
<u>AB</u>	NATCO PHARMA LTD	<u>2.5MG</u>	<u>A200161</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	ROXANE	<u>2.5MG</u>	<u>A090838</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	SUN PHARM INDs LTD	<u>2.5MG</u>	<u>A091466</u>	<u>001</u>	Jun 03, 2011
<u>AB</u>	TEVA PHARMS	<u>2.5MG</u>	<u>A090289</u>	<u>001</u>	Jun 03, 2011

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

<u>AP</u>	+ BEDFORD	<u>EQ 50MG BASE/VIAL</u>	<u>A089384</u>	<u>001</u>	Sep 14, 1987
<u>AP</u>	+	<u>EQ 100MG BASE/VIAL</u>	<u>A089717</u>	<u>001</u>	Mar 28, 1988
<u>AP</u>	TEVA PARENTERAL	<u>EQ 50MG BASE/VIAL</u>	<u>A081278</u>	<u>001</u>	Sep 28, 1993
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A081277</u>	<u>001</u>	Sep 28, 1993
<u>AP</u>		<u>EQ 350MG BASE/VIAL</u>	<u>A040174</u>	<u>001</u>	Jun 12, 1997
		<u>LEUCOVORIN CALCIUM PRESERVATIVE FREE</u>			
<u>AP</u>	APP PHARMS	<u>EQ 200MG BASE/VIAL</u>	<u>A040258</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>	+ BEDFORD	<u>EQ 200MG BASE/VIAL</u>	<u>A040056</u>	<u>001</u>	May 23, 1995
<u>AP</u>	+	<u>EQ 350MG BASE/VIAL</u>	<u>A040335</u>	<u>001</u>	Apr 20, 2000
<u>AP</u>	LUITPOLD	<u>EQ 50MG BASE/VIAL</u>	<u>A040338</u>	<u>001</u>	Jan 31, 2001
		<u>LEUCOVORIN CALCIUM PRESERVATIVE FREE</u>			
	+ APP PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A040286</u>	<u>001</u>	Feb 26, 1999
	+ BEDFORD	<u>EQ 10MG BASE/ML</u>	<u>A040347</u>	<u>001</u>	Apr 25, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 248 (of 424)

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

<u>AB</u>	BARR	<u>EQ 5MG BASE</u>	<u>A071198</u>	<u>001</u>	Sep 24, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A071199</u>	<u>001</u>	Sep 24, 1987
<u>AB</u>	ROXANE	<u>EQ 5MG BASE</u>	<u>A072733</u>	<u>001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A072735</u>	<u>001</u>	Feb 22, 1993
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>A072736</u>	<u>001</u>	Feb 22, 1993
	LEUCOVORIN CALCIUM				
	ROXANE	<u>EQ 10MG BASE</u>	<u>A072734</u>	<u>001</u>	Feb 22, 1993

LEUPROLIDE ACETATE

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

<u>AP</u>	+	SANDOZ	<u>1MIG/0.2ML</u>	<u>A074728</u>	<u>001</u>	Aug 04, 1998
<u>AP</u>		SUN PHARMA GLOBAL	<u>1MIG/0.2ML</u>	<u>A078885</u>	<u>001</u>	Mar 09, 2009
<u>AP</u>		TEVA PARENTERAL	<u>1MIG/0.2ML</u>	<u>A075471</u>	<u>001</u>	Oct 25, 2000
	LUPRON DEPOT					
	+	ABBOTT ENDOCRINE	22.5MIG/VIAL	N020517	001	Dec 22, 1995
	+		30MIG/VIAL	N020517	002	May 30, 1997
			45MIG/VIAL	N020517	003	Jun 17, 2011
	+	ABBOTT LABS	3.75MIG/VIAL	N020011	001	Oct 22, 1990
	+		7.5MIG/VIAL	N019732	001	Jan 26, 1989
	+		11.25MIG/VIAL	N020708	001	Mar 07, 1997
	LUPRON DEPOT-PED					
	+	ABBOTT ENDOCRINE	7.5MIG/VIAL	N020263	002	Apr 16, 1993
	+		11.25MIG/VIAL	N020263	005	Jan 21, 1994
	+		11.25MIG/VIAL	N020263	007	Aug 15, 2011
	+		15MIG/VIAL	N020263	006	Jan 21, 1994
	+		30MIG/VIAL	N020263	008	Aug 15, 2011

INJECTABLE; SUBCUTANEOUS

ELIGARD

+	TOLMAR THERAP	7.5MIG/VIAL	N021343	001	Jan 23, 2002
+		22.5MIG/VIAL	N021379	001	Jul 24, 2002
+		30MIG/VIAL	N021488	001	Feb 13, 2003
+		45MIG/VIAL	N021731	001	Dec 14, 2004

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

<u>AN</u>	DEY	<u>EQ 0.25% BASE</u>	<u>A078309</u>	<u>001</u>	Mar 20, 2009	
<u>AN</u>	WATSON LABS INC	<u>EQ 0.0103% BASE</u>	<u>A077756</u>	<u>003</u>	Apr 09, 2008	
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A077756</u>	<u>001</u>	Apr 09, 2008	
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A077756</u>	<u>002</u>	Apr 09, 2008	
	XOPENEX					
<u>AN</u>	+	SUNOVION	<u>EQ 0.0103% BASE</u>	<u>N020837</u>	<u>003</u>	Jan 30, 2002
<u>AN</u>	+		<u>EQ 0.021% BASE</u>	<u>N020837</u>	<u>001</u>	Mar 25, 1999
<u>AN</u>	+		<u>EQ 0.042% BASE</u>	<u>N020837</u>	<u>002</u>	Mar 25, 1999
<u>AN</u>	+		<u>EQ 0.25% BASE</u>	<u>N020837</u>	<u>004</u>	Jul 18, 2003

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+	SUNOVION	EQ 0.045MIG BASE/INH	N021730	001	Mar 11, 2005
---	----------	----------------------	---------	-----	--------------

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

KEPPRA

<u>AP</u>	+	UCB INC	<u>500MIG/5ML (100MIG/ML)</u>	<u>N021872</u>	<u>001</u>	Jul 31, 2006
-----------	---	---------	-------------------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 249 (of 424)

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

LEVETIRACETAM

<u>AP</u>	HIKMA FARMACEUTICA	<u>500MG/5ML(100MG/ML)</u>	<u>A090981</u> <u>001</u>	Oct 13, 2011
<u>AP</u>	INNOPHARMA LLC	<u>500MG/5ML (100MG/ML)</u>	<u>A091485</u> <u>001</u>	Aug 05, 2011
<u>AP</u>	NEXUS PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A090813</u> <u>001</u>	May 26, 2010
<u>AP</u>	SUN PHARM INDs LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A090754</u> <u>001</u>	Jun 16, 2010
	LEVETIRACETAM IN SODIUM CHLORIDE			
+	HQ SPECIALITY PHARMA	<u>500MG/100ML (5MG/ML)</u>	N202543 <u>001</u>	Nov 09, 2011
+		<u>1000MG/100ML (10MG/ML)</u>	N202543 <u>002</u>	Nov 09, 2011
+		<u>1500MG/100ML (15MG/ML)</u>	N202543 <u>003</u>	Nov 09, 2011

SOLUTION; ORAL

KEPPRA

<u>AA</u>	+ UCB INC	<u>100MG/ML</u>	<u>N021505</u> <u>001</u>	Jul 15, 2003
	<u>LEVETIRACETAM</u>			
<u>AA</u>	ACTAVIS MID ATLANTIC	<u>100MG/ML</u>	<u>A078976</u> <u>001</u>	Jan 15, 2009
<u>AA</u>	AMNEAL PHARMS	<u>100MG/ML</u>	<u>A090992</u> <u>001</u>	Oct 27, 2009
<u>AA</u>	APOTEX	<u>100MG/ML</u>	<u>A090187</u> <u>001</u>	Aug 05, 2011
<u>AA</u>	AUROBINDO PHARM	<u>100MG/ML</u>	<u>A079063</u> <u>001</u>	Jan 15, 2009
<u>AA</u>	CYPRESS PHARM	<u>100MG/ML</u>	<u>A079120</u> <u>001</u>	Jan 16, 2009
<u>AA</u>	LUPIN LTD	<u>100MG/ML</u>	<u>A090893</u> <u>001</u>	Oct 17, 2011
<u>AA</u>	ROXANE	<u>100MG/ML</u>	<u>A078582</u> <u>001</u>	Jan 15, 2009
<u>AA</u>	SILARX	<u>100MG/ML</u>	<u>A090263</u> <u>001</u>	Apr 03, 2009
<u>AA</u>	TARO	<u>100MG/ML</u>	<u>A078774</u> <u>001</u>	Feb 10, 2009
<u>AA</u>	TOLMAR	<u>100MG/ML</u>	<u>A079107</u> <u>001</u>	Jan 15, 2009
<u>AA</u>	TRIS PHARMA INC	<u>100MG/ML</u>	<u>A090461</u> <u>001</u>	Sep 30, 2010
<u>AA</u>	WOCKHARDT	<u>100MG/ML</u>	<u>A090028</u> <u>001</u>	Mar 03, 2010

TABLET; ORAL

KEPPRA

<u>AB</u>	UCB INC	<u>250MG</u>	<u>N021035</u> <u>001</u>	Nov 30, 1999
<u>AB</u>		<u>500MG</u>	<u>N021035</u> <u>002</u>	Nov 30, 1999
<u>AB</u>		<u>750MG</u>	<u>N021035</u> <u>003</u>	Nov 30, 1999
<u>AB</u>	+	<u>1GM</u>	<u>N021035</u> <u>004</u>	Jan 06, 2006
	<u>LEVETIRACETAM</u>			
<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A090843</u> <u>001</u>	Feb 14, 2011
<u>AB</u>		<u>500MG</u>	<u>A090843</u> <u>002</u>	Feb 14, 2011
<u>AB</u>		<u>750MG</u>	<u>A090843</u> <u>003</u>	Feb 14, 2011
<u>AB</u>		<u>1GM</u>	<u>A090843</u> <u>004</u>	Feb 14, 2011
<u>AB</u>	AJANTA PHARMA	<u>250MG</u>	<u>A201293</u> <u>001</u>	Jun 14, 2011
<u>AB</u>		<u>500MG</u>	<u>A201293</u> <u>002</u>	Jun 14, 2011
<u>AB</u>		<u>750MG</u>	<u>A201293</u> <u>003</u>	Jun 14, 2011
<u>AB</u>		<u>1GM</u>	<u>A201293</u> <u>004</u>	Jun 14, 2011
<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A078869</u> <u>001</u>	Mar 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A078869</u> <u>002</u>	Mar 13, 2009
<u>AB</u>		<u>750MG</u>	<u>A078869</u> <u>003</u>	Mar 13, 2009
<u>AB</u>		<u>1GM</u>	<u>A078869</u> <u>004</u>	Mar 13, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A078993</u> <u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078993</u> <u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078993</u> <u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078993</u> <u>004</u>	Jan 15, 2009
<u>AB</u>	BIOKEY	<u>500MG</u>	<u>A090906</u> <u>001</u>	Nov 05, 2010
<u>AB</u>	BOCA PHARMA	<u>250MG</u>	<u>A077319</u> <u>001</u>	Mar 20, 2009
<u>AB</u>		<u>500MG</u>	<u>A077319</u> <u>002</u>	Mar 20, 2009
<u>AB</u>		<u>750MG</u>	<u>A077319</u> <u>003</u>	Mar 20, 2009
<u>AB</u>	BRECKENRIDGE PHARM	<u>250MG</u>	<u>A090511</u> <u>001</u>	Aug 18, 2011
<u>AB</u>		<u>500MG</u>	<u>A090511</u> <u>002</u>	Aug 18, 2011
<u>AB</u>		<u>750MG</u>	<u>A090511</u> <u>003</u>	Aug 18, 2011
<u>AB</u>		<u>1GM</u>	<u>A090511</u> <u>004</u>	Aug 18, 2011
<u>AB</u>	COBALT LABS INC	<u>250MG</u>	<u>A077384</u> <u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A077384</u> <u>002</u>	Jan 15, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 250 (of 424)

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

AB	COBALT LABS INC	<u>750MG</u>	<u>A077384</u> <u>003</u>	Jan 15, 2009
AB	DR REDDYS LABS LTD	<u>250MG</u>	<u>A076920</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A076920</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A076920</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A078904</u> <u>001</u>	Jan 15, 2009
AB	HETERO DRUGS LTD	<u>250MG</u>	<u>A090515</u> <u>001</u>	Oct 08, 2010
AB		<u>500MG</u>	<u>A090515</u> <u>002</u>	Oct 08, 2010
AB		<u>750MG</u>	<u>A090515</u> <u>003</u>	Oct 08, 2010
AB		<u>1GM</u>	<u>A090515</u> <u>004</u>	Oct 08, 2010
AB	INVAGEN PHARMS	<u>250MG</u>	<u>A078234</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A078234</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A078234</u> <u>003</u>	Jan 15, 2009
AB	LUPIN	<u>250MG</u>	<u>A078154</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A078154</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A078154</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A090025</u> <u>001</u>	Jan 15, 2009
AB	METHAPHARM	<u>250MG</u>	<u>A090767</u> <u>001</u>	Jul 28, 2010
AB		<u>500MG</u>	<u>A090767</u> <u>002</u>	Jul 28, 2010
AB		<u>750MG</u>	<u>A090767</u> <u>003</u>	Jul 28, 2010
AB		<u>1GM</u>	<u>A090767</u> <u>004</u>	Jul 28, 2010
AB	MYLAN	<u>250MG</u>	<u>A076919</u> <u>001</u>	Nov 04, 2008
AB		<u>500MG</u>	<u>A076919</u> <u>002</u>	Nov 04, 2008
AB		<u>750MG</u>	<u>A076919</u> <u>003</u>	Nov 04, 2008
AB		<u>1GM</u>	<u>A090261</u> <u>001</u>	Dec 08, 2009
AB	ORCHID HLTHCARE	<u>250MG</u>	<u>A078526</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A078526</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A078526</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A090484</u> <u>001</u>	Aug 05, 2010
AB	ROXANE	<u>250MG</u>	<u>A078042</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A078042</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A078042</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A078042</u> <u>004</u>	Jan 15, 2009
AB	SANDOZ	<u>250MG</u>	<u>A077324</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A077324</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A077324</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A077324</u> <u>004</u>	Jan 15, 2009
AB	SOLCO HLTHCARE	<u>250MG</u>	<u>A078106</u> <u>001</u>	Feb 10, 2009
AB		<u>500MG</u>	<u>A078106</u> <u>002</u>	Feb 10, 2009
AB		<u>750MG</u>	<u>A078106</u> <u>003</u>	Feb 10, 2009
AB		<u>1GM</u>	<u>A078106</u> <u>004</u>	Feb 10, 2009
AB	TARO	<u>250MG</u>	<u>A078960</u> <u>004</u>	Feb 01, 2010
AB		<u>500MG</u>	<u>A078960</u> <u>003</u>	Feb 01, 2010
AB		<u>750MG</u>	<u>A078960</u> <u>002</u>	Feb 01, 2010
AB		<u>1GM</u>	<u>A078960</u> <u>001</u>	Feb 01, 2010
AB	TEVA PHARMS	<u>250MG</u>	<u>A078101</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A078101</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A078101</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A078101</u> <u>004</u>	Jan 15, 2009
AB	TORRENT PHARMS	<u>250MG</u>	<u>A078858</u> <u>001</u>	Jan 15, 2009
AB		<u>500MG</u>	<u>A078858</u> <u>002</u>	Jan 15, 2009
AB		<u>750MG</u>	<u>A078858</u> <u>003</u>	Jan 15, 2009
AB		<u>1GM</u>	<u>A078858</u> <u>004</u>	Jan 15, 2009
AB	VINTAGE PHARMS	<u>250MG</u>	<u>A091491</u> <u>001</u>	Dec 14, 2010
AB		<u>500MG</u>	<u>A091491</u> <u>002</u>	Dec 14, 2010
AB		<u>750MG</u>	<u>A091491</u> <u>003</u>	Dec 14, 2010
AB		<u>1GM</u>	<u>A091491</u> <u>004</u>	Dec 14, 2010
AB	WATSON LABS	<u>1GM</u>	<u>A078797</u> <u>001</u>	Jan 15, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 251 (of 424)

LEVETIRACETAM

TABLET; ORAL

LEVETIRACETAM

<u>AB</u>	WOCHARDT	<u>250MG</u>	<u>A079042</u> <u>001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A079042</u> <u>002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A079042</u> <u>003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A079042</u> <u>004</u>	Jan 15, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A078918</u> <u>001</u>	Apr 29, 2009
<u>AB</u>		<u>1GM</u>	<u>A078918</u> <u>002</u>	Apr 29, 2009

TABLET, EXTENDED RELEASE; ORAL

KEPPRA XR

<u>AB</u>	UCB INC	<u>500MG</u>	<u>N022285</u> <u>001</u>	Sep 12, 2008
<u>AB</u>	+	<u>750MG</u>	<u>N022285</u> <u>002</u>	Feb 12, 2009

LEVETIRACETAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>500MG</u>	<u>A091557</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091557</u> <u>002</u>	Sep 12, 2011
<u>AB</u>	ANCHEN PHARMS	<u>500MG</u>	<u>A091360</u> <u>001</u>	Oct 04, 2011
<u>AB</u>		<u>750MG</u>	<u>A091360</u> <u>002</u>	Oct 04, 2011
<u>AB</u>	APOTEX INC	<u>500MG</u>	<u>A091261</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091261</u> <u>002</u>	Sep 12, 2011
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A091399</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091399</u> <u>002</u>	Sep 12, 2011
<u>AB</u>	MUTUAL PHARM CO INC	<u>500MG</u>	<u>A091285</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091285</u> <u>002</u>	Sep 12, 2011
<u>AB</u>	MYLAN PHARMS INC	<u>500MG</u>	<u>A200475</u> <u>001</u>	Dec 19, 2011
<u>AB</u>		<u>750MG</u>	<u>A200475</u> <u>002</u>	Dec 19, 2011
<u>AB</u>	PAR PHARM	<u>500MG</u>	<u>A091291</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091291</u> <u>002</u>	Sep 12, 2011
<u>AB</u>	TEVA PHARMS	<u>500MG</u>	<u>A091430</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091430</u> <u>002</u>	Sep 12, 2011
<u>AB</u>	WATSON LABS FLORIDA	<u>500MG</u>	<u>A091093</u> <u>001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091093</u> <u>002</u>	Sep 12, 2011

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKBETA

<u>AT</u>	AKORN	<u>0.25%</u>	<u>A074779</u> <u>001</u>	Oct 29, 1996
<u>AT</u>		<u>0.5%</u>	<u>A074780</u> <u>001</u>	Oct 29, 1996

BETAGAN

<u>AT</u>	+ ALLERGAN	<u>0.25%</u>	<u>N019814</u> <u>001</u>	Jun 28, 1989
<u>AT</u>	+	<u>0.5%</u>	<u>N019219</u> <u>002</u>	Dec 19, 1985

LEVOBUNOLOL HYDROCHLORIDE

<u>AT</u>	BAUSCH AND LOMB	<u>0.25%</u>	<u>A074307</u> <u>001</u>	Mar 04, 1994
<u>AT</u>		<u>0.5%</u>	<u>A074326</u> <u>001</u>	Mar 04, 1994
<u>AT</u>	FALCON PHARMS	<u>0.25%</u>	<u>A074851</u> <u>001</u>	Oct 28, 1996
<u>AT</u>		<u>0.5%</u>	<u>A074850</u> <u>001</u>	Oct 28, 1996
<u>AT</u>	NOVEX	<u>0.25%</u>	<u>A075473</u> <u>001</u>	Aug 03, 2000
<u>AT</u>		<u>0.5%</u>	<u>A075475</u> <u>001</u>	Aug 03, 2000

LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR

<u>AP</u>	+ SIGMA TAU	<u>200MG/ML</u>	<u>N020182</u> <u>001</u>	Dec 16, 1992
-----------	-------------	-----------------	---------------------------	--------------

LEVOCARNITINE

<u>AP</u>	BEDFORD	<u>200MG/ML</u>	<u>A075567</u> <u>001</u>	Mar 29, 2001
<u>AP</u>	LUITPOLD	<u>200MG/ML</u>	<u>A075861</u> <u>001</u>	Jun 22, 2001
<u>AP</u>	TEVA PARENTERAL	<u>200MG/ML</u>	<u>A075881</u> <u>001</u>	Mar 29, 2001

SOLUTION; ORAL

CARNITOR

<u>AA</u>	+ SIGMA TAU	<u>1GM/10ML</u>	<u>N019257</u> <u>001</u>	Apr 10, 1986
-----------	-------------	-----------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 252 (of 424)

LEVOCARNITINE

SOLUTION; ORAL				
	CARNITOR SF			
AA	SIGMA TAU	<u>1GM/10ML</u>	<u>N019257 002</u>	Mar 28, 2007
	LEVOCARNITINE			
AA	HI TECH PHARMA	<u>1GM/10ML</u>	<u>A077399 001</u>	Oct 25, 2007
AA	LYNE	<u>1GM/10ML</u>	<u>A076851 001</u>	Aug 10, 2004
TABLET; ORAL				
	CARNITOR			
AB	+ SIGMA TAU	<u>330MG</u>	<u>N018948 001</u>	Dec 27, 1985
	LEVOCARNITINE			
AB	COREPHARMA	<u>330MG</u>	<u>A076858 001</u>	Sep 20, 2004

LEVOCETIRIZINE DIHYDROCHLORIDE

SOLUTION; ORAL				
	LEVOCETIRIZINE DIHYDROCHLORIDE			
AA	SYNTTHON PHARMS	<u>2.5MG/5ML</u>	<u>A091263 001</u>	Nov 07, 2011
	XYZAL			
AA	+ UCB INC	<u>2.5MG/5ML</u>	<u>N022157 001</u>	Jan 28, 2008
TABLET; ORAL				
	LEVOCETIRIZINE DIHYDROCHLORIDE			
AB	DR REDDYS LABS LTD	<u>5MG</u>	<u>A090392 001</u>	Feb 24, 2011
AB	GLENMARK GENERICS	<u>5MG</u>	<u>A090385 001</u>	Feb 24, 2011
AB	SYNTTHON PHARMS	<u>5MG</u>	<u>A090229 001</u>	Nov 26, 2010
AB	TEVA PHARMS	<u>5MG</u>	<u>A090199 001</u>	Aug 22, 2011
	XYZAL			
AB	+ UCB INC	<u>5MG</u>	<u>N022064 001</u>	May 25, 2007

LEVOFLOXACIN

INJECTABLE; INJECTION				
	LEVAQUIN			
AP	+ JANSSEN PHARMS	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>N020635 001</u>	Dec 20, 1996
AP	+ JANSSEN PHARMS	<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>N020635 004</u>	Dec 20, 1996
	LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER			
AP	+ JANSSEN PHARMS	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>N020635 002</u>	Dec 20, 1996
AP	+ JANSSEN PHARMS	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>N020635 003</u>	Dec 20, 1996
AP	+ JANSSEN PHARMS	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>N020635 005</u>	Dec 20, 1996
	LEVOFLOXACIN			
AP	AKORN	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A091644 001</u>	Jun 20, 2011
AP	AKORN	<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A091644 002</u>	Jun 20, 2011
AP	SAGENT PHARMS	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A200560 001</u>	Jun 20, 2011
AP	SAGENT PHARMS	<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A200560 002</u>	Jun 20, 2011
	LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER			
AP	ACS DOBFAR INFO SA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A090343 001</u>	Jul 07, 2011
AP	ACS DOBFAR INFO SA	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A090343 002</u>	Jul 07, 2011
AP	HIKMA FARMACEUTICA	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A090343 003</u>	Jul 07, 2011
AP	HIKMA FARMACEUTICA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091375 001</u>	Sep 16, 2011
AP	HIKMA FARMACEUTICA	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091375 002</u>	Sep 16, 2011
AP	HIKMA FARMACEUTICA	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091375 003</u>	Sep 16, 2011
SOLUTION; ORAL				
	LEVAQUIN			
AA	+ JANSSEN PHARMS	<u>250MG/10ML</u>	<u>N021721 001</u>	Oct 21, 2004
	LEVOFLOXACIN			
AA	HI TECH PHARMA	<u>250MG/10ML</u>	<u>A091678 001</u>	Jun 20, 2011
SOLUTION/DROPS; OPHTHALMIC				
	LEVOFLOXACIN			
AT	AKORN	<u>0.5%</u>	<u>A090268 001</u>	Dec 20, 2010
AT	APOTEX	<u>0.5%</u>	<u>A078282 001</u>	Dec 20, 2010
AT	HI TECH PHARMA	<u>0.5%</u>	<u>A076826 001</u>	Feb 10, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 253 (of 424)

LEVOFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

LEVOFLOXACIN

<u>AT</u>	NEXUS PHARMS	<u>0.5%</u>	<u>A077700</u> <u>001</u>	Dec 20, 2010
<u>AT</u>	<u>QUIXIN</u>	<u>0.5%</u>	<u>N021199</u> <u>001</u>	Aug 18, 2000
<u>AT</u>	+ SANTEN	<u>1.5%</u>	<u>N021571</u> <u>001</u>	Mar 01, 2004
	IQUIX			
	+ SANTEN			

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
<u>AB</u>	<u>LEVOFLOXACIN</u>			
<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
<u>AB</u>		<u>500MG</u>	<u>A201043</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A201043</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	DR REDDYS LABS INC	<u>250MG</u>	<u>A076710</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076710</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076710</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	GLENMARK GENERICS	<u>250MG</u>	<u>A200250</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A200250</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A200250</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A078424</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A078424</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A078424</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A076276</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076276</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A077097</u> <u>001</u>	Jun 20, 2011
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A077438</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A077438</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A077438</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	TEVA	<u>250MG</u>	<u>A076361</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076361</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076361</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	TORRENT PHARMS	<u>250MG</u>	<u>A090722</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A090722</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A090722</u> <u>003</u>	Jun 20, 2011
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A090367</u> <u>001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A090367</u> <u>002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A090367</u> <u>003</u>	Jun 20, 2011

LEVOLEUCOVORIN CALCIUM

POWDER; IV (INFUSION)

FUSILEV

+ SPECTRUM PHARMS	EQ 50MG BASE/VIAL	<u>N020140</u> <u>001</u>	Mar 07, 2008
-------------------	-------------------	---------------------------	--------------

SOLUTION; IV (INFUSION)

+ SPECTRUM PHARMS	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)	<u>N020140</u> <u>002</u>	Apr 29, 2011
	EQ 250MG BASE/25ML (EQ 10MG BASE/ML)	<u>N020140</u> <u>003</u>	Apr 29, 2011

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ISOCATINE HYDROCHLORIDE W/ LEVONORDEFRIN

<u>AP</u>	NOVOCOL	<u>0.05MG/ML;2%</u>	<u>A084697</u> <u>001</u>
<u>AP</u>	<u>SCANDONEST L</u>		
<u>AP</u>	DEPROCO	<u>0.05MG/ML;2%</u>	<u>A088388</u> <u>001</u>

Oct 10, 1984

PRESCRIPTION DRUG PRODUCT LIST

3 - 254 (of 424)

LEVONORGESTREL

INTRAUTERINE DEVICE; INTRAUTERINE MIRENA				
+ BAYER HLTHCARE	52MG		N021225 001	Dec 06, 2000
TABLET; ORAL <u>LEVONORGESTREL</u>				
<u>AB</u> PERRIGO R AND D	<u>0.75MG</u>		<u>A090740 001</u>	Dec 30, 2010
<u>AB</u> + WATSON LABS	<u>0.75MG</u>		<u>A078665 001</u>	Aug 28, 2009
<u>AB</u>	<u>0.75MG</u>		<u>A078666 001</u>	Jun 24, 2009
PLAN B				
<u>AB</u> + TEVA WOMENS	<u>0.75MG</u>		<u>N021045 002</u>	Aug 24, 2006
PLAN B ONE-STEP				
+ DURAMED	1.5MG		N021998 001	Jul 10, 2009

LEVORPHANOL TARTRATE

TABLET; ORAL LEVORPHANOL TARTRATE				
ROXANE	2MG		A074278 001	Mar 31, 2000

LEVOOTHYROXINE SODIUM**

Refer to Preface Section 1.8 Levothyroxine Sodium for amplifying information

CAPSULE; ORAL TIROSINT				
INST BIOCHIMIQUE	0.013MG		N022121 001	Aug 01, 2007
INSTITUT BIOCHIMIQUE	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
POWDER; INTRAVENOUS LEVOOTHYROXINE SODIUM				
+ APP PHARMS	100MCG/VIAL		N202231 001	Jun 24, 2011
+	200MCG/VIAL		N202231 002	Jun 24, 2011
+	500MCG/VIAL		N202231 003	Jun 24, 2011
TABLET; ORAL LEVO-T				
--> ALARA PHARM	--> AB1,AB2,AB3 0.025MG		N021342 001	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.05MG		N021342 002	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.075MG		N021342 003	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.088MG		N021342 004	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.1MG		N021342 005	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.112MG		N021342 006	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.125MG		N021342 007	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.137MG		N021342 012	Dec 08, 2003
-->	--> AB1,AB2,AB3 0.15MG		N021342 008	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.175MG		N021342 009	Mar 01, 2002
-->	--> AB1,AB2,AB3 0.2MG		N021342 010	Mar 01, 2002
--> +	--> AB1,AB2,AB3 0.3MG		N021342 011	Mar 01, 2002
LEVOOTHYROXINE SODIUM				
--> MERCK KGAA	--> AB2,AB3 0.025MG		A076752 001	Jun 16, 2005
-->	--> AB2,AB3 0.05MG		A076752 002	Jun 16, 2005
-->	--> AB2,AB3 0.075MG		A076752 003	Jun 16, 2005
-->	--> AB2,AB3 0.088MG		A076752 004	Jun 16, 2005
-->	--> AB2,AB3 0.1MG		A076752 005	Jun 16, 2005
-->	--> AB2,AB3 0.112MG		A076752 006	Jun 16, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 255 (of 424)

LEVOOTHYROXINE SODIUM**

Refer to Preface Section 1.8 Levothyroxine Sodium for amplifying information

TABLET; ORAL

LEVOOTHYROXINE SODIUM

-->	MERCK KGAA	--> AB2,AB3 0.125MG	A076752 007	Jun 16, 2005
-->		--> AB2,AB3 0.15MG	A076752 008	Jun 16, 2005
-->		--> AB2,AB3 0.175MG	A076752 009	Jun 16, 2005
-->		--> AB2,AB3 0.2MG	A076752 010	Jun 16, 2005
-->		--> AB2,AB3 0.3MG	A076752 011	Jun 16, 2005
-->	MYLAN	--> AB1,AB2,AB3,AB4 0.025MG	A076187 001	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.05MG	A076187 002	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.075MG	A076187 003	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.088MG	A076187 004	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.1MG	A076187 005	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.112MG	A076187 006	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.125MG	A076187 007	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.137MG	A076187 012	Dec 13, 2006
-->		--> AB1,AB2,AB3,AB4 0.15MG	A076187 008	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.175MG	A076187 009	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.2MG	A076187 010	Jun 05, 2002
-->		--> AB1,AB2,AB3,AB4 0.3MG	A076187 011	Jun 05, 2002
	LEVOXYL			
-->	KING PHARMS	--> AB1,AB3 0.025MG	N021301 001	May 25, 2001
-->		--> AB1,AB3 0.05MG	N021301 002	May 25, 2001
-->		--> AB1,AB3 0.075MG	N021301 003	May 25, 2001
-->		--> AB1,AB3 0.088MG	N021301 004	May 25, 2001
-->		--> AB1,AB3 0.1MG	N021301 005	May 25, 2001
-->		--> AB1,AB3 0.112MG	N021301 006	May 25, 2001
-->		--> 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
	SYNTHROID			
-->	ABBOTT	--> AB1,AB2 0.025MG	N021402 001	Jul 24, 2002
-->		--> AB1,AB2 0.05MG	N021402 002	Jul 24, 2002
-->		--> AB1,AB2 0.075MG	N021402 003	Jul 24, 2002
-->		--> AB1,AB2 0.088MG	N021402 004	Jul 24, 2002
-->		--> AB1,AB2 0.1MG	N021402 005	Jul 24, 2002
-->		--> AB1,AB2 0.112MG	N021402 006	Jul 24, 2002
-->		--> AB1,AB2 0.125MG	N021402 007	Jul 24, 2002
-->		--> AB1,AB2 0.137MG	N021402 008	Jul 24, 2002
-->		--> AB1,AB2 0.15MG	N021402 009	Jul 24, 2002
-->		--> AB1,AB2 0.175MG	N021402 010	Jul 24, 2002
-->		--> AB1,AB2 0.2MG	N021402 012	Jul 24, 2002
--> +		--> AB1,AB2 0.3MG	N021402 011	Jul 24, 2002
	UNITHROID			
-->	STEVENS J	--> AB1,AB2,AB3 0.025MG	N021210 001	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.05MG	N021210 002	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.075MG	N021210 003	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.088MG	N021210 004	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.1MG	N021210 005	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.112MG	N021210 006	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.125MG	N021210 007	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.137MG	N021210 012	Feb 08, 2008
-->		--> AB1,AB2,AB3 0.15MG	N021210 008	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.175MG	N021210 009	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.2MG	N021210 010	Aug 21, 2000
--> +		--> AB1,AB2,AB3 0.3MG	N021210 011	Aug 21, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 256 (of 424)

LEVOOTHYROXINE SODIUM**

Refer to Preface Section 1.8 Levothyroxine Sodium for amplifying information

TABLET; ORAL

LEVOTHROID

<u>AB4</u>	LLOYD	<u>0.025MG</u>	<u>N021116</u> <u>001</u>	Oct 24, 2002
<u>AB4</u>		<u>0.05MG</u>	<u>N021116</u> <u>002</u>	Oct 24, 2002
<u>AB4</u>		<u>0.075MG</u>	<u>N021116</u> <u>003</u>	Oct 24, 2002
<u>AB4</u>		<u>0.088MG</u>	<u>N021116</u> <u>010</u>	Oct 24, 2002
<u>AB4</u>		<u>0.1MG</u>	<u>N021116</u> <u>004</u>	Oct 24, 2002
<u>AB4</u>		<u>0.112MG</u>	<u>N021116</u> <u>011</u>	Oct 24, 2002
<u>AB4</u>		<u>0.125MG</u>	<u>N021116</u> <u>005</u>	Oct 24, 2002
<u>AB4</u>		<u>0.137MG</u>	<u>N021116</u> <u>012</u>	Dec 07, 2004
<u>AB4</u>		<u>0.15MG</u>	<u>N021116</u> <u>006</u>	Oct 24, 2002
<u>AB4</u>		<u>0.175MG</u>	<u>N021116</u> <u>007</u>	Oct 24, 2002
<u>AB4</u>		<u>0.2MG</u>	<u>N021116</u> <u>008</u>	Oct 24, 2002
<u>AB4</u> +		<u>0.3MG</u>	<u>N021116</u> <u>009</u>	Oct 24, 2002

LIDOCAINE

OINTMENT; TOPICAL

LIDOCAINE

<u>AT</u> +	FOUGERA	<u>5%</u>	<u>A080198</u> <u>001</u>	
<u>AT</u>	NOVOCOL INC	<u>5%</u>	<u>A040911</u> <u>001</u>	May 23, 2011
<u>AT</u>	TARO	<u>5%</u>	<u>A086724</u> <u>001</u>	

PATCH; TOPICAL

LIDODERM

+ TEIKOKU PHARMA USA	5%	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>	HOSPIRA	<u>0.5%</u>	<u>A088328</u> <u>001</u>	May 17, 1984
<u>AP</u>		<u>1%</u>	<u>A083158</u> <u>001</u>	
<u>AP</u>		<u>1%</u>	<u>A088329</u> <u>001</u>	May 17, 1984
<u>AP</u>		<u>2%</u>	<u>A040078</u> <u>001</u>	Jun 23, 1995
<u>AP</u>		<u>2%</u>	<u>A083158</u> <u>002</u>	
<u>AP</u>		<u>2%</u>	<u>A088294</u> <u>001</u>	May 17, 1984
<u>AP</u>		<u>20%</u>	<u>A083158</u> <u>003</u>	
<u>AP</u>	LUITPOLD	<u>1%</u>	<u>A080850</u> <u>001</u>	
<u>AP</u>		<u>2%</u>	<u>A083198</u> <u>001</u>	
<u>AP</u>	STRIDES ARCOLAB LTD	<u>0.5%</u>	<u>A091056</u> <u>001</u>	Dec 08, 2010
<u>AP</u>		<u>0.5%</u>	<u>A091058</u> <u>001</u>	Sep 30, 2010
<u>AP</u>		<u>1%</u>	<u>A091056</u> <u>002</u>	Dec 08, 2010
<u>AP</u>		<u>1%</u>	<u>A091058</u> <u>002</u>	Sep 30, 2010

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>200MG/100ML</u>	<u>N019830</u> <u>002</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML</u>	<u>N018461</u> <u>002</u>	

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>400MG/100ML</u>	<u>N019830</u> <u>003</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>400MG/100ML</u>	<u>N018461</u> <u>003</u>	

LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>400MG/100ML</u>	<u>N018388</u> <u>002</u>	
<u>AP</u>		<u>400MG/100ML</u>	<u>N018388</u> <u>003</u>	

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>800MG/100ML</u>	<u>N019830</u> <u>004</u>	Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>800MG/100ML</u>	<u>N018461</u> <u>004</u>	Feb 22, 1982

LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>800MG/100ML</u>	<u>N018388</u> <u>003</u>	Nov 05, 1982
-----------	---------	--------------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 257 (of 424)

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	APP PHARMS	<u>1%</u>	<u>A088586</u> <u>001</u>	Jul 24, 1985
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088325</u> <u>001</u>	Jul 31, 1984
<u>AP</u>		<u>1%</u>	<u>A088299</u> <u>001</u>	Jul 31, 1984
<u>AP</u>		<u>2%</u>	<u>A088327</u> <u>001</u>	Jul 31, 1984

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	ABRAXIS PHARM	<u>2%</u>	<u>N017584</u> <u>001</u>	
<u>AP</u>		<u>4%</u>	<u>N017584</u> <u>002</u>	
<u>AP</u>	APP PHARMS	<u>1%</u>	<u>A080404</u> <u>002</u>	
<u>AP</u>		<u>2%</u>	<u>A080404</u> <u>003</u>	
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A080408</u> <u>001</u>	
<u>AP</u>		<u>1.5%</u>	<u>A080408</u> <u>002</u>	
<u>AP</u>		<u>4%</u>	<u>A088295</u> <u>001</u>	May 17, 1984
<u>AP</u>	INTL MEDICATION	<u>20%</u>	<u>N017702</u> <u>001</u>	
<u>AP</u>	STRIDES ARCOLAB LTD	<u>2%</u>	<u>A090665</u> <u>001</u>	Sep 27, 2010

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A040302</u> <u>001</u>	Sep 28, 1998
<u>AP</u>		<u>2%</u>	<u>A040302</u> <u>002</u>	Sep 28, 1998

LIDOPEN

<u>AP</u>	MERIDIAN MEDCL TECHN	<u>10%</u>	<u>N017549</u> <u>001</u>	
-----------	----------------------	------------	---------------------------	--

XYLOCAINE

<u>AP</u>	+ APP PHARMS	<u>0.5%</u>	<u>N006488</u> <u>008</u>	
<u>AP</u>	+	<u>1%</u>	<u>N006488</u> <u>007</u>	
<u>AP</u>	+	<u>1.5%</u>	<u>N006488</u> <u>010</u>	
<u>AP</u>	+	<u>2%</u>	<u>N006488</u> <u>002</u>	

XYLOCAINE 4% PRESERVATIVE FREE

<u>AP</u>	+ APP PHARMS	<u>4%</u>	<u>N010417</u> <u>001</u>	
-----------	--------------	-----------	---------------------------	--

XYLOCAINE DENTAL

<u>AP</u>	+ DENTSPLY PHARM	<u>2%</u>	<u>N021380</u> <u>001</u>	
-----------	------------------	-----------	---------------------------	--

XYLOCAINE PRESERVATIVE FREE

<u>AP</u>	+ APP PHARMS	<u>1%</u>	<u>N016801</u> <u>005</u>	Jan 19, 1988
<u>AP</u>	+	<u>2%</u>	<u>N016801</u> <u>001</u>	
<u>AP</u>	+	<u>4%</u>	<u>N016801</u> <u>002</u>	
<u>AP</u>	+	<u>10%</u>	<u>N016801</u> <u>003</u>	
<u>AP</u>	+	<u>20%</u>	<u>N016801</u> <u>004</u>	

INJECTABLE; SPINAL

LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%

+ HOSPIRA 5%

A083914 001

JELLY; TOPICAL

ANESTACON

<u>AT</u>	POLYMEDICA	<u>2%</u>	<u>A080429</u> <u>001</u>	
-----------	------------	-----------	---------------------------	--

LIDOCAINE HYDROCHLORIDE

<u>AT</u>	AKORN	<u>2%</u>	<u>A040433</u> <u>001</u>	Feb 12, 2003
<u>AT</u>	HI TECH PHARMA	<u>2%</u>	<u>A040837</u> <u>001</u>	Mar 23, 2011
<u>AT</u>	INTL MEDICATION	<u>2%</u>	<u>A086283</u> <u>001</u>	
<u>AT</u>	TEVA PHARMS	<u>2%</u>	<u>A081318</u> <u>001</u>	Apr 29, 1993

XYLOCAINE

<u>AT</u>	+ OAK PHARMS	<u>2%</u>	<u>N008816</u> <u>001</u>	
-----------	--------------	-----------	---------------------------	--

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE

<u>AT</u>	HI TECH PHARMA	<u>2%</u>	<u>A040014</u> <u>001</u>	Jul 10, 1995
<u>AT</u>	WOCKHARDT	<u>2%</u>	<u>A087872</u> <u>001</u>	Nov 18, 1982

LIDOCAINE HYDROCHLORIDE VISCOSUS

<u>AT</u>	VINTAGE	<u>2%</u>	<u>A040708</u> <u>001</u>	Feb 27, 2007
-----------	---------	-----------	---------------------------	--------------

LIDOCAINE VISCOSUS

<u>AT</u>	ROXANE	<u>2%</u>	<u>A088802</u> <u>001</u>	Apr 26, 1985
-----------	--------	-----------	---------------------------	--------------

XYLOCAINE VISCOSUS

<u>AT</u>	+ APP PHARMS	<u>2%</u>	<u>N009470</u> <u>001</u>	
-----------	--------------	-----------	---------------------------	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 258 (of 424)

LIDOCAINE HYDROCHLORIDE

SOLUTION; TOPICAL

LARYNG-O-JET KIT

<u>AT</u>	INTL MEDICATION	<u>4%</u>	<u>A086364</u>	<u>001</u>	
	<u>LIDOCAINE HYDROCHLORIDE</u>				
<u>AT</u>	ROXANE	<u>4%</u>	<u>A088803</u>	<u>001</u>	Apr 03, 1985
<u>AT</u>	VINTAGE	<u>4%</u>	<u>A040710</u>	<u>001</u>	Feb 27, 2007
<u>AT</u>	WOCKHARDT	<u>4%</u>	<u>A087881</u>	<u>001</u>	Nov 18, 1982
	<u>LTA II KIT</u>				
<u>AT</u>	HOSPIRA	<u>4%</u>	<u>A080409</u>	<u>001</u>	
	<u>XYLOCAINE 4% PRESERVATIVE FREE</u>				
<u>AT</u>	+ APP PHARMS	<u>4%</u>	<u>N010417</u>	<u>002</u>	

LIDOCAINE; PRILOCAINE

CREAM; TOPICAL

EMLA

<u>AB</u>	+ OAK PHARMS	<u>2.5%;2.5%</u>	<u>N019941</u>	<u>001</u>	Dec 30, 1992
	<u>LIDOCAINE AND PRILOCAINE</u>				
<u>AB</u>	HI TECH PHARMA	<u>2.5%;2.5%</u>	<u>A076290</u>	<u>001</u>	Sep 25, 2003
<u>AB</u>	NYCOMED US	<u>2.5%;2.5%</u>	<u>A076453</u>	<u>001</u>	Aug 18, 2003

GEL; PERIODONTAL

ORAQIX

+ DENTSPLY PHARM 2.5%;2.5%

N021451 001 Dec 19, 2003

LIDOCAINE; TETRACAIN

CREAM; TOPICAL

LIDOCAINE AND TETRACAIN

+ GALDERMA LABS LP 7%;7%

N021717 001 Jun 29, 2006

PATCH; TOPICAL

SYNERA

+ ZARS PHARM 70MG;70MG

N021623 001 Jun 23, 2005

LINAGLIPTIN

TABLET; ORAL

TRADJENTA

+ BOEHRINGER INGELHEIM 5MG

N201280 001 May 02, 2011

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCIN

+ PHARMACIA AND UPJOHN EQ 300MG BASE/ML

N050317 001

LINDANE

LOTION; TOPICAL

LINDANE

<u>AT</u>	OLTA PHARMS	<u>1%</u>	<u>A087313</u>	<u>001</u>	
<u>AT</u>	+ WOCKHARDT	<u>1%</u>	<u>A088190</u>	<u>001</u>	Aug 16, 1984

SHAMPOO; TOPICAL

LINDANE

<u>AT</u>	OLTA PHARMS	<u>1%</u>	<u>A087266</u>	<u>001</u>	
<u>AT</u>	+ WOCKHARDT	<u>1%</u>	<u>A088191</u>	<u>001</u>	Sep 18, 1984

LINEZOLID

FOR SUSPENSION; ORAL

ZYVOX

+ PHARMACIA AND UPJOHN 100MG/5ML

N021132 001 Apr 18, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 259 (of 424)

LINEZOLID

INJECTABLE; INJECTION ZYVOX		
+ PHARMACIA AND UPJOHN 200MG/100ML	N021131 001	Apr 18, 2000
TABLET; ORAL ZYVOX		
+ PHARMACIA AND UPJOHN 600MG	N021130 002	Apr 18, 2000

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION <u>LIOTHYRONINE SODIUM</u>		
<u>AP X GEN PHARMS</u>	<u>EQ 0.01MG BASE/ML</u>	<u>A076923 001</u> Aug 17, 2005
<u>AP + JHP PHARMS</u>	<u>EQ 0.01MG BASE/ML</u>	<u>N020105 001</u> Dec 31, 1991
TABLET; ORAL <u>CYTOMEL</u>		
<u>AB KING PHARMS</u>	<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>
<u>LIOTHYRONINE SODIUM</u>		
<u>AB COASTAL PHARMS</u>	<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 MYLAN</u>	<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

LIOTRIX (T4;T3)

TABLET; ORAL THYROLAR-0.25	0.0125MG;0.0031MG	N016807 001
FOREST LABS		
THYROLAR-0.5	0.025MG;0.0063MG	N016807 005
FOREST LABS		
THYROLAR-1	0.05MG;0.0125MG	N016807 004
FOREST LABS		
THYROLAR-2	0.1MG;0.025MG	N016807 002
FOREST LABS		
THYROLAR-3	0.15MG;0.0375MG	N016807 003
+ FOREST LABS		

LIRAGLUTIDE RECOMBINANT

SOLUTION; SUBCUTANEOUS VICTOZA		
+ NOVO NORDISK INC	18MG/3ML (6MG/ML)	N022341 001 Jan 25, 2010

LISDEXAMFETAMINE Dimesylate

CAPSULE; ORAL VYVANSE		
SHIRE DEVELOPMENT	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

LISINOPRIL

TABLET; ORAL <u>LISINOPRIL</u>	<u>2.5MG</u>	<u>A076102 001</u> Sep 30, 2002
<u>AB APOTEX INC</u>		

PRESCRIPTION DRUG PRODUCT LIST

3 - 260 (of 424)

LISINOPRIL

TABLET; ORAL

LISINOPRIL

AB	APOTEX INC	<u>5MG</u>	<u>A076102</u>	<u>002</u>	Sep 30, 2002
AB		<u>10MG</u>	<u>A076102</u>	<u>003</u>	Sep 30, 2002
AB		<u>20MG</u>	<u>A076102</u>	<u>004</u>	Sep 30, 2002
AB		<u>30MG</u>	<u>A076102</u>	<u>005</u>	Sep 30, 2002
AB		<u>40MG</u>	<u>A076102</u>	<u>006</u>	Sep 30, 2002
AB	AUROBINDO	<u>2.5MG</u>	<u>A077622</u>	<u>001</u>	Feb 22, 2006
AB		<u>5MG</u>	<u>A077622</u>	<u>002</u>	Feb 22, 2006
AB		<u>10MG</u>	<u>A077622</u>	<u>003</u>	Feb 22, 2006
AB		<u>20MG</u>	<u>A077622</u>	<u>004</u>	Feb 22, 2006
AB		<u>30MG</u>	<u>A077622</u>	<u>005</u>	Feb 22, 2006
AB		<u>40MG</u>	<u>A077622</u>	<u>006</u>	Feb 22, 2006
AB	IVAX SUB TEVA PHARMS	<u>2.5MG</u>	<u>A075752</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A075752</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075752</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A075752</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A075752</u>	<u>005</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A075752</u>	<u>006</u>	Jul 01, 2002
AB	LEK PHARMS	<u>2.5MG</u>	<u>A075999</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A075999</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075999</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A075999</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A075999</u>	<u>005</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A075999</u>	<u>006</u>	Jul 01, 2002
AB	LUPIN	<u>2.5MG</u>	<u>A077321</u>	<u>001</u>	Sep 09, 2005
AB		<u>5MG</u>	<u>A077321</u>	<u>002</u>	Sep 09, 2005
AB		<u>10MG</u>	<u>A077321</u>	<u>003</u>	Sep 09, 2005
AB		<u>20MG</u>	<u>A077321</u>	<u>004</u>	Sep 09, 2005
AB		<u>30MG</u>	<u>A077321</u>	<u>005</u>	Sep 09, 2005
AB		<u>40MG</u>	<u>A077321</u>	<u>006</u>	Sep 09, 2005
AB	MYLAN	<u>2.5MG</u>	<u>A076071</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A076071</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A076071</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A076071</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A076071</u>	<u>005</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A076071</u>	<u>006</u>	Jul 01, 2002
AB	PRINSTON INC	<u>2.5MG</u>	<u>A076180</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A076180</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A076180</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A076164</u>	<u>001</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A076164</u>	<u>002</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A076164</u>	<u>003</u>	Jul 01, 2002
AB	RANBAXY	<u>2.5MG</u>	<u>A075944</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A075944</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075944</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A075944</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A075944</u>	<u>006</u>	Feb 11, 2003
AB		<u>40MG</u>	<u>A075944</u>	<u>005</u>	Jul 01, 2002
AB	SANDOZ	<u>2.5MG</u>	<u>A075903</u>	<u>001</u>	Jul 01, 2002
AB		<u>2.5MG</u>	<u>A075994</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A075903</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075994</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075903</u>	<u>003</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075994</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A075903</u>	<u>004</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A075994</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A075903</u>	<u>005</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A075994</u>	<u>005</u>	Jul 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 261 (of 424)

LISINOPRIL

TABLET; ORAL

LISINOPRIL

AB	SANDOZ	<u>40MG</u>	<u>A075903</u>	<u>006</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A075994</u>	<u>006</u>	Jul 01, 2002
AB	VINTAGE	<u>2.5MG</u>	<u>A075743</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A075743</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A075743</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A075743</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A075743</u>	<u>005</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A075743</u>	<u>006</u>	Jul 01, 2002
AB	WATSON LABS	<u>2.5MG</u>	<u>A076059</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A076059</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A076059</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A076059</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A076059</u>	<u>005</u>	Jul 01, 2002
AB		<u>40MG</u>	<u>A076059</u>	<u>006</u>	Jul 01, 2002
AB	WEST WARD	<u>2.5MG</u>	<u>A076063</u>	<u>001</u>	Jul 01, 2002
AB		<u>5MG</u>	<u>A076063</u>	<u>002</u>	Jul 01, 2002
AB		<u>10MG</u>	<u>A076063</u>	<u>003</u>	Jul 01, 2002
AB		<u>20MG</u>	<u>A076063</u>	<u>004</u>	Jul 01, 2002
AB		<u>30MG</u>	<u>A076063</u>	<u>006</u>	Jun 27, 2003
AB		<u>40MG</u>	<u>A076063</u>	<u>005</u>	Jul 01, 2002
AB	WOCKHARDT	<u>2.5MG</u>	<u>A078402</u>	<u>001</u>	Apr 19, 2007
AB		<u>5MG</u>	<u>A078402</u>	<u>002</u>	Apr 19, 2007
AB		<u>10MG</u>	<u>A078402</u>	<u>003</u>	Apr 19, 2007
AB		<u>20MG</u>	<u>A078402</u>	<u>004</u>	Apr 19, 2007
AB		<u>30MG</u>	<u>A078402</u>	<u>005</u>	Apr 19, 2007
AB		<u>40MG</u>	<u>A078402</u>	<u>006</u>	Apr 19, 2007

PRINIVIL

AB	MERCK	<u>5MG</u>	<u>N019558</u>	<u>001</u>	Dec 29, 1987
AB		<u>10MG</u>	<u>N019558</u>	<u>002</u>	Dec 29, 1987
AB		<u>20MG</u>	<u>N019558</u>	<u>003</u>	Dec 29, 1987
AB		<u>40MG</u>	<u>N019558</u>	<u>004</u>	Oct 25, 1988

ZESTRIL

AB	ASTRAZENECA	<u>2.5MG</u>	<u>N019777</u>	<u>005</u>	Apr 29, 1993
AB		<u>5MG</u>	<u>N019777</u>	<u>001</u>	May 19, 1988
AB		<u>10MG</u>	<u>N019777</u>	<u>002</u>	May 19, 1988
AB		<u>20MG</u>	<u>N019777</u>	<u>003</u>	May 19, 1988
AB		<u>30MG</u>	<u>N019777</u>	<u>006</u>	Jan 20, 1999
AB	+	<u>40MG</u>	<u>N019777</u>	<u>004</u>	May 19, 1988

LITHIUM CARBONATE

CAPSULE; ORAL

LITHIUM CARBONATE

AB	ALEMBIC LTD	<u>150MG</u>	<u>A079159</u>	<u>001</u>	Jan 12, 2009
AB		<u>300MG</u>	<u>A079159</u>	<u>002</u>	Jan 12, 2009
AB		<u>600MG</u>	<u>A079159</u>	<u>003</u>	Jan 12, 2009
AB	APOTEX INC	<u>300MG</u>	<u>A076795</u>	<u>001</u>	Nov 22, 2004
AB	GLENMARK GENERICS	<u>150MG</u>	<u>A079139</u>	<u>001</u>	Feb 03, 2009
AB		<u>300MG</u>	<u>A079139</u>	<u>002</u>	Feb 03, 2009
AB		<u>600MG</u>	<u>A079139</u>	<u>003</u>	Feb 03, 2009
AB	HETERO DRUGS LTD	<u>150MG</u>	<u>A090702</u>	<u>001</u>	Sep 25, 2009
AB		<u>300MG</u>	<u>A090702</u>	<u>002</u>	Sep 25, 2009
AB		<u>600MG</u>	<u>A090702</u>	<u>003</u>	Sep 25, 2009
AB	ROXANE	<u>150MG</u>	<u>N017812</u>	<u>002</u>	Jan 28, 1987
AB		<u>300MG</u>	<u>N017812</u>	<u>001</u>	
AB	+	<u>600MG</u>	<u>N017812</u>	<u>003</u>	Jan 28, 1987
AB	WEST WARD	<u>150MG</u>	<u>A076243</u>	<u>002</u>	Feb 24, 2003
AB		<u>300MG</u>	<u>A076243</u>	<u>001</u>	Jun 27, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 262 (of 424)

LITHIUM CARBONATE

CAPSULE; ORAL

LITHIUM CARBONATE

<u>AB</u>	WEST WARD	<u>600MG</u>	<u>A078763</u> <u>001</u>	Apr 15, 2008
TABLET; ORAL				
		<u>LITHIUM CARBONATE</u>		
<u>AB</u>	+ ROXANE	<u>300MG</u>	<u>N018558</u> <u>001</u>	Jan 29, 1982
<u>AB</u>	SUN PHARM INDs INC	<u>300MG</u>	<u>A091027</u> <u>001</u>	Jun 24, 2010
<u>AB</u>	WEST WARD	<u>300MG</u>	<u>A078715</u> <u>001</u>	Dec 28, 2010
TABLET, EXTENDED RELEASE; ORAL				
		<u>LITHIUM CARBONATE</u>		
<u>AB</u>	GLENMARK GENERICS	<u>300MG</u>	<u>A091544</u> <u>001</u>	Dec 27, 2010
<u>AB</u>		<u>450MG</u>	<u>A091616</u> <u>001</u>	Feb 14, 2011
<u>AB</u>	ROXANE	<u>300MG</u>	<u>A076832</u> <u>001</u>	Oct 28, 2004
<u>AB</u>	+ WEST WARD	<u>450MG</u>	<u>A076691</u> <u>001</u>	Jan 05, 2004
<u>AB</u>		<u>450MG</u>	<u>A076490</u> <u>001</u>	Jun 17, 2003
		<u>LITHOBID</u>		
<u>AB</u>	+ NOVEN THERAP	<u>300MG</u>	<u>N018027</u> <u>001</u>	

LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

<u>AA</u>	+ ROXANE	<u>EQ 300MG CARBONATE/5ML</u>	<u>N018421</u> <u>001</u>	
<u>AA</u>	WOCKHARDT	<u>EQ 300MG CARBONATE/5ML</u>	<u>A070755</u> <u>001</u>	May 21, 1986

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS; OPHTHALMIC

+ ALCON

EQ 0.1% BASE

N020191 001 Sep 23, 1993

LOMUSTINE

CAPSULE; ORAL

CEENU

BRISTOL MYERS SQUIBB	10MG	N017588 001
	40MG	N017588 002
+	100MG	N017588 003

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

IMODIUM

<u>AB</u>	+ MCNEIL CONS	<u>2MG</u>	<u>N017694</u> <u>001</u>	
<u>LOPERAMIDE HYDROCHLORIDE</u>				
<u>AB</u>	MYLAN	<u>2MG</u>	<u>A072741</u> <u>001</u>	Sep 18, 1991
<u>AB</u>	TEVA	<u>2MG</u>	<u>A073192</u> <u>001</u>	Apr 30, 1992

LOPINAVIR; RITONAVIR

CAPSULE; ORAL

KALETRA

+ ABBOTT

133.3MG;33.3MG

N021226 001 Sep 15, 2000

SOLUTION; ORAL

KALETRA

+ ABBOTT

80MG/ML;20MG/ML

N021251 001 Sep 15, 2000

TABLET; ORAL

KALETRA

ABBOTT

100MG;25MG

N021906 002 Nov 09, 2007

+

200MG;50MG

N021906 001 Oct 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 263 (of 424)

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>	PADDICK LLC	<u>2MG/ML</u>	<u>A079244</u>	<u>001</u>	Apr 28, 2009
<u>AA</u>	PHARM ASSOC	<u>2MG/ML</u>	<u>A090260</u>	<u>001</u>	Jun 15, 2010
	<u>LORAZEPAM INTENSOL</u>				
<u>AA</u>	+ ROXANE	<u>2MG/ML</u>	<u>A072755</u>	<u>001</u>	Jun 28, 1991

INJECTABLE; INJECTION

ATIVAN

<u>AP</u>	+ BAXTER HLTHCARE CORP	<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>	BEDFORD	<u>2MG/ML</u>	<u>A077076</u>	<u>001</u>	Jul 13, 2005
<u>AP</u>		<u>4MG/ML</u>	<u>A077076</u>	<u>002</u>	Jul 13, 2005
<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>2MG/ML</u>	<u>A074300</u>	<u>001</u>	Apr 12, 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
<u>AP</u>	TAYLOR	<u>2MG/ML</u>	<u>A075025</u>	<u>001</u>	Jul 23, 1998
<u>AP</u>	WATSON LABS	<u>2MG/ML</u>	<u>A074276</u>	<u>001</u>	Apr 15, 1994
<u>AP</u>		<u>4MG/ML</u>	<u>A074276</u>	<u>002</u>	Apr 15, 1994

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>	ACTAVIS ELIZABETH	<u>0 .5MG</u>	<u>A071403</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>		<u>1MG</u>	<u>A071404</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>		<u>2MG</u>	<u>A071141</u>	<u>001</u>	Apr 21, 1987
<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>	EXCELLIUM	<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>	IVAX SUB TEVA PHARMS	<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>	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
<u>AB</u>		<u>1MG</u>	<u>A071590</u>	<u>001</u>	Oct 13, 1987
<u>AB</u>		<u>1MG</u>	<u>A077657</u>	<u>002</u>	Mar 16, 2006
<u>AB</u>		<u>2MG</u>	<u>A071591</u>	<u>001</u>	Oct 13, 1987
<u>AB</u>		<u>2MG</u>	<u>A077657</u>	<u>003</u>	Mar 16, 2006
<u>AB</u>	RANBAXY	<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
<u>AB</u>		<u>2MG</u>	<u>A076045</u>	<u>003</u>	Aug 29, 2001
<u>AB</u>	SANDOZ	<u>0 .5MG</u>	<u>A071193</u>	<u>001</u>	Apr 15, 1988
<u>AB</u>		<u>1MG</u>	<u>A071194</u>	<u>001</u>	Apr 15, 1988
<u>AB</u>		<u>2MG</u>	<u>A071195</u>	<u>001</u>	Apr 15, 1988
<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

PRESCRIPTION DRUG PRODUCT LIST

3 - 264 (of 424)

LORAZEPAM

TABLET; ORAL

LORAZEPAM

<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>A071118</u>	<u>001</u>	Jul 24, 1986
<u>AB</u>		<u>1MG</u>	<u>A072927</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>		<u>2MG</u>	<u>A072928</u>	<u>001</u>	Oct 31, 1991

LOSARTAN POTASSIUM

TABLET; ORAL

COZAAR

<u>AB</u>	MERCK	<u>25MG</u>	<u>N020386</u>	<u>001</u>	Apr 14, 1995
<u>AB</u>		<u>50MG</u>	<u>N020386</u>	<u>002</u>	Apr 14, 1995
<u>AB</u>	+	<u>100MG</u>	<u>N020386</u>	<u>003</u>	Oct 13, 1998

LOSARTAN POTASSIUM

<u>AB</u>	ACTAVIS INC	<u>25MG</u>	<u>A090382</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090382</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090382</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090428</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090428</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090428</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	APOTEX CORP	<u>25MG</u>	<u>A090790</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090790</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090790</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A090083</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090083</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090083</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A078232</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078232</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078232</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A091590</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A091590</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A091590</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A091497</u>	<u>001</u>	Jun 06, 2011
<u>AB</u>		<u>50MG</u>	<u>A091497</u>	<u>002</u>	Jun 06, 2011
<u>AB</u>		<u>100MG</u>	<u>A091497</u>	<u>003</u>	Jun 06, 2011
<u>AB</u>	ROXANE	<u>25MG</u>	<u>A077459</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A077459</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A077459</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A077424</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A077424</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A077424</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	TEVA	<u>25MG</u>	<u>A076958</u>	<u>001</u>	Apr 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A076958</u>	<u>002</u>	Apr 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A076958</u>	<u>003</u>	Apr 06, 2010
<u>AB</u>	TORRENT PHARMS	<u>25MG</u>	<u>A090467</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090467</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090467</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	UPSHER SMITH	<u>25MG</u>	<u>A090544</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A090544</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A090544</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A091129</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A091129</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A091129</u>	<u>003</u>	Oct 06, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078243</u>	<u>001</u>	Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078243</u>	<u>002</u>	Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078243</u>	<u>003</u>	Oct 06, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 265 (of 424)

LOTEPREDNOL ETABONATE

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			
+ BAUSCH AND LOMB	0.5%;0.3%	N050804 001	Dec 14, 2004

LOVASTATIN

TABLET; ORAL

LOVASTATIN

AB ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075828 001</u>	Dec 17, 2001
AB	<u>20MG</u>	<u>A075828 002</u>	Dec 17, 2001
AB	<u>40MG</u>	<u>A075828 003</u>	Dec 17, 2001
AB APOTEX INC	<u>10MG</u>	<u>A077748 001</u>	Feb 28, 2007
AB	<u>20MG</u>	<u>A077748 002</u>	Feb 28, 2007
AB	<u>40MG</u>	<u>A077748 003</u>	Feb 28, 2007
AB CARLSBAD	<u>10MG</u>	<u>A075991 001</u>	Jun 05, 2002
AB	<u>20MG</u>	<u>A075991 002</u>	Jun 05, 2002
AB	<u>40MG</u>	<u>A075991 003</u>	Jun 05, 2002
AB LUPIN	<u>10MG</u>	<u>A078296 001</u>	Mar 14, 2008
AB	<u>20MG</u>	<u>A078296 002</u>	Nov 01, 2007
AB	<u>40MG</u>	<u>A078296 003</u>	Nov 01, 2007
AB MUTUAL PHARM	<u>10MG</u>	<u>A077520 001</u>	Apr 14, 2006
AB	<u>20MG</u>	<u>A077520 002</u>	Apr 14, 2006
AB	<u>40MG</u>	<u>A077520 003</u>	Apr 14, 2006
AB MYLAN	<u>10MG</u>	<u>A075451 001</u>	Dec 17, 2001
AB	<u>10MG</u>	<u>A075935 001</u>	Dec 17, 2001
AB	<u>20MG</u>	<u>A075451 002</u>	Dec 17, 2001
AB	<u>20MG</u>	<u>A075935 002</u>	Dec 17, 2001
AB	<u>40MG</u>	<u>A075451 003</u>	Dec 17, 2001
AB	<u>40MG</u>	<u>A075935 003</u>	Dec 17, 2001
AB SANDOZ	<u>10MG</u>	<u>A075300 001</u>	Dec 17, 2001
AB	<u>10MG</u>	<u>A075636 001</u>	Dec 17, 2001
AB	<u>20MG</u>	<u>A075300 002</u>	Dec 17, 2001
AB	<u>20MG</u>	<u>A075636 002</u>	Dec 17, 2001
AB	<u>40MG</u>	<u>A075300 003</u>	Dec 17, 2001
AB	<u>40MG</u>	<u>A075636 003</u>	Dec 17, 2001
AB TEVA	<u>10MG</u>	<u>A075551 003</u>	Dec 17, 2001
AB	<u>20MG</u>	<u>A075551 002</u>	Dec 17, 2001
AB	<u>40MG</u>	<u>A075551 001</u>	Dec 17, 2001
<u>MEVACOR</u>			
AB MERCK	<u>20MG</u>	<u>N019643 003</u>	Aug 31, 1987
AB +	<u>40MG</u>	<u>N019643 004</u>	Dec 14, 1988

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV			
ANDRX LABS LLC	20MG	N021316 002	Jun 26, 2002
	40MG	N021316 003	Jun 26, 2002
+	60MG	N021316 004	Jun 26, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 266 (of 424)

LOVASTATIN; NIACIN

TABLET, EXTENDED RELEASE; ORAL

ADVICOR

+ ABBOTT	20MG;500MG	N021249 001	Dec 17, 2001
+	20MG;750MG	N021249 002	Dec 17, 2001
+	20MG;1GM	N021249 003	Dec 17, 2001
+	40MG;1GM	N021249 004	Apr 27, 2006

LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE SUCCINATE

<u>AB</u>	LANNETT HOLDINGS INC	<u>EQ 5MG BASE</u>	<u>A090695 001</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090695 002</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A090695 003</u>	Sep 26, 2011
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090695 004</u>	Sep 26, 2011
<u>AB</u>	MYLAN	<u>EQ 5MG BASE</u>	<u>A076762 001</u>	Nov 01, 2004
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076762 002</u>	Nov 01, 2004
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A076762 003</u>	Nov 01, 2004
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076762 004</u>	Nov 01, 2004
<u>AB</u>	WATSON LABS	<u>EQ 5MG BASE</u>	<u>A072204 001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A072205 001</u>	Jun 15, 1988
<u>AB</u> +		<u>EQ 25MG BASE</u>	<u>A072206 001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A072062 001</u>	Jun 15, 1988

LUBIPROSTONE

CAPSULE; ORAL

AMITIZA

SUCAMPO PHARMS	8MCG	N021908 002	Apr 29, 2008
+	24MCG	N021908 001	Jan 31, 2006

LURASIDONE HYDROCHLORIDE

TABLET; ORAL

LATUDA

SUNOVION	40MG	N200603 001	Oct 28, 2010
+	80MG	N200603 002	Oct 28, 2010

LUTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

LUVERIS

+ EMD SERONO	75 IU/VIAL	N021322 001	Oct 08, 2004
--------------	------------	-------------	--------------

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYRON

+ UDL LABS	EQ 85MG BASE/GM	N016763 001
------------	-----------------	-------------

FOR SOLUTION; TOPICAL

SULFAMYRON

+ UDL LABS	5%	N019832 003	Jun 05, 1998
------------	----	-------------	--------------

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
-----------------	------------------------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 267 (of 424)

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
0ML

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 30MG/100ML;37MG/100ML;368MG/100ML;526MG
/100ML;502MG/100ML N017378 001

PLASMA-LYTE A IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 30MG/100ML;37MG/100ML;368MG/100ML;526MG
/100ML;502MG/100ML N017378 002 Nov 22, 1982

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN 30MG/100ML;37MG/100ML;370MG/100ML;530MG
/100ML;500MG/100ML

NORMOSOL-R IN PLASTIC CONTAINER

HOSPIRA 30MG/100ML;37MG/100ML;222MG/100ML;526MG
/100ML;502MG/100ML

SOLUTION; IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

B BRAUN 30MG/100ML;37MG/100ML;370MG/100ML;530MG
/100ML;500MG/100ML

PHYSISOL IN PLASTIC CONTAINER

HOSPIRA 30MG/100ML;37MG/100ML;222MG/100ML;526MG
/100ML;502MG/100ML

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 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

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATE

AP + ABRAXIS PHARM 500MG/ML N019316 001 Sep 08, 1986

AP HOSPIRA 500MG/ML A075151 001 Apr 25, 2000

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ HOSPIRA 1GM/100ML N020488 001 Jul 11, 1995

+ 2GM/100ML N020488 002 Jul 11, 1995

MAGNESIUM SULFATE IN PLASTIC CONTAINER

HOSPIRA 2GM/50ML (40MG/ML) N020309 003 Jan 26, 2007

+ 4GM/100ML (40MG/ML) N020309 001 Jun 24, 1994

+ 4GM/50ML (80MG/ML) N020309 002 Jun 24, 1994

MAGNESIUM SULFATE ANHYDROUS; POTASSIUM SULFATE; SODIUM SULFATE

SOLUTION; ORAL

SUPREP BOWEL PREP KIT

+ BRAINTREE LABS 1.6GM/BOT;3.13GM/BOT;17.5GM/BOT N022372 001 Aug 05, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 268 (of 424)

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; 800M G/100ML; 8.75MG/100ML</u>	<u>N018508 001</u>	Feb 19, 1982
<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800M G/100ML; 8.75MG/100ML</u>	<u>N018336 001</u>	

MALATHION

LOTION; TOPICAL

MALATHION

<u>AT</u>	SYNERX PHARMA	<u>0.5%</u>	<u>A078743 001</u>	Mar 06, 2009
<u>AT</u>	+ TARO PHARMS NORTH	<u>0.5%</u>	<u>N018613 001</u>	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%

<u>AP</u>	B BRAUN	<u>10GM/100ML</u>	<u>N016080 002</u>	
<u>AP</u>	<u>MANNITOL 10% IN PLASTIC CONTAINER</u>			
<u>AP</u>	B BRAUN	<u>10GM/100ML</u>	<u>N020006 002</u>	Jul 26, 1993
<u>AP</u>	HOSPIRA	<u>10GM/100ML</u>	<u>N019603 002</u>	Jan 08, 1987

MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER

<u>AP</u>	B BRAUN	<u>10GM/100ML</u>	<u>N016080 006</u>	
<u>AP</u>	<u>MANNITOL 15%</u>		<u>N016080 003</u>	
<u>AP</u>	B BRAUN	<u>15GM/100ML</u>	<u>N020006 003</u>	Jul 26, 1993
<u>AP</u>	<u>MANNITOL 15% IN PLASTIC CONTAINER</u>		<u>N019603 003</u>	Jan 08, 1990

MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%

<u>AP</u>	B BRAUN	<u>15GM/100ML</u>	<u>N016080 005</u>	
<u>AP</u>	<u>MANNITOL 20%</u>		<u>N014738 001</u>	
<u>AP</u>	B BRAUN	<u>20GM/100ML</u>	<u>N016080 004</u>	
<u>AP</u>	<u>MANNITOL 20% IN PLASTIC CONTAINER</u>			

<u>AP</u>	B BRAUN	<u>20GM/100ML</u>	<u>N020006 004</u>	Jul 26, 1993
<u>AP</u>	HOSPIRA	<u>20GM/100ML</u>	<u>N019603 004</u>	Jan 08, 1990
<u>AP</u>	<u>MANNITOL 25%</u>			
<u>AP</u>	APP PHARMS	<u>12.5GM/50ML</u>	<u>A080677 001</u>	

<u>AP</u>	HOSPIRA	<u>12.5GM/50ML</u>	<u>N016269 006</u>	Aug 25, 1994
<u>AP</u>	INTL MEDICATION	<u>12.5GM/50ML</u>	<u>A083051 001</u>	
<u>AP</u>	LUITPOLD	<u>12.5GM/50ML</u>	<u>A087409 001</u>	Jan 21, 1982
<u>AP</u>	<u>MANNITOL 5%</u>			

<u>AP</u>	B BRAUN	<u>5GM/100ML</u>	<u>N016080 001</u>	
<u>AP</u>	<u>MANNITOL 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	B BRAUN	<u>5GM/100ML</u>	<u>N020006 001</u>	Jul 26, 1993
<u>AP</u>	HOSPIRA	<u>5GM/100ML</u>	<u>N019603 001</u>	Jan 08, 1987

<u>AP</u>	<u>MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%</u>			
<u>AP</u>	B BRAUN	<u>5GM/100ML</u>	<u>N016080 007</u>	

<u>AP</u>	<u>OSMITROL 10% IN WATER</u>		<u>N013684 002</u>	
<u>AP</u>	BAXTER HLTHCARE	<u>10GM/100ML</u>		
<u>AP</u>	<u>OSMITROL 10% IN WATER IN PLASTIC CONTAINER</u>		<u>N013684 006</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 269 (of 424)

MANNITOL

INJECTABLE; INJECTION		
<u>OSMITROL 15% IN WATER</u>		
AP BAXTER HLTHCARE <u>15GM/100ML</u>	<u>N013684 004</u>	
<u>OSMITROL 15% IN WATER IN PLASTIC CONTAINER</u>		
AP BAXTER HLTHCARE <u>15GM/100ML</u>	<u>N013684 008</u>	
<u>OSMITROL 20% IN WATER</u>		
AP BAXTER HLTHCARE <u>20GM/100ML</u>	<u>N013684 003</u>	
<u>OSMITROL 20% IN WATER IN PLASTIC CONTAINER</u>		
AP BAXTER HLTHCARE <u>20GM/100ML</u>	<u>N013684 007</u>	
<u>OSMITROL 5% IN WATER</u>		
AP BAXTER HLTHCARE <u>5GM/100ML</u>	<u>N013684 001</u>	
<u>OSMITROL 5% IN WATER IN PLASTIC CONTAINER</u>		
AP BAXTER HLTHCARE <u>5GM/100ML</u>	<u>N013684 005</u>	
POWDER; INHALATION		
ARIDOL KIT		
+ PHARMAXIS N/A, 5MG, 10MG, 20MG, 40MG	N022368 001	Oct 05, 2010
SOLUTION; IRRIGATION		
RESECTISOL IN PLASTIC CONTAINER		
B BRAUN 5GM/100ML	N016772 002	

MANNITOL; SORBITOL

SOLUTION; IRRIGATION		
SORBITOL-MANNITOL IN PLASTIC CONTAINER		
HOSPIRA 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

TABLET; ORAL		
SELZENTRY		
VIVI HLTHCARE 150MG	N022128 001	Aug 06, 2007
+ 300MG	N022128 002	Aug 06, 2007

MEBENDAZOLE

TABLET, CHEWABLE; ORAL		
MEBENDAZOLE		
+ TEVA PHARMS 100MG	A073580 001	Jan 04, 1995

MECASERMIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS		
INCRELEX		
+ TERCICA 40MG/4ML (10MG/ML)	N021839 001	Aug 30, 2005

MECHLORETHAMINE HYDROCHLORIDE

INJECTABLE; INJECTION		
MUSTARGEN		
+ LUNDBECK INC 10MG/VIAL	N006695 001	

MECLIZINE HYDROCHLORIDE

TABLET; ORAL		
<u>ANTIVERT</u>		
AA + PFIZER <u>12.5MG</u>	<u>N010721 006</u>	
AA + <u>25MG</u>	<u>N010721 004</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 270 (of 424)

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

ANTIVERT

<u>AA</u>	+ PFIZER	<u>50MG</u>	<u>N010721</u>	<u>001</u>	Jan 20, 1982
<u>MECLIZINE HYDROCHLORIDE</u>					
<u>AA</u>	AMNEAL PHARMS	<u>12.5MG</u>	<u>A201451</u>	<u>001</u>	Feb 23, 2011
<u>AA</u>		<u>25MG</u>	<u>A201451</u>	<u>002</u>	Feb 23, 2011
<u>AA</u>		<u>50MG</u>	<u>A201451</u>	<u>003</u>	Feb 23, 2011
<u>AA</u>	JUBILANT CADISTA	<u>12.5MG</u>	<u>A040659</u>	<u>001</u>	Jun 04, 2010
<u>AA</u>		<u>25MG</u>	<u>A040659</u>	<u>002</u>	Jun 04, 2010
<u>AA</u>	PAR PHARM	<u>12.5MG</u>	<u>A087127</u>	<u>001</u>	
<u>AA</u>		<u>25MG</u>	<u>A087128</u>	<u>001</u>	
<u>AA</u>		<u>50MG</u>	<u>A089674</u>	<u>001</u>	Mar 31, 1988
<u>AA</u>	SANDOZ	<u>12.5MG</u>	<u>A084843</u>	<u>002</u>	May 22, 1989
<u>AA</u>		<u>25MG</u>	<u>A084092</u>	<u>003</u>	May 22, 1989
<u>AA</u>	VINTAGE PHARMS	<u>12.5MG</u>	<u>A040179</u>	<u>001</u>	Jan 30, 1997
<u>AA</u>		<u>25MG</u>	<u>A040179</u>	<u>002</u>	Jan 30, 1997
<u>AA</u>	WATSON LABS	<u>25MG</u>	<u>A085740</u>	<u>001</u>	

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLOFENAMATE SODIUM

<u>AB</u>	MYLAN	<u>EQ 50MG BASE</u>	<u>A071081</u>	<u>002</u>	Sep 03, 1986
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>A071081</u>	<u>001</u>	Sep 03, 1986
<u>AB</u>	WATSON LABS	<u>EQ 50MG BASE</u>	<u>A071468</u>	<u>001</u>	Apr 15, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A071469</u>	<u>001</u>	Apr 15, 1987

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>150MG/ML</u>	<u>N020246</u>	<u>001</u>	Oct 29, 1992
<u>MEDROXYPROGESTERONE ACETATE</u>					
<u>AB</u>	SANDOZ	<u>150MG/ML</u>	<u>A078711</u>	<u>001</u>	May 20, 2009
<u>AB</u>	TEVA PARENTERAL	<u>150MG/ML</u>	<u>A076552</u>	<u>001</u>	Oct 27, 2004
<u>AB</u>		<u>150MG/ML</u>	<u>A076553</u>	<u>001</u>	Jul 28, 2004

DEPO-PROVERA

+ PHARMACIA AND UPJOHN 400MG/ML

N012541 003

INJECTABLE; SUBCUTANEOUS

DEPO-SUBQ PROVERA 104

+ PHARMACIA AND UPJOHN 104MG/0.65ML

N021583 001 Dec 17, 2004

TABLET; ORAL

MEDROXYPROGESTERONE ACETATE

<u>AB</u>	BARR	<u>2.5MG</u>	<u>A040159</u>	<u>001</u>	Aug 09, 1996
<u>AB</u>		<u>5MG</u>	<u>A040159</u>	<u>002</u>	Aug 09, 1996
<u>AB</u>		<u>10MG</u>	<u>A040159</u>	<u>003</u>	Aug 09, 1996
<u>PROVERA</u>					
<u>AB</u>	PHARMACIA AND UPJOHN	<u>2.5MG</u>	<u>N011839</u>	<u>001</u>	
<u>AB</u>		<u>5MG</u>	<u>N011839</u>	<u>003</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N011839</u>	<u>004</u>	
MEDROXYPROGESTERONE ACETATE					
<u>BP</u>	USL PHARMA	<u>10MG</u>	<u>A088484</u>	<u>001</u>	Jul 26, 1984

MEFENAMIC ACID

CAPSULE; ORAL

MEFENAMIC ACID

<u>AB</u>	LUPIN LTD	<u>250MG</u>	<u>A091322</u>	<u>001</u>	Jul 22, 2011
<u>AB</u>	MICRO LABS LTD	<u>250MG</u>	<u>A090562</u>	<u>001</u>	Nov 19, 2010
<u>PONSTEL</u>					
<u>AB</u>	+ SHIONOGI INC	<u>250MG</u>	<u>N015034</u>	<u>003</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 271 (of 424)

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>250MG</u>	<u>A076392</u>	<u>001</u>	Dec 29, 2003
<u>AB</u>	ROXANE	<u>250MG</u>	<u>A076523</u>	<u>001</u>	Oct 01, 2004
<u>AB</u> +	SANDOZ	<u>250MG</u>	<u>A076175</u>	<u>001</u>	Feb 20, 2002
<u>AB</u>	WEST WARD	<u>250MG</u>	<u>A077699</u>	<u>001</u>	Apr 21, 2010

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE

<u>AB</u> +	BRISTOL MYERS SQUIBB	<u>40MG/ML</u>	<u>N020264</u>	<u>001</u>	Sep 10, 1993
		<u>MEGESTROL ACETATE</u>			
<u>AB</u>	APOTEX INC	<u>40MG/ML</u>	<u>A077404</u>	<u>001</u>	Feb 16, 2006
<u>AB</u>	PAR PHARM	<u>40MG/ML</u>	<u>A075671</u>	<u>001</u>	Jul 25, 2001
<u>AB</u>	ROXANE	<u>40MG/ML</u>	<u>A075997</u>	<u>001</u>	Feb 15, 2002
<u>AB</u>	TEVA PHARMS	<u>40MG/ML</u>	<u>A075681</u>	<u>001</u>	May 05, 2003
<u>AB</u>	WOCKHARDT	<u>40MG/ML</u>	<u>A076721</u>	<u>001</u>	Nov 01, 2004
	MEGACE ES				
+ PAR PHARM		125MG/ML	N021778	001	Jul 05, 2005

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>	ROXANE	<u>20MG</u>	<u>A074458</u>	<u>001</u>	Sep 29, 1995
<u>AB</u>		<u>40MG</u>	<u>A074458</u>	<u>002</u>	Sep 29, 1995

MELOXICAM

SUSPENSION; ORAL

MOBIC

+ BOEHRINGER INGELHEIM	7.5MG/5ML	N021530	001	Jun 01, 2004
------------------------	-----------	---------	-----	--------------

TABLET; ORAL

MELOXICAM

<u>AB</u>	APOTEX INC	<u>7.5MG</u>	<u>A077882</u>	<u>001</u>	Jul 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077882</u>	<u>002</u>	Jul 20, 2006
<u>AB</u>	AUROBINDO PHARMA	<u>7.5MG</u>	<u>A078008</u>	<u>001</u>	Oct 02, 2006
<u>AB</u>		<u>15MG</u>	<u>A078008</u>	<u>002</u>	Oct 02, 2006
<u>AB</u>	BEIJING DOUBLE CRANE	<u>7.5MG</u>	<u>A078039</u>	<u>001</u>	Dec 14, 2006
<u>AB</u>		<u>15MG</u>	<u>A078039</u>	<u>002</u>	Dec 14, 2006
<u>AB</u>	BRECKENRIDGE PHARM	<u>7.5MG</u>	<u>A077920</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077920</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	CARACO	<u>7.5MG</u>	<u>A077937</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077937</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	CARLSBAD	<u>7.5MG</u>	<u>A077918</u>	<u>001</u>	Dec 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A077918</u>	<u>002</u>	Dec 07, 2006
<u>AB</u>	COREPHARMA	<u>7.5MG</u>	<u>A077930</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077930</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	DR REDDYS LABS INC	<u>7.5MG</u>	<u>A077931</u>	<u>001</u>	Jul 25, 2006
<u>AB</u>		<u>15MG</u>	<u>A077931</u>	<u>002</u>	Jul 25, 2006
<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A077932</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077932</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	LUPIN PHARMS	<u>7.5MG</u>	<u>A077944</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077944</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	MYLAN	<u>7.5MG</u>	<u>A077923</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077934</u>	<u>001</u>	Jul 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077923</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077934</u>	<u>002</u>	Jul 20, 2006
<u>AB</u>	PURACAP PHARM	<u>7.5MG</u>	<u>A077938</u>	<u>001</u>	Jul 19, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 272 (of 424)

MELOXICAM

TABLET; ORAL

MELOXICAM

<u>AB</u>	PURACAP PHARM	<u>15MG</u>	<u>A077938</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	STRIDES ARCOLAB LTD	<u>7.5MG</u>	<u>A077928</u>	<u>001</u>	May 13, 2009
<u>AB</u>		<u>15MG</u>	<u>A077928</u>	<u>002</u>	May 13, 2009
<u>AB</u>	TARO	<u>7.5MG</u>	<u>A078102</u>	<u>001</u>	Nov 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A078102</u>	<u>002</u>	Nov 07, 2006
<u>AB</u>	TEVA PHARMS	<u>7.5MG</u>	<u>A077936</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077936</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	UNICHEM	<u>7.5MG</u>	<u>A077927</u>	<u>001</u>	Dec 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077927</u>	<u>002</u>	Dec 20, 2006
<u>AB</u>	WATSON LABS	<u>7.5MG</u>	<u>A077929</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077929</u>	<u>002</u>	Jul 19, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>7.5MG</u>	<u>A077921</u>	<u>001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077921</u>	<u>002</u>	Jul 19, 2006
	<u>MOBIC</u>				
<u>AB</u>	BOEHRINGER INGELHEIM	<u>7.5MG</u>	<u>N020938</u>	<u>001</u>	Apr 13, 2000
<u>AB</u>	+	<u>15MG</u>	<u>N020938</u>	<u>002</u>	Aug 23, 2000

MELPHALAN

TABLET; ORAL

ALKERAN

+ GLAXOSMITHKLINE 2MG

N014691 002

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

<u>AP</u>	+	GLAXOSMITHKLINE	<u>EQ 50MG BASE/VIAL</u>	<u>N020207</u>	<u>001</u>	Nov 18, 1992
		<u>MELPHALAN HYDROCHLORIDE</u>				
<u>AP</u>		BEDFORD LABS	<u>EQ 50MG BASE/VIAL</u>	<u>A090303</u>	<u>001</u>	Oct 28, 2010
<u>AP</u>		BIONICHE PHARMA USA	<u>EQ 50MG BASE/VIAL</u>	<u>A090299</u>	<u>001</u>	Oct 27, 2009
<u>AP</u>		SYNERX	<u>EQ 50MG BASE/VIAL</u>	<u>A090270</u>	<u>001</u>	Jun 09, 2009

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NAMENDA XR

FOREST LABS	7MG	N022525 001	Jun 21, 2010
	14MG	N022525 002	Jun 21, 2010
	21MG	N022525 003	Jun 21, 2010
+	28MG	N022525 004	Jun 21, 2010

SOLUTION; ORAL

NAMENDA

+ FOREST LABS 2MG/ML N021627 001 Apr 18, 2005

TABLET; ORAL

NAMENDA

FOREST LABS	5MG	N021487 001	Oct 16, 2003
+	10MG	N021487 002	Oct 16, 2003

MENOTROPINS (FSH:LH)

INJECTABLE; IM-SC

REPRONEX

+ FERRING 75 IU/VIAL;75 IU/VIAL N021047 001 Aug 27, 1999

INJECTABLE; SUBCUTANEOUS

MENOPUR

+ FERRING 75 IU/VIAL;75 IU/VIAL N021663 001 Oct 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 273 (of 424)

MEPENZOLATE BROMIDE

TABLET; ORAL
 CANTIL
 + SANOFI AVENTIS US 25MG N010679 003

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

<u>AP</u>	+ HOSPIRA	<u>25MG/ML</u>	<u>N021171 001</u>
<u>AP</u>	+	<u>50MG/ML</u>	<u>N021171 002</u>
<u>AP</u>	+	<u>75MG/ML</u>	<u>N021171 003</u>
<u>AP</u>	+	<u>100MG/ML</u>	<u>N021171 004</u>

MEPERIDINE HYDROCHLORIDE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>25MG/ML</u>	<u>A080455 007</u>
<u>AP</u>		<u>50MG/ML</u>	<u>A080455 008</u>
<u>AP</u>		<u>75MG/ML</u>	<u>A080455 009</u>
<u>AP</u>		<u>100MG/ML</u>	<u>A080455 010</u>
<u>AP</u>	ELKINS SINK	<u>25MG/ML</u>	<u>A080445 001</u>
<u>AP</u>		<u>50MG/ML</u>	<u>A080445 002</u>
<u>AP</u>		<u>75MG/ML</u>	<u>A080445 003</u>
<u>AP</u>		<u>100MG/ML</u>	<u>A080445 004</u>
<u>AP</u>	WATSON LABS	<u>50MG/ML</u>	<u>A073444 001</u>
<u>AP</u>		<u>100MG/ML</u>	<u>A073445 001</u>

Mar 17, 1992
Mar 17, 1992

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>10MG/ML</u>	<u>A081002 001</u>
<u>AP</u>	+ HOSPIRA	<u>10MG/ML</u>	<u>A088432 001</u>
<u>AP</u>	INTL MEDICATION	<u>10MG/ML</u>	<u>A081309 001</u>
<u>AP</u>	WATSON LABS	<u>10MG/ML</u>	<u>A073443 001</u>

Jul 30, 1993
Aug 16, 1984
Aug 30, 1993
Mar 17, 1992

SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

+ ROXANE 50MG/5ML A088744 001 Jan 30, 1985

TABLET; ORAL

DEMEROL

<u>AA</u>	+ SANOFI AVENTIS US	<u>50MG</u>	<u>N005010 001</u>
<u>AA</u>	+	<u>100MG</u>	<u>N005010 004</u>

MEPERIDINE HYDROCHLORIDE

<u>AA</u>	BARR	<u>50MG</u>	<u>A088639 001</u>
<u>AA</u>		<u>100MG</u>	<u>A088640 001</u>
<u>AA</u>	CARACO	<u>50MG</u>	<u>A040446 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040446 002</u>
<u>AA</u>	EPIC PHARMA	<u>50MG</u>	<u>A040331 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040331 002</u>
<u>AA</u>	MALLINCKRODT	<u>50MG</u>	<u>A040352 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040352 002</u>
<u>AA</u>	MIKART	<u>50MG</u>	<u>A040893 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040893 003</u>
<u>AA</u>	ROXANE	<u>50MG</u>	<u>A040110 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040110 002</u>
<u>AA</u>	VINTAGE PHARMS	<u>50MG</u>	<u>A040191 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040191 002</u>
<u>AA</u>	WATSON LABS	<u>50MG</u>	<u>A040186 001</u>
<u>AA</u>		<u>100MG</u>	<u>A040186 002</u>

MEPERIDINE HYDROCHLORIDE

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>
-----------	-----------	-----------	--------------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 274 (of 424)

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

<u>AP</u> + HOSPIRA	<u>1.5%</u>	<u>N012250</u> <u>005</u>	
<u>AP</u> +	<u>2%</u>	<u>N012250</u> <u>002</u>	
<u>AP</u> + NOVOCOL	<u>3%</u>	<u>A080925</u> <u>001</u>	
<u>MEPIVACAINE HYDROCHLORIDE</u>			
<u>AP</u> HOSPIRA INC	<u>3%</u>	<u>A040806</u> <u>001</u>	Apr 28, 2008
<u>AP</u> WATSON LABS	<u>1%</u>	<u>A088769</u> <u>001</u>	Nov 20, 1984
<u>AP</u>	<u>2%</u>	<u>A088770</u> <u>001</u>	Nov 20, 1984
<u>POLOCAINE</u>			
<u>AP</u> APP PHARMS	<u>1%</u>	<u>A089407</u> <u>001</u>	Dec 01, 1986
<u>AP</u>	<u>2%</u>	<u>A089410</u> <u>001</u>	Dec 01, 1986
<u>AP</u> DENTSPLY PHARM	<u>3%</u>	<u>A088653</u> <u>001</u>	Aug 21, 1984
<u>POLOCAINE-MPF</u>			
<u>AP</u> APP PHARMS	<u>1%</u>	<u>A089406</u> <u>001</u>	Dec 01, 1986
<u>AP</u>	<u>1.5%</u>	<u>A089408</u> <u>001</u>	Dec 01, 1986
<u>AP</u>	<u>2%</u>	<u>A089409</u> <u>001</u>	Dec 01, 1986
<u>SCANDONEST PLAIN</u>			
<u>AP</u> + DEPROCO	<u>3%</u>	<u>A088387</u> <u>001</u>	Oct 10, 1984

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

<u>AA</u> ALEMBIC PHARMS LTD	<u>200MG</u>	<u>A090122</u> <u>001</u>	Feb 18, 2009
<u>AA</u>	<u>400MG</u>	<u>A090122</u> <u>002</u>	Feb 18, 2009
<u>AA</u> INVAGEN PHARMS	<u>200MG</u>	<u>A040797</u> <u>001</u>	Feb 27, 2008
<u>AA</u>	<u>400MG</u>	<u>A040797</u> <u>002</u>	Feb 27, 2008
<u>AA</u> TARO	<u>200MG</u>	<u>A200998</u> <u>001</u>	May 23, 2011
<u>AA</u>	<u>400MG</u>	<u>A200998</u> <u>002</u>	May 23, 2011
<u>AA</u> + WATSON LABS	<u>200MG</u>	<u>A083304</u> <u>001</u>	
<u>AA</u> +	<u>400MG</u>	<u>A083308</u> <u>001</u>	

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL

SOLAGE

+ AQUA PHARMS	2%;0.01%	<u>N020922</u> <u>001</u>	Dec 10, 1999
---------------	----------	---------------------------	--------------

MERCAPTOPURINE

TABLET; ORAL

MERCAPTOPURINE

<u>AB</u> MYLAN	<u>50MG</u>	<u>A040594</u> <u>001</u>	Jul 01, 2005
<u>AB</u> PROMETHEUS LABS	<u>50MG</u>	<u>A040461</u> <u>001</u>	Feb 11, 2004
<u>AB</u> ROXANE	<u>50MG</u>	<u>A040528</u> <u>001</u>	Feb 13, 2004
<u>PURINETHOL</u>			
<u>AB</u> + TEVA	<u>50MG</u>	<u>N009053</u> <u>002</u>	

MEROPENEM

INJECTABLE; INJECTION

MEROPENEM

<u>AP</u> ACS DOBFAR	<u>500MG/VIAL</u>	<u>A091404</u> <u>001</u>	Oct 26, 2011
<u>AP</u>	<u>1GM/VIAL</u>	<u>A091404</u> <u>002</u>	Oct 26, 2011
<u>AP</u> HOSPIRA INC	<u>500MG/VIAL</u>	<u>A090940</u> <u>001</u>	Jun 22, 2010
<u>AP</u>	<u>1GM/VIAL</u>	<u>A090940</u> <u>002</u>	Jun 22, 2010
<u>AP</u> SANDOZ	<u>500MG/VIAL</u>	<u>A091201</u> <u>001</u>	Mar 29, 2011
<u>AP</u>	<u>1GM/VIAL</u>	<u>A091201</u> <u>002</u>	Mar 29, 2011
<u>MERREM</u>			
<u>AP</u> + ASTRAZENECA	<u>500MG/VIAL</u>	<u>N050706</u> <u>003</u>	Jun 21, 1996
<u>AP</u> +	<u>1GM/VIAL</u>	<u>N050706</u> <u>001</u>	Jun 21, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 275 (of 424)

MESALAMINE

CAPSULE, EXTENDED RELEASE; ORAL APRISO				
+ SALIX PHARMS	375MG	N022301 001	Oct 31, 2008	
PENTASA				
SHIRE	250MG	N020049 001	May 10, 1993	
+	500MG	N020049 002	Jul 08, 2004	
ENEMA; RECTAL <u>MESALAMINE</u>				
<u>AB</u> PERRIGO ISRAEL	<u>4GM/60ML</u>	<u>A076751 001</u>	Sep 17, 2004	
<u>AB</u> TEVA	<u>4GM/60ML</u>	<u>A076841 001</u>	Sep 30, 2004	
<u>ROWASA</u>				
<u>AB</u> + MEDA PHARMS	<u>4GM/60ML</u>	<u>N019618 001</u>	Dec 24, 1987	
<u>SFROWASA</u>				
<u>AB</u> MEDA PHARMS	<u>4GM/60ML</u>	<u>N019618 002</u>	Jun 20, 2008	
SUPPOSITORY; RECTAL CANASA				
+ AXCAN	1GM	N021252 002	Nov 05, 2004	
TABLET, DELAYED RELEASE; ORAL ASACOL				
+ WARNER CHILCOTT LLC	400MG	N019651 001	Jan 31, 1992	
ASACOL HD				
+ WARNER CHILCOTT LLC	800MG	N021830 001	May 29, 2008	
LIALDA				
+ SHIRE	1.2GM	N022000 001	Jan 16, 2007	

MESNA

INJECTABLE; INTRAVENOUS <u>MESNA</u>				
<u>AP</u> APP PHARMS	<u>100MG/ML</u>	<u>A075811 001</u>	Apr 26, 2001	
<u>AP</u> BEDFORD	<u>100MG/ML</u>	<u>A075739 001</u>	Jan 09, 2004	
<u>AP</u> SAGENT STRIDES	<u>100MG/ML</u>	<u>A090913 001</u>	Apr 13, 2010	
<u>AP</u> TEVA PARENTERAL	<u>100MG/ML</u>	<u>A075764 001</u>	Apr 27, 2001	
<u>MESNEX</u>				
<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			
+ WATSON LABS	0.05MG;1MG	N016659 001	

METAPROTERENOL SULFATE

SOLUTION; INHALATION <u>METAPROTERENOL SULFATE</u>				
<u>AN</u> + DEY	<u>0.4%</u>	<u>A071786 001</u>	Aug 05, 1988	
<u>AN</u> +	<u>0.6%</u>	<u>A070804 001</u>	Aug 17, 1987	
<u>AN</u> NOVEX	<u>0.4%</u>	<u>A075402 001</u>	Feb 28, 2001	
<u>AN</u>	<u>0.6%</u>	<u>A075403 001</u>	Feb 28, 2001	
<u>AN</u> WOCKHARDT	<u>0.4%</u>	<u>A075586 001</u>	May 30, 2002	
<u>AN</u>	<u>0.6%</u>	<u>A075586 002</u>	May 30, 2002	
SYRUP; ORAL <u>METAPROTERENOL SULFATE</u>				
<u>AA</u> NOVEX	<u>10MG/5ML</u>	<u>A075235 001</u>	Jan 27, 2000	
<u>AA</u> + SILARX	<u>10MG/5ML</u>	<u>A073632 001</u>	Jul 22, 1992	

PRESCRIPTION DRUG PRODUCT LIST

3 - 276 (of 424)

METAPROTERENOL SULFATE

TABLET; ORAL

METAPROTERENOL SULFATE

<u>AB</u>	PAR PHARM	<u>10MG</u>	<u>A072024</u>	<u>001</u>	Jun 28, 1988
<u>AB</u>	+	<u>20MG</u>	<u>A072025</u>	<u>001</u>	Jun 28, 1988
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A073013</u>	<u>001</u>	Jan 31, 1991
<u>AB</u>		<u>20MG</u>	<u>A072795</u>	<u>001</u>	Jan 31, 1991

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

METARAMINOL BITARTRATE

+ APP PHARMS EQ 10MG BASE/ML

A080722 001

METAXALONE

TABLET; ORAL

METAXALONE

<u>AB</u>	SANDOZ	<u>800MG</u>	<u>A040445</u>	<u>001</u>	Mar 31, 2010
<u>AB</u>	<u>SKELAXIN</u>	<u>800MG</u>	<u>N013217</u>	<u>003</u>	Aug 30, 2002

METFORMIN HYDROCHLORIDE

SOLUTION; ORAL

RIOMET

+ RANBAXY 500MG/5ML

N021591 001 Sep 11, 2003

TABLET; ORAL

GLUCOPHAGE

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N020357</u>	<u>001</u>	Mar 03, 1995
<u>AB</u>		<u>850MG</u>	<u>N020357</u>	<u>002</u>	Mar 03, 1995
<u>AB</u>	+	<u>1GM</u>	<u>N020357</u>	<u>005</u>	Nov 05, 1998

METFORMIN HYDROCHLORIDE

<u>AB</u>	ALKEM	<u>500MG</u>	<u>A091184</u>	<u>001</u>	Nov 01, 2010
<u>AB</u>		<u>850MG</u>	<u>A091184</u>	<u>002</u>	Nov 01, 2010
<u>AB</u>		<u>1GM</u>	<u>A091184</u>	<u>003</u>	Nov 01, 2010
<u>AB</u>	ALPHAPHARM	<u>500MG</u>	<u>A075969</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>850MG</u>	<u>A075969</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>		<u>1GM</u>	<u>A075969</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>	ALVOGEN	<u>500MG</u>	<u>A076033</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A076033</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A076033</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A077880</u>	<u>001</u>	Jun 05, 2006
<u>AB</u>		<u>850MG</u>	<u>A077880</u>	<u>002</u>	Jun 05, 2006
<u>AB</u>		<u>1GM</u>	<u>A077880</u>	<u>003</u>	Jun 05, 2006
<u>AB</u>	APOTEX	<u>500MG</u>	<u>A075984</u>	<u>001</u>	Apr 23, 2002
<u>AB</u>		<u>500MG</u>	<u>A090666</u>	<u>001</u>	Dec 07, 2011
<u>AB</u>		<u>850MG</u>	<u>A075984</u>	<u>002</u>	Apr 23, 2002
<u>AB</u>		<u>850MG</u>	<u>A090666</u>	<u>002</u>	Dec 07, 2011
<u>AB</u>		<u>1GM</u>	<u>A075984</u>	<u>003</u>	Apr 23, 2002
<u>AB</u>		<u>1GM</u>	<u>A090666</u>	<u>003</u>	Dec 07, 2011
<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>A077095</u>	<u>001</u>	Jan 14, 2005
<u>AB</u>		<u>850MG</u>	<u>A077095</u>	<u>002</u>	Jan 14, 2005
<u>AB</u>		<u>1GM</u>	<u>A077095</u>	<u>003</u>	Jan 14, 2005
<u>AB</u>	CARACO	<u>500MG</u>	<u>A075967</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>850MG</u>	<u>A075967</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>		<u>1GM</u>	<u>A075967</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>	DR REDDYS LABS INC	<u>500MG</u>	<u>A077787</u>	<u>001</u>	Aug 23, 2006
<u>AB</u>		<u>850MG</u>	<u>A077787</u>	<u>002</u>	Aug 23, 2006
<u>AB</u>		<u>1GM</u>	<u>A077787</u>	<u>003</u>	Aug 23, 2006
<u>AB</u>	GLENMARK GENERICS	<u>500MG</u>	<u>A078170</u>	<u>001</u>	May 23, 2008
<u>AB</u>		<u>850MG</u>	<u>A078170</u>	<u>002</u>	May 23, 2008
<u>AB</u>		<u>1GM</u>	<u>A078170</u>	<u>003</u>	May 23, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 277 (of 424)

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

AB	GRANULES INDIA	500MG	A090564 001	Apr 22, 2010
AB		850MG	A090564 002	Apr 22, 2010
AB		1GM	A090564 003	Apr 22, 2010
AB	INDICUS PHARMA	500MG	A079148 001	Nov 25, 2008
AB		850MG	A079148 002	Nov 25, 2008
AB		1GM	A079148 003	Nov 25, 2008
AB	IVAX SUB TEVA PHARMS	500MG	A075972 001	Jan 24, 2002
AB		850MG	A075972 002	Jan 24, 2002
AB		1GM	A075972 003	Jan 24, 2002
AB	MUTUAL PHARMA	500MG	A076038 001	Feb 21, 2002
AB		850MG	A076038 002	Feb 21, 2002
AB		1GM	A076038 003	Feb 21, 2002
AB	MYLAN	500MG	A075973 001	Jan 25, 2002
AB		500MG	A075976 001	Jan 24, 2002
AB		850MG	A075973 002	Jan 25, 2002
AB		850MG	A075976 002	Jan 24, 2002
AB		1GM	A075973 003	Jan 25, 2002
AB		1GM	A075976 003	Jan 24, 2002
AB	PROVIDENT PHARM	500MG	A077853 001	Jul 28, 2006
AB		850MG	A077853 002	Jul 28, 2006
AB		1GM	A077853 003	Jul 28, 2006
AB	SANDOZ	500MG	A075965 001	Jan 25, 2002
AB		500MG	A075985 001	Jan 25, 2002
AB		850MG	A075965 002	Jan 25, 2002
AB		850MG	A075985 002	Jan 25, 2002
AB		1GM	A075965 003	Jan 25, 2002
AB		1GM	A075985 003	Jan 25, 2002
AB	TEVA	500MG	A075978 001	Jan 25, 2002
AB		850MG	A075978 002	Jan 25, 2002
AB		1GM	A075978 003	Nov 05, 2002
AB	TORRENT PHARMS	500MG	A077711 001	Jan 24, 2007
AB		850MG	A077711 002	Jan 24, 2007
AB		1GM	A077711 003	Jan 24, 2007
AB	WATSON LABS	500MG	A075979 001	Jan 24, 2002
AB		850MG	A075979 002	Jan 24, 2002
AB		1GM	A075979 003	Jan 24, 2002
AB	WATSON LABS FLORIDA	500MG	A075961 001	Jan 25, 2002
AB		850MG	A075961 002	Jan 25, 2002
AB		1GM	A075961 003	Jan 25, 2002
AB	ZYDUS PHARMS USA	500MG	A077064 001	Apr 18, 2005
AB		850MG	A077064 002	Apr 18, 2005
AB		1GM	A077064 003	Apr 18, 2005
	METFORMIN HYDROCHLORIDE			
	IVAX SUB TEVA PHARMS	625MG	A075972 005	Jan 24, 2002
		750MG	A075972 004	Jan 24, 2002

TABLET, EXTENDED RELEASE; ORAL

FORTAMET

AB	+ ANDRX LABS LLC	1GM	N021574 002	Apr 27, 2004
AB	+ BRISTOL MYERS SQUIBB	750MG	N021202 004	Apr 11, 2003

METFORMIN HYDROCHLORIDE

AB	ALVOGEN	750MG	A078321 002	Apr 17, 2008
AB	AMNEAL PHARMS NY	750MG	A078596 002	Jan 03, 2008
AB	APOTEX	750MG	A076706 002	Dec 29, 2005
AB	BARR	750MG	A076863 001	Oct 14, 2004
AB	IMPAX LABS	750MG	A076985 001	Sep 13, 2005
AB	LUPIN LTD	1GM	A090692 002	Jun 29, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 278 (of 424)

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>750MG</u>	<u>A077113</u>	<u>001</u>	Sep 08, 2005
<u>AB</u>	RANBAXY	<u>750MG</u>	<u>A077211</u>	<u>001</u>	Jun 29, 2005
<u>AB</u>	SUN PHARM INDS (IN)	<u>750MG</u>	<u>A077336</u>	<u>002</u>	Feb 09, 2006
<u>AB</u>	TEVA	<u>750MG</u>	<u>A076864</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	TORRENT PHARMS	<u>750MG</u>	<u>A079226</u>	<u>001</u>	Feb 18, 2010
<u>AB</u>	WATSON LABS FLORIDA	<u>750MG</u>	<u>A076869</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>750MG</u>	<u>A077078</u>	<u>001</u>	Apr 21, 2005

GLUCOPHAGE XR

<u>AB1</u>	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N021202</u>	<u>001</u>	Oct 13, 2000
------------	----------------------	--------------	----------------	------------	--------------

METFORMIN HYDROCHLORIDE

<u>AB1</u>	ALVOGEN	<u>500MG</u>	<u>A078321</u>	<u>001</u>	Apr 17, 2008
<u>AB1</u>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A078596</u>	<u>001</u>	Jan 03, 2008
<u>AB1</u>	APOTEX	<u>500MG</u>	<u>A076706</u>	<u>001</u>	Dec 14, 2004
<u>AB1</u>	IMPAX LABS	<u>500MG</u>	<u>A076249</u>	<u>001</u>	Jul 30, 2004
<u>AB1</u>	MYLAN	<u>500MG</u>	<u>A076650</u>	<u>001</u>	Sep 13, 2005
<u>AB1</u>	NOSTRUM	<u>500MG</u>	<u>A076756</u>	<u>001</u>	Jul 26, 2006
<u>AB1</u>	RANBAXY LABS LTD	<u>500MG</u>	<u>A076413</u>	<u>001</u>	Jun 18, 2004
<u>AB1</u>	SANDOZ	<u>500MG</u>	<u>A076873</u>	<u>001</u>	Dec 14, 2004
<u>AB1</u>	SUN PHARM INDS (IN)	<u>500MG</u>	<u>A077336</u>	<u>001</u>	Feb 09, 2006
<u>AB1</u>	TEVA	<u>500MG</u>	<u>A076269</u>	<u>001</u>	Jun 18, 2004
<u>AB1</u>	TORRENT PHARM	<u>500MG</u>	<u>A090014</u>	<u>001</u>	Dec 30, 2009
<u>AB1</u>	WATSON LABS FLORIDA	<u>500MG</u>	<u>A076172</u>	<u>001</u>	Jun 16, 2004
<u>AB1</u>	WATSON LABS INC	<u>500MG</u>	<u>A076818</u>	<u>001</u>	Dec 14, 2004
<u>AB1</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077060</u>	<u>001</u>	Apr 20, 2005

FORTAMET

<u>AB2</u>	ANDRX LABS LLC	<u>500MG</u>	<u>N021574</u>	<u>001</u>	Apr 27, 2004
------------	----------------	--------------	----------------	------------	--------------

METFORMIN HYDROCHLORIDE

<u>AB2</u>	LUPIN LTD	<u>500MG</u>	<u>A090692</u>	<u>001</u>	Jun 29, 2011
BX	GLUMETZA				

BX	SANTARUS	500MG	N021748	001	Jun 03, 2005
BX	+	1GM	N021748	002	Jun 03, 2005

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOPLUS MET

<u>AB</u>	TAKEDA GLOBAL	<u>500MG; EQ 15MG BASE</u>	<u>N021842</u>	<u>001</u>	Aug 29, 2005
<u>AB</u>	+	<u>850MG; EQ 15MG BASE</u>	<u>N021842</u>	<u>002</u>	Aug 29, 2005

PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>500MG; EQ 15MG BASE</u>	<u>A090406</u>	<u>001</u>	Feb 25, 2011
<u>AB</u>	+	<u>850MG; EQ 15MG BASE</u>	<u>A090406</u>	<u>002</u>	Feb 25, 2011

TABLET, EXTENDED RELEASE; ORAL

ACTOPLUS MET XR

TAKEDA GLOBAL	1GM;EQ 15MG BASE	N022024	001	May 12, 2009
+	1GM;EQ 30MG BASE	N022024	002	May 12, 2009

METFORMIN HYDROCHLORIDE; REPAGLINIDE

TABLET; ORAL

PRANDIMET

NOVO NORDISK INC	500MG;1MG	N022386	001	Jun 23, 2008
+	500MG;2MG	N022386	002	Jun 23, 2008

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDAMET

SB PHARMCO	500MG;EQ 2MG BASE	N021410	002	Oct 10, 2002
	500MG;EQ 4MG BASE	N021410	003	Oct 10, 2002
	1GM;EQ 2MG BASE	N021410	004	Aug 25, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 279 (of 424)

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL
 AVANDAMET
 + SB PHARMCO 1GM;EQ 4MG BASE N021410 005 Aug 25, 2003

METFORMIN HYDROCHLORIDE; SAXAGLIPTIN

TABLET, EXTENDED RELEASE; ORAL
 KOMBIGLYZE XR
 BRISTOL MYERS SQUIBB 500MG;5MG N200678 001 Nov 05, 2010
 1GM;2.5MG N200678 003 Nov 05, 2010
 + 1GM;5MG N200678 002 Nov 05, 2010

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET; ORAL
 JANUMET
 MERCK 500MG;EQ 50MG BASE N022044 001 Mar 30, 2007
 + 1GM;EQ 50MG BASE N022044 002 Mar 30, 2007

METHACHOLINE CHLORIDE

FOR SOLUTION; INHALATION
 PROVOCHOLINE
 + METHAPHARM 100MG/VIAL N019193 001 Oct 31, 1986

METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL
METHADONE HYDROCHLORIDE
AA ROXANE 10MG/ML A040180 001 Apr 30, 1998
AA VISTAPHARM 10MG/ML A040088 001 Nov 30, 1994
METHADONE HYDROCHLORIDE INTENSOL
AA ROXANE 10MG/ML A089897 001 Sep 06, 1988
METHADOSE
AA + MALLINCKRODT 10MG/ML N017116 002

INJECTABLE; INJECTION
 DOLOPHINE HYDROCHLORIDE
 + MYLAN INSTITUTIONAL 10MG/ML N021624 001

POWDER; FOR RX COMPOUNDING
 METHADONE HYDROCHLORIDE
 MALLINCKRODT 50GM/BOT N006383 002
 100GM/BOT N006383 003
 500GM/BOT N006383 004

SOLUTION; ORAL

METHADONE HYDROCHLORIDE
AA + ROXANE 5MG/5ML A087393 001
AA + 10MG/5ML A087997 001 Aug 30, 1982
AA VISTAPHARM 5MG/5ML A090707 001 Jun 30, 2010
AA + 10MG/5ML A090707 002 Jun 30, 2010

TABLET; ORAL

DOLOPHINE HYDROCHLORIDE
AA + ROXANE 5MG N006134 002
AA + 10MG N006134 010
METHADONE HYDROCHLORIDE
AA MALLINCKRODT 5MG A040517 001 Apr 27, 2004
AA 10MG A040517 002 Apr 27, 2004
AA 40MG A077142 001 Jul 12, 2005
AA + ROXANE 40MG N017058 001
AA SANDOZ 10MG A040241 002 May 29, 1998
AA 40MG A075082 001 Mar 25, 1998
AA THE PHARMANETWORK 10MG A090635 001 Nov 25, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 280 (of 424)

METHADONE HYDROCHLORIDE

TABLET; ORAL

METHADOSE

<u>AA</u>	MALLINCKRODT	<u>5MG</u>	<u>A040050</u> <u>001</u>	Apr 15, 1993
<u>AA</u>		<u>10MG</u>	<u>A040050</u> <u>002</u>	Apr 15, 1993
<u>AA</u>		<u>40MG</u>	<u>A074184</u> <u>001</u>	Apr 29, 1993

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DESOXYN

<u>AA</u>	+ LUNDBECK INC	<u>5MG</u>	<u>N005378</u> <u>002</u>	
<u>AA</u>	METHAMPHETAMINE HYDROCHLORIDE			
<u>AA</u>	COASTAL PHARMS	<u>5MG</u>	<u>A091189</u> <u>001</u>	Apr 21, 2010

METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

<u>AB</u>	MIKART	<u>25MG</u>	<u>A040062</u> <u>001</u>	Jan 27, 1994
<u>AB</u>	+	<u>50MG</u>	<u>A040062</u> <u>002</u>	Jan 27, 1994
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A040036</u> <u>001</u>	Jun 30, 1993
<u>AB</u>		<u>50MG</u>	<u>A040036</u> <u>002</u>	Jun 30, 1993
<u>AB</u>	TEVA PHARMS	<u>25MG</u>	<u>A040001</u> <u>001</u>	Jun 30, 1993
<u>AB</u>		<u>50MG</u>	<u>A040001</u> <u>002</u>	Jun 30, 1993

METHENAMINE HIPPURATE

TABLET; ORAL

HIPREX

<u>AB</u>	+ SANOFI AVENTIS US	<u>1GM</u>	<u>N017681</u> <u>001</u>	
<u>AB</u>	METHENAMINE HIPPURATE			
<u>AB</u>	COREPHARMA	<u>1GM</u>	<u>A076411</u> <u>001</u>	Jun 20, 2003

UREX

<u>AB</u>	CNTY LINE PHARMS	<u>1GM</u>	<u>N016151</u> <u>001</u>
-----------	------------------	------------	---------------------------

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

<u>AB</u>	CARACO	<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>AB</u>	CEDAR PHARMS	<u>5MG</u>	<u>A040547</u> <u>001</u>	Feb 18, 2005
<u>AB</u>		<u>10MG</u>	<u>A040547</u> <u>002</u>	Feb 18, 2005
<u>AB</u>	EMCURE PHARMS USA	<u>5MG</u>	<u>A040734</u> <u>001</u>	Dec 14, 2007
<u>AB</u>		<u>10MG</u>	<u>A040734</u> <u>002</u>	Dec 14, 2007
<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>	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

TAPAZOLE

<u>AB</u>	KING PHARMS	<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

INJECTABLE; INJECTION

METHOCARBAMOL

<u>AP</u>	WATSON LABS	<u>100MG/ML</u>	<u>A086459</u> <u>001</u>
-----------	-------------	-----------------	---------------------------

ROBAXIN

<u>AP</u>	+ BAXTER HLTHCARE CORP	<u>100MG/ML</u>	<u>N011790</u> <u>001</u>
-----------	------------------------	-----------------	---------------------------

TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	AUSTARPHARMA LLC	<u>500MG</u>	<u>A200958</u> <u>001</u>	Oct 21, 2011
-----------	------------------	--------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 281 (of 424)

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	AUSTARPHARMA LLC	<u>750MG</u>	<u>A200958</u>	<u>002</u>	Oct 21, 2011
<u>AA</u>	HETERO DRUGS	<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>	LANNETT HOLDINGS INC	<u>500MG</u>	<u>A084756</u>	<u>002</u>	Mar 31, 2003
<u>AA</u>		<u>750MG</u>	<u>A084756</u>	<u>001</u>	
<u>AA</u>	SANDOZ	<u>500MG</u>	<u>A084616</u>	<u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A084615</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>500MG</u>	<u>A085180</u>	<u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A084276</u>	<u>002</u>	
<u>AA</u>		<u>750MG</u>	<u>A085192</u>	<u>001</u>	
<u>AA</u>	WEST WARD	<u>500MG</u>	<u>A085159</u>	<u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A085123</u>	<u>001</u>	
	<u>ROBAXIN</u>				
<u>AA</u>	+ SCHWARZ PHARMA	<u>500MG</u>	<u>N011011</u>	<u>004</u>	
	<u>ROBAXIN-750</u>				
<u>AA</u>	+ SCHWARZ PHARMA	<u>750MG</u>	<u>N011011</u>	<u>006</u>	

METHOHEXITAL SODIUMINJECTABLE; INJECTION
BREVITAL SODIUM

+ JHP PHARMS	500MG/VIAL	N011559 001
+	2.5GM/VIAL	N011559 002

METHOTREXATE SODIUMINJECTABLE; INJECTION
METHOTREXATE SODIUM

<u>AP</u>	+ APP PHARMS	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040263</u>	<u>001</u>	Feb 26, 1999
<u>AP</u>	+	<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040263</u>	<u>002</u>	Feb 26, 1999
<u>AP</u>	+ HOSPIRA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>010</u>	Dec 15, 2004
	<u>METHOTREXATE SODIUM PRESERVATIVE FREE</u>				
<u>AP</u>	+ BEDFORD	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A089340</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	+	<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A089343</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>	+ BIONICHE PHARMA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040767</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	+ BIONICHE PHARMA USA	<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A040768</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	+	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040716</u>	<u>001</u>	Apr 30, 2007
<u>AP</u>	EBEWE PHARMA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A090039</u>	<u>001</u>	Mar 31, 2009
<u>AP</u>		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	<u>A090039</u>	<u>002</u>	Mar 31, 2009
<u>AP</u>		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A090029</u>	<u>001</u>	Mar 31, 2009
<u>AP</u>	+ HOSPIRA	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>N011719</u>	<u>012</u>	Apr 13, 2005
<u>AP</u>	PHARMACHEMIE BV	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	<u>A040850</u>	<u>001</u>	Jan 11, 2010
<u>AP</u>		<u>EQ 250MG/10ML (EQ 25MG BASE/ML)</u>	<u>A040853</u>	<u>001</u>	Jan 11, 2010
<u>AP</u>		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	<u>A040843</u>	<u>001</u>	Jan 11, 2010

METHOTREXATE SODIUM

+ BEDFORD	EQ 100MG BASE/4ML (EQ 25MG BASE/ML)
+	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)

METHOTREXATE SODIUM PRESERVATIVE FREE

+ BEDFORD	EQ 1GM BASE/VIAL
-----------	------------------

A040632 001 Aug 12, 2005

TABLET; ORAL

METHOTREXATE SODIUM

<u>AB</u>	BARR	<u>EQ 2.5MG BASE</u>	<u>A081099</u>	<u>001</u>	Oct 15, 1990
<u>AB</u>	+ DAVA PHARMS INC	<u>EQ 2.5MG BASE</u>	<u>N008085</u>	<u>002</u>	
<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE</u>	<u>A081235</u>	<u>001</u>	May 15, 1992
<u>AB</u>	ROXANE	<u>EQ 2.5MG BASE</u>	<u>A040054</u>	<u>001</u>	Aug 01, 1994

PRESCRIPTION DRUG PRODUCT LIST

3 - 282 (of 424)

METHOTREXATE SODIUM

TABLET; ORAL 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 8-MOP + VALEANT PHARM INTL	10MG	N009048 001
OXSORALEN-ULTRA		
+ VALEANT PHARM INTL	10MG	N019600 001 Oct 30, 1986
INJECTABLE; INJECTION UVADEX		
+ THERAKOS	0.02MG/ML	N020969 001 Feb 25, 1999
LOTION; TOPICAL OXSORALEN		
+ VALEANT PHARM INTL	1%	N009048 002

METHSCOPOLAMINE BROMIDE

TABLET; ORAL METHSCOPOLAMINE BROMIDE		
AA BOCA PHARMA	<u>2.5MG</u>	<u>A040624 001</u> Dec 28, 2006
AA	<u>5MG</u>	<u>A040624 002</u> Dec 28, 2006
AA BRECKENRIDGE PHARM	<u>2.5MG</u>	<u>A040642 001</u> Dec 06, 2011
AA	<u>5MG</u>	<u>A040642 002</u> Dec 06, 2011
PAMINE		
AA + FOUGERA PHARMS	<u>2.5MG</u>	<u>N008848 001</u>
PAMINE FORTE		
AA + FOUGERA PHARMS	<u>5MG</u>	<u>N008848 002</u> Mar 25, 2003

METHSUXIMIDE

CAPSULE; ORAL CELONTIN		
PARKE DAVIS	150MG	N010596 007
+	300MG	N010596 008

METHYCLOTHIAZIDE

TABLET; ORAL ENDURON		
AB + ABBOTT	<u>5MG</u>	<u>N012524 004</u>
METHYCLOTHIAZIDE		
AB MYLAN	<u>5MG</u>	<u>A087672 001</u> Aug 17, 1982
AB WATSON LABS	<u>5MG</u>	<u>A088724 001</u> Sep 06, 1984
ENDURON		
ABBOTT	2.5MG	N012524 001

METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL METVIXIA		
+ GALDERMA LABS LP	EQ 16.8% BASE	N021415 001 Jul 27, 2004

METHYLDOPA

TABLET; ORAL METHYLDOPA		
AB ACCORD HLTH	<u>250MG</u>	<u>A070084 001</u> Oct 15, 1985
AB	<u>500MG</u>	<u>A070085 001</u> Oct 15, 1985

PREScription DRUG PRODUCT LIST

3 - 283 (of 424)

METHYLDOPA

TABLET; ORAL

METHYLDOPA

<u>AB</u>	IVAX	SUB	TEVA	PHARMS	<u>250MG</u>	<u>A070098</u>	<u>001</u>	Feb 20, 1986
<u>AB</u>					<u>500MG</u>	<u>A070343</u>	<u>001</u>	Feb 20, 1986
<u>AB</u>			MYLAN		<u>250MG</u>	<u>A070075</u>	<u>001</u>	Apr 18, 1985
<u>AB</u>	+				<u>500MG</u>	<u>A070076</u>	<u>001</u>	Apr 18, 1985
<u>AB</u>			WATSON LABS		<u>500MG</u>	<u>A070625</u>	<u>001</u>	Jun 06, 1986

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

METHYLDOPATE HYDROCHLORIDE

AP + LUITPOLD 50MG/ML A071279 001 Oct 02, 1987
AP TEVA PARENTERAL 50MG/ML A072974 001 Nov 22, 1991

METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHERGINE

AP + NOVARTIS 0.2MG/ML N006035 004
METHYLERGONOVINE MALEATE

AP ERGOJECT 0.2MG/ML A040889 001 Sep 13, 2010

AP LUITPOLD 0.2MG/ML A090193 001 Nov 24, 2008

TABLET; ORAL

METHERGINE

AB + NOVARTIS 0 .2MG N006035 003
METHYLERGONOVINE MALEATE
AB NOVEL LABS INC 0 .2MG A091577 001 May 02, 2011

METHYLNALTREXONE BROMIDE

INJECTABLE; SUBCUTANEOUS

RELISTOR

PROGENICS 8MG/0.4ML N021964 002 Sep 27, 2010

SOLUTION; SUBCUTANEOUS

RELISTOR

+ PROGENICS 12MG/0.6ML (12MG/0.6ML) N021964 001 Apr 24, 2008

METHYLPHENIDATE

FILM, EXTENDED RELEASE; TRANSDERMAL

DAYTRANA

NOVEN PHARMS INC	10MG/9HR (1.1MG/HR)	N021514 001	Apr 06, 2006
	15MG/9HR (1.6MG/HR)	N021514 002	Apr 06, 2006
	20MG/9HR (2.2MG/HR)	N021514 003	Apr 06, 2006
+	30MG/9HR (3.3MG/HR)	N021514 004	Apr 06, 2006

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ACTAVIS	<u>20MG</u>	<u>A078458</u>	<u>001</u>	Dec 01, 2011
<u>AB</u>		<u>30MG</u>	<u>A078458</u>	<u>002</u>	Dec 01, 2011
<u>AB</u>		<u>40MG</u>	<u>A078458</u>	<u>003</u>	Dec 01, 2011
	<u>RITALIN LA</u>				
<u>AB</u>	NOVARTIS	<u>20MG</u>	<u>N021284</u>	<u>001</u>	Jun 05, 2002
<u>AB</u>		<u>30MG</u>	<u>N021284</u>	<u>002</u>	Jun 05, 2002
<u>AB</u>	+	<u>40MG</u>	<u>N021284</u>	<u>003</u>	Jun 05, 2002
	METADATE CD				
BX	UCB INC	10MG	<u>N021259</u>	<u>003</u>	May 27, 2003
BX		20MG	<u>N021259</u>	<u>001</u>	Apr 03, 2001
BX		30MG	<u>N021259</u>	<u>002</u>	Jun 19, 2003
BX		40MG	<u>N021259</u>	<u>004</u>	Feb 19, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 284 (of 424)

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL RITALIN LA			
BX	NOVARTIS	10MG	N021284 004 Apr 10, 2004
	METADATE CD		
	UCB INC	50MG	N021259 005 Feb 19, 2006
+		60MG	N021259 006 Feb 19, 2006
SOLUTION; ORAL <u>METHYLIN</u>			
AA	+ MALLINCKRODT	<u>5MG/5ML</u>	<u>N021419 001</u> Dec 19, 2002
AA	+	<u>10MG/5ML</u>	<u>N021419 002</u> Dec 19, 2002
<u>METHYLPHENIDATE HYDROCHLORIDE</u>			
AA	TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A091601 001</u> Jul 23, 2010
AA		<u>10MG/5ML</u>	<u>A091601 002</u> Jul 23, 2010
TABLET; ORAL <u>METHYLPHENIDATE HYDROCHLORIDE</u>			
AB	MALLINCKRODT	<u>5MG</u>	<u>A040300 001</u> Nov 27, 1998
AB		<u>10MG</u>	<u>A040300 002</u> Nov 27, 1998
AB		<u>20MG</u>	<u>A040300 003</u> Nov 27, 1998
AB	UCB INC	<u>5MG</u>	<u>A086429 001</u>
AB		<u>10MG</u>	<u>A085799 001</u>
AB		<u>20MG</u>	<u>A086428 001</u>
AB	WATSON LABS	<u>5MG</u>	<u>A040220 001</u> Aug 29, 1997
AB		<u>10MG</u>	<u>A040220 002</u> Aug 29, 1997
AB		<u>20MG</u>	<u>A040220 003</u> Aug 29, 1997
<u>RITALIN</u>			
AB	NOVARTIS	<u>5MG</u>	<u>N010187 003</u>
AB		<u>10MG</u>	<u>N010187 006</u>
AB	+	<u>20MG</u>	<u>N010187 010</u>
TABLET, CHEWABLE; ORAL METHYLIN			
	MALLINCKRODT	2.5MG	N021475 001 Apr 15, 2003
		5MG	N021475 002 Apr 15, 2003
+		10MG	N021475 003 Apr 15, 2003
TABLET, EXTENDED RELEASE; ORAL <u>METADATE ER</u>			
AB	UCB INC	<u>10MG</u>	<u>A040306 001</u> Oct 20, 1999
AB	+	<u>20MG</u>	<u>A089601 001</u> Jun 01, 1988
<u>METHYLIN ER</u>			
AB	MALLINCKRODT	<u>10MG</u>	<u>A075629 001</u> May 09, 2000
AB		<u>20MG</u>	<u>A075629 002</u> May 09, 2000
<u>METHYLPHENIDATE HYDROCHLORIDE</u>			
AB	WATSON LABS	<u>20MG</u>	<u>A040410 001</u> Feb 09, 2001
<u>RITALIN-SR</u>			
AB	NOVARTIS	<u>20MG</u>	<u>N018029 001</u> Mar 30, 1982
CONCERTA			
	JANSSEN PHARMS	18MG	N021121 001 Aug 01, 2000
		27MG	N021121 004 Apr 01, 2002
		36MG	N021121 002 Aug 01, 2000
+		54MG	N021121 003 Dec 08, 2000

METHYLPREDNISOLONE

TABLET; ORAL <u>MEDROL</u>			
AB	PHARMACIA AND UPJOHN	<u>4MG</u>	<u>N011153 001</u>
AB		<u>8MG</u>	<u>N011153 004</u>
AB		<u>16MG</u>	<u>N011153 003</u>
AB	+	<u>32MG</u>	<u>N011153 006</u>
<u>METHYLPREDNISOLONE</u>			
AB	DURAMED PHARMS BARR	<u>4MG</u>	<u>A088497 001</u> Feb 21, 1984

PREScription DRUG PRODUCT LIST

3 - 285 (of 424)

METHYLPREDNISOLONE

TABLET; ORAL

METHYLPREDNISOLONE

<u>AB</u>	JUBILANT CADISTA	<u>4MG</u>	<u>A040189</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>		<u>8MG</u>	<u>A040189</u>	<u>002</u>	Oct 31, 1997
<u>AB</u>		<u>16MG</u>	<u>A040189</u>	<u>003</u>	Jul 20, 2007
<u>AB</u>		<u>32MG</u>	<u>A040189</u>	<u>004</u>	Jul 20, 2007
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A040194</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	VINTAGE PHARMS	<u>4MG</u>	<u>A040183</u>	<u>001</u>	Dec 22, 1998
<u>AB</u>	WATSON LABS	<u>4MG</u>	<u>A040232</u>	<u>001</u>	Oct 16, 1997

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

DEPO-MEDROL

<u>AB</u>	<u>+</u>	PHARMACIA AND UPJOHN	<u>40MG/ML</u>	<u>N011757</u>	<u>001</u>	
<u>AB</u>	<u>+</u>		<u>80MG/ML</u>	<u>N011757</u>	<u>004</u>	
<u>METHYLPREDNISOLONE ACETATE</u>						
<u>AB</u>		SANDOZ	<u>40MG/ML</u>	<u>A040719</u>	<u>001</u>	Jan 29, 2009
<u>AB</u>			<u>40MG/ML</u>	<u>A040794</u>	<u>001</u>	Mar 05, 2009
<u>AB</u>			<u>80MG/ML</u>	<u>A040719</u>	<u>002</u>	Jan 29, 2009
<u>AB</u>			<u>80MG/ML</u>	<u>A040794</u>	<u>002</u>	Mar 05, 2009
<u>AB</u>		TEVA PARENTERAL	<u>40MG/ML</u>	<u>A040557</u>	<u>001</u>	Feb 23, 2005
<u>AB</u>			<u>40MG/ML</u>	<u>A040620</u>	<u>001</u>	Oct 27, 2006
<u>AB</u>			<u>80MG/ML</u>	<u>A040557</u>	<u>002</u>	Feb 23, 2005
<u>AB</u>			<u>80MG/ML</u>	<u>A040620</u>	<u>002</u>	Oct. 27, 2006

DEPO-MEDROTI

+ PHARMACIA AND UPTON 20MG/ML N011757 002

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

<u>AP</u>	HEMOFARM	<u>EQ 40MG BASE/VIAL</u>	<u>A040793</u>	<u>001</u>	Nov 25, 2008
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040827</u>	<u>001</u>	Nov 25, 2008
<u>AP</u>	HOSPIRA	<u>EQ 40MG BASE/VIAL</u>	<u>A040664</u>	<u>001</u>	Dec 20, 2005
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040665</u>	<u>001</u>	Dec 20, 2005

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>	APP PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>A040583</u>	<u>001</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040583</u>	<u>002</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040612</u>	<u>001</u>	Aug 12, 2004
<u>AP</u>	BEDFORD LABS	<u>EQ 40MG BASE/VIAL</u>	<u>A040662</u>	<u>001</u>	Feb 21, 2007
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040641</u>	<u>002</u>	Feb 21, 2007
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A040641</u>	<u>003</u>	Feb 21, 2007
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A040709</u>	<u>001</u>	Feb 21, 2007
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040641</u>	<u>004</u>	Feb 21, 2007
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040709</u>	<u>002</u>	Feb 21, 2007
<u>AP</u>	MUSTAFA NEVSAT	<u>EQ 40MG BASE/VIAL</u>	<u>A040888</u>	<u>001</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040888</u>	<u>002</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A040888</u>	<u>003</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040888</u>	<u>004</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A040888</u>	<u>005</u>	Jul 18, 2011
<u>AP</u>	TEVA PARENTERAL	<u>EQ 125MG BASE/VIAL</u>	<u>A081266</u>	<u>001</u>	Nov 30, 1992

SOT-II=MEDBOT.

<u>AP</u> + PHARMACIA AND UPJOHN	<u>EQ 40MG BASE/VIAL</u>	<u>N011856</u>	<u>003</u>
<u>AP</u> +	<u>EQ 125MG BASE/VIAL</u>	<u>N011856</u>	<u>004</u>
<u>AP</u> +	<u>EQ 500MG BASE/VIAL</u>	<u>N011856</u>	<u>005</u>
<u>AP</u> +	<u>EQ 1GM BASE/VIAL</u>	<u>N011856</u>	<u>006</u>
<u>AP</u> +	<u>EQ 2GM BASE/VIAL</u>	<u>N011856</u>	<u>007</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 286 (of 424)

METHYLTESTOSTERONE

CAPSULE; ORAL TESTRED		
+ VALEANT PHARM INTL	10MG	A083976 001
TABLET; ORAL ANDROID 10		
BP VALEANT PHARM INTL	10MG	A086450 001
ANDROID 25		
BP + VALEANT PHARM INTL	25MG	A087147 001
METHYLTESTOSTERONE		
BP IMPAX LABS	10MG	A080767 002
BP	25MG	A084310 001

METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC <u>METIPRANOLOL</u>		
AT FALCON PHARMS	<u>0.3%</u>	<u>A075720 001</u> Aug 06, 2001
<u>OPTIPRANOLOL</u>		
AT + BAUSCH AND LOMB	<u>0.3%</u>	<u>N019907 001</u> Dec 29, 1989

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION <u>METOCLOPRAMIDE HYDROCHLORIDE</u>		
AP HOSPIRA	<u>EQ 5MG BASE/ML</u>	<u>A073118 001</u> Jan 17, 1991
AP TEVA PARENTERAL	<u>EQ 5MG BASE/ML</u>	<u>A073135 001</u> Nov 27, 1991
<u>REGLAN</u>		
AP + BAXTER HLTHCARE CORP	<u>EQ 5MG BASE/ML</u>	<u>N017862 001</u>
SOLUTION; ORAL <u>METOCLOPRAMIDE HYDROCHLORIDE</u>		
AA ANI PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A071402 001</u> Jun 25, 1993
AA PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>A072744 001</u> May 28, 1991
AA SILARK	<u>EQ 5MG BASE/5ML</u>	<u>A073680 001</u> Oct 27, 1992
AA VISTAPHARM	<u>EQ 5MG BASE/5ML</u>	<u>A075051 001</u> Jan 26, 2001
AA + WOCKHARDT	<u>EQ 5MG BASE/5ML</u>	<u>A074703 001</u> Oct 31, 1997
TABLET; ORAL <u>METOCLOPRAMIDE HYDROCHLORIDE</u>		
AB ACTAVIS ELIZABETH	<u>EQ 10MG BASE</u>	<u>A070581 001</u> Oct 17, 1985
AB IPCA LABS LTD	<u>EQ 5MG BASE</u>	<u>A078807 001</u> Jun 12, 2008
AB	<u>EQ 10MG BASE</u>	<u>A078807 002</u> Jun 12, 2008
AB MUTUAL PHARM	<u>EQ 5MG BASE</u>	<u>A071536 002</u> Jan 16, 1997
AB NORTHSTAR HLTHCARE	<u>EQ 5MG BASE</u>	<u>A078374 001</u> Nov 30, 2007
AB	<u>EQ 10MG BASE</u>	<u>A078374 002</u> Nov 30, 2007
AB TEVA	<u>EQ 5MG BASE</u>	<u>A072801 001</u> Jun 15, 1993
AB	<u>EQ 10MG BASE</u>	<u>A070184 001</u> Jul 29, 1985
AB VINTAGE PHARMS	<u>EQ 5MG BASE</u>	<u>A077878 001</u> Aug 28, 2006
AB	<u>EQ 10MG BASE</u>	<u>A077878 002</u> Aug 28, 2006
AB WATSON LABS	<u>EQ 5MG BASE</u>	<u>A072750 001</u> Dec 28, 1995
AB	<u>EQ 10MG BASE</u>	<u>A071250 001</u> Feb 03, 1988
<u>REGLAN</u>		
AB ANI PHARMS	<u>EQ 5MG BASE</u>	<u>N017854 002</u> May 05, 1987
AB +	<u>EQ 10MG BASE</u>	<u>N017854 001</u>
TABLET, ORALLY DISINTEGRATING; ORAL METOZOLV ODT		
SALIX PHARMS	EQ 5MG BASE	N022246 001 Sep 04, 2009
+	EQ 10MG BASE	N022246 002 Sep 04, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 287 (of 424)

METOLAZONE

TABLET; ORAL

METOLAZONE

<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A076698</u>	<u>001</u>	Dec 23, 2003
<u>AB</u>		<u>5MG</u>	<u>A076698</u>	<u>002</u>	Oct 19, 2004
<u>AB</u>		<u>10MG</u>	<u>A076698</u>	<u>003</u>	Oct 19, 2004
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>A076732</u>	<u>001</u>	Dec 19, 2003
<u>AB</u>		<u>5MG</u>	<u>A076466</u>	<u>001</u>	Dec 19, 2003
<u>AB</u>		<u>10MG</u>	<u>A076466</u>	<u>002</u>	Dec 19, 2003
<u>AB</u>	TEVA	<u>2.5MG</u>	<u>A076600</u>	<u>001</u>	Jan 06, 2004
<u>AB</u>		<u>5MG</u>	<u>A076833</u>	<u>001</u>	Mar 01, 2004
<u>AB</u>		<u>10MG</u>	<u>A075543</u>	<u>003</u>	Dec 24, 2003
	<u>ZAROXOLYN</u>				
<u>AB</u>	UCB INC	<u>2.5MG</u>	<u>N017386</u>	<u>001</u>	
<u>AB</u>	+	<u>5MG</u>	<u>N017386</u>	<u>002</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N017386</u>	<u>003</u>	

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 25MG TARTRATE</u>	<u>A202033</u>	<u>001</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A202033</u>	<u>002</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A202033</u>	<u>003</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A202033</u>	<u>004</u>	Dec 15, 2011
<u>AB</u>	NESHER PHARMS	<u>EQ 25MG TARTRATE</u>	<u>A077779</u>	<u>001</u>	Mar 20, 2008
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A077176</u>	<u>001</u>	May 14, 2008
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A076640</u>	<u>002</u>	May 18, 2007
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A076640</u>	<u>001</u>	May 18, 2007
<u>AB</u>	SANDOZ	<u>EQ 25MG TARTRATE</u>	<u>A076969</u>	<u>001</u>	Jul 31, 2006
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A076969</u>	<u>002</u>	May 18, 2007
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A076969</u>	<u>003</u>	Mar 20, 2008
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A076969</u>	<u>004</u>	Mar 20, 2008
<u>AB</u>	WATSON LABS FLORIDA	<u>EQ 25MG TARTRATE</u>	<u>A077118</u>	<u>001</u>	Aug 03, 2009
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A076862</u>	<u>001</u>	Aug 03, 2009
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A077298</u>	<u>001</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A077298</u>	<u>002</u>	Apr 15, 2010
<u>AB</u>	WOCKHARDT	<u>EQ 25MG TARTRATE</u>	<u>A090615</u>	<u>001</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A090615</u>	<u>002</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A090615</u>	<u>003</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A090615</u>	<u>004</u>	Jul 22, 2010
	<u>TOPROL-XL</u>				
<u>AB</u>	ASTRAZENECA	<u>EQ 25MG TARTRATE</u>	<u>N019962</u>	<u>004</u>	Feb 05, 2001
<u>AB</u>	+	<u>EQ 50MG TARTRATE</u>	<u>N019962</u>	<u>001</u>	Jan 10, 1992
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>N019962</u>	<u>002</u>	Jan 10, 1992
<u>AB</u>	+	<u>EQ 200MG TARTRATE</u>	<u>N019962</u>	<u>003</u>	Jan 10, 1992

METOPROLOL TARTRATE

INJECTABLE; INJECTION

LOPRESSOR

<u>AP</u>	+	NOVARTIS	<u>1MG/ML</u>	<u>N018704</u>	<u>001</u>	Mar 30, 1984
		<u>METOPROLOL TARTRATE</u>				
<u>AP</u>		APP PHARMS	<u>1MG/ML</u>	<u>A091045</u>	<u>001</u>	Oct 25, 2010
<u>AP</u>		BEDFORD LABS	<u>1MG/ML</u>	<u>A076495</u>	<u>001</u>	Jul 07, 2003
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077761</u>	<u>001</u>	May 30, 2007
<u>AP</u>		HOSPIRA	<u>1MG/ML</u>	<u>A074133</u>	<u>001</u>	Dec 21, 1993
<u>AP</u>			<u>1MG/ML</u>	<u>A075160</u>	<u>001</u>	Jul 06, 1998
<u>AP</u>			<u>1MG/ML</u>	<u>A078085</u>	<u>001</u>	Apr 29, 2008
<u>AP</u>		LUITPOLD	<u>1MG/ML</u>	<u>A090386</u>	<u>001</u>	Sep 30, 2009
<u>AP</u>			<u>1MG/ML</u>	<u>A091307</u>	<u>001</u>	Dec 29, 2010
<u>AP</u>		SAGENT STRIDES	<u>1MG/ML</u>	<u>A090317</u>	<u>001</u>	Apr 19, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 288 (of 424)

METOPROLOL TARTRATE

INJECTABLE; INJECTION			
<u>METOPROLOL TARTRATE</u>			
<u>AP</u>	SANDOZ	<u>1MG/ML</u>	<u>A077360 001</u> Oct 02, 2007
<u>AP</u>	WATSON LABS	<u>1MG/ML</u>	<u>A074032 001</u> Dec 21, 1993
TABLET; ORAL			
<u>LOPRESSOR</u>			
<u>AB</u>	NOVARTIS	<u>50MG</u>	<u>N017963 001</u>
<u>AB</u>		<u>100MG</u>	<u>N017963 002</u>
<u>METOPROLOL TARTRATE</u>			
<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>	CARACO	<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>	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>	MUTUAL PHARM	<u>25MG</u>	<u>A073654 002</u> Jul 15, 2009
<u>AB</u>		<u>50MG</u>	<u>A073653 001</u> Dec 21, 1993
<u>AB</u>		<u>100MG</u>	<u>A073654 001</u> Dec 21, 1993
<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>	SANDOZ	<u>50MG</u>	<u>A073288 001</u> Mar 25, 1994
<u>AB</u>		<u>100MG</u>	<u>A073289 001</u> Mar 25, 1994
<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

METRONIDAZOLE

CAPSULE; ORAL			
<u>FLAGYL</u>			
<u>AB</u>	+ GD SEARLE LLC	<u>375MG</u>	<u>N020334 001</u> May 03, 1995
<u>METRONIDAZOLE</u>			
<u>AB</u>	ALEMBIC 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			
<u>METROCREAM</u>			
<u>AB</u>	+ GALDERMA LABS LP	<u>0.75%</u>	<u>N020531 001</u> Sep 20, 1995
<u>METRONIDAZOLE</u>			
<u>AB</u>	ALTANA	<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>	+ SANOFI AVENTIS US	<u>1%</u>	<u>N020743 001</u> Sep 26, 1997
GEL; TOPICAL			
<u>METROGEL</u>			
<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
<u>METRONIDAZOLE</u>			
<u>AB</u>	ALTANA	<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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 289 (of 424)

METRONIDAZOLE

GEL; VAGINAL

METROGEL-VAGINAL

<u>AB</u>	+ MEDICIS	<u>0.75%</u>	<u>N020208</u>	<u>001</u>	Aug 17, 1992
<u>AB</u>	<u>METRONIDAZOLE</u>				
<u>AB</u>	TOLMAR	<u>0.75%</u>	<u>A077264</u>	<u>001</u>	Oct 31, 2006
VANDAZOLE					
BX	TEVA PHARMS	0.75%	N021806	001	May 20, 2005

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

<u>AP</u>	+ BAXTER HLTHCARE	<u>500MG/100ML</u>	<u>N018657</u>	<u>001</u>	
<u>AP</u>	+ PFIZER	<u>500MG/100ML</u>	<u>N018353</u>	<u>002</u>	
<u>METRO I.V. IN PLASTIC CONTAINER</u>					
<u>AP</u>	+ B BRAUN	<u>500MG/100ML</u>	<u>N018900</u>	<u>001</u>	Sep 29, 1983
<u>METRONIDAZOLE IN PLASTIC CONTAINER</u>					
<u>AP</u>	CLARIS LIFESCIENCES	<u>500MG/100ML</u>	<u>A078084</u>	<u>001</u>	Mar 31, 2008
<u>AP</u>	+ HOSPIRA	<u>500MG/100ML</u>	<u>N018890</u>	<u>002</u>	Nov 18, 1983

LOTION; TOPICAL

METROLOTION

<u>AB</u>	+ GALDERMA LABS LP	<u>0.75%</u>	<u>N020901</u>	<u>001</u>	Nov 24, 1998
<u>AB</u>	<u>METRONIDAZOLE</u>				
<u>AB</u>	ALTANA	<u>0.75%</u>	<u>A077197</u>	<u>001</u>	May 24, 2006

TABLET; ORAL

FLAGYL

<u>AB</u>	GD SEARLE LLC	<u>250MG</u>	<u>N012623</u>	<u>001</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N012623</u>	<u>003</u>	
<u>METRONIDAZOLE</u>					
<u>AB</u>	ALEMBIC PHARMS LTD	<u>250MG</u>	<u>A079067</u>	<u>001</u>	Mar 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A079067</u>	<u>002</u>	Mar 13, 2009
<u>AB</u>	MUTUAL PHARM	<u>250MG</u>	<u>A070772</u>	<u>001</u>	Jul 16, 1986
<u>AB</u>		<u>500MG</u>	<u>A070773</u>	<u>001</u>	Jul 16, 1986
<u>AB</u>	PLIVA	<u>250MG</u>	<u>A070027</u>	<u>001</u>	Nov 06, 1984
<u>AB</u>		<u>500MG</u>	<u>A070033</u>	<u>001</u>	Dec 06, 1984
<u>AB</u>	TEVA	<u>500MG</u>	<u>A070044</u>	<u>001</u>	Feb 08, 1985
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A070035</u>	<u>001</u>	Dec 20, 1984
<u>AB</u>		<u>250MG</u>	<u>N018764</u>	<u>001</u>	Sep 17, 1982
<u>AB</u>		<u>500MG</u>	<u>N018764</u>	<u>002</u>	Dec 20, 1982

TABLET, EXTENDED RELEASE; ORAL

FLAGYL ER

<u>AB</u>	+ GD SEARLE LLC	<u>750MG</u>	<u>N020868</u>	<u>001</u>	Nov 26, 1997
<u>METRONIDAZOLE</u>			<u>A090222</u>	<u>001</u>	May 05, 2010
<u>AB</u>	ALEMBIC LTD	<u>750MG</u>			

METYRAPONE

CAPSULE; ORAL

METOPIRONE

+	NOVARTIS	250MG	N012911	002	Aug 09, 1996
---	----------	-------	---------	-----	--------------

METYROSINE

CAPSULE; ORAL

DEMSER

+	ATON	250MG	N017871	001	
---	------	-------	---------	-----	--

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

<u>AB</u>	TEVA	<u>150MG</u>	<u>A074377</u>	<u>001</u>	May 16, 1995
<u>AB</u>		<u>200MG</u>	<u>A074377</u>	<u>002</u>	May 16, 1995
<u>AB</u>	+	<u>250MG</u>	<u>A074377</u>	<u>003</u>	May 16, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 290 (of 424)

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A074711</u>	<u>001</u>	Feb 26, 1997
<u>AB</u>		<u>150MG</u>	<u>A074865</u>	<u>001</u>	Apr 13, 1998
<u>AB</u>		<u>200MG</u>	<u>A074711</u>	<u>002</u>	Feb 26, 1997
<u>AB</u>		<u>200MG</u>	<u>A074865</u>	<u>002</u>	Apr 13, 1998
<u>AB</u>		<u>250MG</u>	<u>A074711</u>	<u>003</u>	Feb 26, 1997
<u>AB</u>		<u>250MG</u>	<u>A074865</u>	<u>003</u>	Apr 13, 1998

MICAFUNGIN SODIUM

INJECTABLE; IV (INFUSION)

+ ASTELLAS	50MG/VIAL	N021506	002	Mar 16, 2005
+	100MG/VIAL	N021506	003	Jun 27, 2006

MICONAZOLE

TABLET; Buccal

ORAVIG

+ BIOALLIANCE PHARMA	50MG
----------------------	------

N022404 001 Apr 16, 2010

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>200MG</u>	<u>A073508</u>	<u>001</u>	Nov 19, 1993
<u>AB</u>	+ INSIGHT PHARMS	<u>200MG</u>	<u>N018888</u>	<u>001</u>	Aug 15, 1984

MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+ STIEFEL LABS INC	0.25%;81.35%;15%
--------------------	------------------

N021026 001 Feb 16, 2006

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>	APP PHARMS	<u>EQ 1MG BASE/ML</u>	<u>A075154</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075154</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 1MG BASE/ML</u>	<u>A075243</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075243</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>A075324</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075324</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>	BEDFORD	<u>EQ 1MG BASE/ML</u>	<u>A075247</u>	<u>002</u>	Jun 23, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075247</u>	<u>001</u>	Jun 23, 2000
<u>AP</u>	BEN VENUE	<u>EQ 1MG BASE/ML</u>	<u>A075421</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075421</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>	CLARIS LIFESCIENCES	<u>EQ 1MG BASE/ML</u>	<u>A075637</u>	<u>001</u>	Oct 31, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075637</u>	<u>002</u>	Oct 31, 2000
<u>AP</u>	+ HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075293</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075856</u>	<u>001</u>	Jun 13, 2002
<u>AP</u>	+	<u>EQ 5MG BASE/ML</u>	<u>A075293</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075856</u>	<u>002</u>	Jun 13, 2002
<u>AP</u>	INTL MEDICATED	<u>EQ 1MG BASE/ML</u>	<u>A076144</u>	<u>001</u>	Jan 26, 2005
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A076144</u>	<u>002</u>	Jan 26, 2005
<u>AP</u>	INTL MEDICATION	<u>EQ 1MG BASE/ML</u>	<u>A076020</u>	<u>001</u>	Jul 16, 2004
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A076020</u>	<u>002</u>	Jul 16, 2004
<u>AP</u>	TAYLOR	<u>EQ 1MG BASE/ML</u>	<u>A075494</u>	<u>001</u>	Jun 30, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075481</u>	<u>001</u>	Jun 30, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A075494</u>	<u>002</u>	Jun 30, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 291 (of 424)

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>	WOCKHARDT	<u>EQ 1MG BASE/ML</u>	<u>A078141 001</u>	May 30, 2008
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A078511 001</u>	Nov 10, 2008
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A078141 002</u>	May 30, 2008
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A078511 002</u>	Nov 10, 2008
		<u>MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE</u>		
<u>AP</u>	+ HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>A075857 001</u>	Jul 22, 2002
<u>AP</u>	+ SAGENT STRIDES	<u>EQ 1MG BASE/ML</u>	<u>A075857 002</u>	Jul 22, 2002
<u>AP</u>	SAGENT STRIDES	<u>EQ 5MG BASE/ML</u>	<u>A090315 001</u>	Nov 29, 2010
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A090315 002</u>	Nov 29, 2010
		<u>MIDOZALAM HYDROCHLORIDE</u>		
<u>AP</u>	SAGENT STRIDES	<u>EQ 1MG BASE/ML</u>	<u>A090316 001</u>	May 04, 2011
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>A090316 002</u>	May 04, 2011

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

<u>AA</u>	APOTEX INC	<u>EQ 2MG BASE/ML</u>	<u>A077115 001</u>	Sep 09, 2005
<u>AA</u>	HI TECH PHARMA	<u>EQ 2MG BASE/ML</u>	<u>A075958 001</u>	Sep 04, 2003
<u>AA</u>	PADDOCK LLC	<u>EQ 2MG BASE/ML</u>	<u>A076379 001</u>	May 02, 2005
<u>AA</u>	RANBAXY	<u>EQ 2MG BASE/ML</u>	<u>A076058 001</u>	Mar 15, 2002
<u>AA</u>	+ ROXANE	<u>EQ 2MG BASE/ML</u>	<u>A075873 001</u>	Apr 30, 2002

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>2.5MG</u>	<u>A077746 001</u>	Sep 12, 2006
<u>AB</u>		<u>5MG</u>	<u>A077746 002</u>	Sep 12, 2006
<u>AB</u>		<u>10MG</u>	<u>A077746 003</u>	Sep 12, 2006
<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	<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	<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
		<u>PROAMATINE</u>		
<u>AB</u>	SHIRE	<u>2.5MG</u>	<u>N019815 001</u>	Sep 06, 1996
<u>AB</u>	+	<u>5MG</u>	<u>N019815 002</u>	Sep 06, 1996
<u>AB</u>		<u>10MG</u>	<u>N019815 003</u>	Mar 20, 2002

MIFEPRISTONE

TABLET; ORAL

MIFEPREX

+ DANCO LABS LLC	200MG	N020687 001	Sep 28, 2000
------------------	-------	-------------	--------------

MIGLITOL

TABLET; ORAL

GLYSET

PHARMACIA AND UPJOHN	25MG	N020682 001	Dec 18, 1996
	50MG	N020682 002	Dec 18, 1996
+	100MG	N020682 003	Dec 18, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 292 (of 424)

MIGLUSTAT

CAPSULE; ORAL
 ZAVESCA
 + ACTELION PHARMS LTD 100MG N021348 001 Jul 31, 2003

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL
 SAVELLA
 CYPRESS BIOSCIENCE 12.5MG N022256 001 Jan 14, 2009
 25MG N022256 002 Jan 14, 2009
 + 50MG N022256 003 Jan 14, 2009
 100MG N022256 004 Jan 14, 2009

MILRINONE LACTATE

INJECTABLE; INJECTION
MILRINONE LACTATE
 AP APP PHARMS EQ 1MG BASE/ML A075936 001 May 28, 2002
 AP BAXTER HLTHCARE EQ 1MG BASE/ML A075530 001 May 28, 2002
 AP + BEDFORD EQ 1MG BASE/ML A075660 001 May 28, 2002
 AP CLARIS LIFESCIENCES EQ 1MG BASE/ML A076427 001 Sep 21, 2004
 AP GLAND PHARMA LTD EQ 1MG BASE/ML A077190 001 Oct 31, 2006
 AP HIKMA FARMACEUTICA EQ 1MG BASE/ML A077966 001 Dec 03, 2010
 AP INTL MEDICATED EQ 1MG BASE/ML A076013 001 Aug 02, 2002
MILRINONE LACTATE IN DEXTROSE 5%
 AP CLARIS LIFESCIENCES EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A077151 002 Jul 20, 2005
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER
 AP B BRAUN EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A076414 001 Aug 18, 2004
 AP + BAXTER HLTHCARE EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A075834 001 May 28, 2002
 AP + EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A075834 002 May 28, 2002
 AP BEDFORD LABS EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A078113 001 May 21, 2008
 AP EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A078113 002 May 21, 2008
 AP CLARIS LIFESCIENCES EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A077151 001 Jul 20, 2005
 AP HOSPIRA EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A075885 001 May 28, 2002
 AP EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A075885 002 May 28, 2002
MILRINONE LACTATE IN PLASTIC CONTAINER
 AP HIKMA FARMACEUTICA EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML) A090038 001 Jan 21, 2010
 AP EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML) A090038 002 Jan 21, 2010

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL
DYNACIN
 AB MEDICIS EQ 75MG BASE A063067 002 Sep 15, 1999
 AB EQ 100MG BASE A063067 001 Jul 31, 1990
MINOCIN
 AB TRIAX PHARMS LLC EQ 50MG BASE N050649 001 May 31, 1990
 AB EQ 100MG BASE N050649 002 May 31, 1990
MINOCYCLINE HYDROCHLORIDE
 AB AUROBINDO PHARMA EQ 50MG BASE A065470 001 Mar 11, 2008
 AB EQ 75MG BASE A065470 002 Mar 11, 2008
 AB EQ 100MG BASE A065470 003 Mar 11, 2008
 AB IMPAX LABS EQ 50MG BASE A065005 001 Mar 23, 1999
 AB EQ 75MG BASE A065005 003 Apr 18, 2001
 AB EQ 100MG BASE A065005 002 Mar 23, 1999
 AB RANBAXY EQ 50MG BASE A065062 001 Nov 30, 2000
 AB EQ 75MG BASE A065062 002 Nov 30, 2000
 AB EQ 100MG BASE A065062 003 Nov 30, 2000
 AB TEVA EQ 50MG BASE A063011 001 Mar 02, 1992
 AB EQ 75MG BASE A063009 002 Aug 12, 2003
 AB + EQ 100MG BASE A063009 001 Mar 02, 1992
 AB WATSON LABS EQ 50MG BASE A063181 001 Dec 30, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 293 (of 424)

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

MINOCYCLINE HYDROCHLORIDE

<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

INJECTABLE; INJECTION

MINOCIN

+ TRIAX PHARMS LLC 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>	+ RANBAXY	<u>EQ 100MG BASE</u>	<u>A065131 003</u>	Apr 16, 2003
<u>AB</u>		<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
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065156 003</u>	Jan 06, 2004

TABLET, EXTENDED RELEASE; ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>EQ 45MG BASE</u>	<u>A065485 001</u>	Mar 17, 2009
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A065485 002</u>	Mar 17, 2009
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A065485 003</u>	Mar 17, 2009
<u>AB</u>	IMPAX LABS INC	<u>EQ 45MG BASE</u>	<u>A090024 001</u>	Feb 03, 2009
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A090024 002</u>	Feb 03, 2009
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A090024 003</u>	Feb 03, 2009
<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>	MATRIX LABS LTD	<u>EQ 45MG BASE</u>	<u>A090911 001</u>	Jul 20, 2010
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A090911 002</u>	Jul 20, 2010
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A090911 003</u>	Jul 20, 2010
<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>SOLODYN</u>			
<u>AB</u>	MEDICIS	<u>EQ 45MG BASE</u>	<u>N050808 001</u>	May 08, 2006
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>N050808 002</u>	May 08, 2006
<u>AB</u>	+ SOLODYN	<u>EQ 135MG BASE</u>	<u>N050808 003</u>	May 08, 2006
	MEDICIS	EQ 55MG BASE	N050808 008	Aug 27, 2010
		EQ 65MG BASE	N050808 004	Jul 23, 2009
		EQ 80MG BASE	N050808 007	Aug 27, 2010
		EQ 105MG BASE	N050808 006	Aug 27, 2010
		EQ 115MG BASE	N050808 005	Jul 23, 2009

MINOXIDIL

TABLET; ORAL

MINOXIDIL

<u>AB</u>	MUTUAL PHARM	<u>2.5MG</u>	<u>A072708 001</u>	Dec 14, 1995
<u>AB</u>		<u>10MG</u>	<u>A072709 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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 294 (of 424)

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

<u>AB</u>	ALPHAPHARM	<u>15MG</u>	<u>A076176</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076176</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076176</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	APOTEX INC	<u>15MG</u>	<u>A077666</u>	<u>001</u>	Aug 22, 2007
<u>AB</u>		<u>30MG</u>	<u>A077666</u>	<u>002</u>	Aug 22, 2007
<u>AB</u>		<u>45MG</u>	<u>A077666</u>	<u>003</u>	Aug 22, 2007
<u>AB</u>	AUROBINDO	<u>7.5MG</u>	<u>A076921</u>	<u>001</u>	Oct 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076921</u>	<u>002</u>	Oct 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076921</u>	<u>003</u>	Oct 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076921</u>	<u>004</u>	Oct 22, 2004
<u>AB</u>	CARACO	<u>7.5MG</u>	<u>A076541</u>	<u>004</u>	Apr 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076541</u>	<u>001</u>	Apr 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076541</u>	<u>002</u>	Apr 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076541</u>	<u>003</u>	Apr 22, 2004
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A076122</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076122</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076122</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>A076219</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076219</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076219</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	TEVA	<u>15MG</u>	<u>A076119</u>	<u>001</u>	Jan 24, 2003
<u>AB</u>		<u>30MG</u>	<u>A076119</u>	<u>002</u>	Jan 24, 2003
<u>AB</u>		<u>45MG</u>	<u>A076119</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>A076312</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076312</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076312</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	WATSON LABS FLORIDA	<u>15MG</u>	<u>A076336</u>	<u>001</u>	Jun 20, 2003
<u>AB</u>		<u>30MG</u>	<u>A076336</u>	<u>002</u>	Jun 20, 2003
<u>AB</u>		<u>45MG</u>	<u>A076336</u>	<u>003</u>	Jun 20, 2003
<u>REMERON</u>					
<u>AB</u>	+ ORGANON USA INC	<u>15MG</u>	<u>N020415</u>	<u>001</u>	Jun 14, 1996
<u>AB</u>		<u>30MG</u>	<u>N020415</u>	<u>002</u>	Jun 14, 1996
<u>AB</u>		<u>45MG</u>	<u>N020415</u>	<u>003</u>	Mar 17, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>15MG</u>	<u>A077959</u>	<u>001</u>	Feb 14, 2011
<u>AB</u>		<u>30MG</u>	<u>A077959</u>	<u>002</u>	Feb 14, 2011
<u>AB</u>		<u>45MG</u>	<u>A077959</u>	<u>003</u>	Feb 14, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>15MG</u>	<u>A077376</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>A077376</u>	<u>003</u>	Dec 08, 2005
<u>AB</u>		<u>45MG</u>	<u>A077376</u>	<u>004</u>	Feb 28, 2006
<u>AB</u>	TEVA	<u>15MG</u>	<u>A076901</u>	<u>001</u>	Jun 28, 2005
<u>AB</u>		<u>30MG</u>	<u>A076901</u>	<u>002</u>	Jun 28, 2005
<u>AB</u>		<u>45MG</u>	<u>A076901</u>	<u>003</u>	Jun 28, 2005
<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>A076307</u>	<u>001</u>	Dec 17, 2003
<u>AB</u>		<u>30MG</u>	<u>A076307</u>	<u>002</u>	Dec 17, 2003
<u>AB</u>		<u>45MG</u>	<u>A076307</u>	<u>003</u>	Feb 28, 2006
<u>REMERON SOLTAB</u>					
<u>AB</u>	+ ORGANON USA INC	<u>15MG</u>	<u>N021208</u>	<u>001</u>	Jan 12, 2001
<u>AB</u>		<u>30MG</u>	<u>N021208</u>	<u>002</u>	Jan 12, 2001
<u>AB</u>		<u>45MG</u>	<u>N021208</u>	<u>003</u>	Jan 12, 2001

MISOPROSTOL

TABLET; ORAL

CYTOTEC

<u>AB</u>	GD SEARLE LLC	<u>0.1MG</u>	<u>N019268</u>	<u>003</u>	Sep 21, 1990
<u>AB</u>	+	<u>0.2MG</u>	<u>N019268</u>	<u>001</u>	Dec 27, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 295 (of 424)

MISOPROSTOL

TABLET; ORAL

MISOPROSTOL

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>0.1MG</u>	<u>A076095</u>	<u>001</u>	Jul 10, 2002
<u>AB</u>		<u>0.2MG</u>	<u>A076095</u>	<u>002</u>	Jul 10, 2002

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>	+ ACCORD HLTHCARE	<u>5MG/VIAL</u>	<u>A064144</u>	<u>001</u>	Apr 30, 1998
<u>AP</u>	+	<u>20MG/VIAL</u>	<u>A064144</u>	<u>002</u>	Apr 30, 1998
<u>AP</u>	+	<u>40MG/VIAL</u>	<u>A064144</u>	<u>003</u>	Aug 11, 2009
<u>AP</u>	BAXTER HLTHCARE	<u>5MG/VIAL</u>	<u>A064180</u>	<u>001</u>	Dec 23, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A064180</u>	<u>002</u>	Dec 23, 1999
<u>AP</u>	BEDFORD	<u>5MG/VIAL</u>	<u>A064117</u>	<u>001</u>	Apr 19, 1995
<u>AP</u>		<u>20MG/VIAL</u>	<u>A064117</u>	<u>002</u>	Apr 19, 1995

MITOTANE

TABLET; ORAL

LYSODREN

+ BRISTOL MYERS SQUIBB 500MG

N016885 001

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

<u>AP</u>	APP PHARMS	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077496</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077496</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077496</u>	<u>003</u>	Apr 11, 2006
<u>AP</u>	BEDFORD	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076611</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076611</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076611</u>	<u>003</u>	Apr 11, 2006
<u>AP</u>	FRESENIUS KABI ONCOL	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A078606</u>	<u>001</u>	May 14, 2008
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A078606</u>	<u>002</u>	May 14, 2008
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A078606</u>	<u>003</u>	May 14, 2008
<u>AP</u>	+ HOSPIRA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076871</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>	+	<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076871</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>	+	<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076871</u>	<u>003</u>	Apr 11, 2006
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A078980</u>	<u>001</u>	Apr 13, 2009
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A078980</u>	<u>002</u>	Apr 13, 2009
<u>AP</u>	TEVA PARENTERAL	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077356</u>	<u>001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077356</u>	<u>002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077356</u>	<u>003</u>	Apr 11, 2006

MODAFINIL

TABLET; ORAL

PROVIGIL

CEPHALON

100MG

N020717 001 Dec 24, 1998

+

200MG

N020717 002 Dec 24, 1998

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>7.5MG</u>	<u>A078454</u>	<u>001</u>	Jun 02, 2008
<u>AB</u>		<u>15MG</u>	<u>A078454</u>	<u>002</u>	Jun 02, 2008
<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A090416</u>	<u>001</u>	Mar 30, 2010
<u>AB</u>		<u>15MG</u>	<u>A090416</u>	<u>002</u>	Mar 30, 2010
<u>AB</u>	PADDICK LLC	<u>7.5MG</u>	<u>A077536</u>	<u>001</u>	Nov 30, 2006
<u>AB</u>		<u>15MG</u>	<u>A077536</u>	<u>002</u>	Nov 30, 2006
<u>AB</u>	TEVA	<u>7.5MG</u>	<u>A076204</u>	<u>001</u>	May 08, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 296 (of 424)

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	TEVA	<u>15MG</u>	<u>A076204</u> <u>002</u>	May 08, 2003
<u>AB</u>	<u>UNIVASC</u>			
<u>AB</u>	UCB INC	<u>7.5MG</u>	<u>N020312</u> <u>001</u>	Apr 19, 1995

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCON

<u>AB</u>	+ SCHERING	<u>0.1%</u>	<u>N019625</u> <u>001</u>	May 06, 1987
	<u>MOMETASONE FUROATE</u>			
<u>AB</u>	ALTANA	<u>0.1%</u>	<u>A076171</u> <u>001</u>	Apr 08, 2005
<u>AB</u>	G AND W LABS	<u>0.1%</u>	<u>A077447</u> <u>001</u>	May 22, 2006
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A078541</u> <u>001</u>	May 28, 2008
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076679</u> <u>001</u>	Dec 21, 2004
<u>AB</u>	TOLMAR	<u>0.1%</u>	<u>A076591</u> <u>001</u>	Apr 18, 2007

LOTION; TOPICAL

ELOCON

<u>AB</u>	+ SCHERING	<u>0.1%</u>	<u>N019796</u> <u>001</u>	Mar 30, 1989
	<u>MOMETASONE FUROATE</u>			
<u>AB</u>	G AND W LABS	<u>0.1%</u>	<u>A077678</u> <u>001</u>	Nov 21, 2007
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A090506</u> <u>001</u>	Aug 09, 2010
<u>AB</u>	NYCOMED US	<u>0.1%</u>	<u>A075919</u> <u>001</u>	Nov 29, 2007
<u>AB</u>	PERRIGO	<u>0.1%</u>	<u>A077180</u> <u>001</u>	Apr 06, 2005
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076788</u> <u>001</u>	Mar 15, 2006
<u>AB</u>	TOLMAR	<u>0.1%</u>	<u>A076499</u> <u>001</u>	Nov 21, 2007

OINTMENT; TOPICAL

ELOCON

<u>AB</u>	+ SCHERING	<u>0.1%</u>	<u>N019543</u> <u>001</u>	Apr 30, 1987
	<u>MOMETASONE FUROATE</u>			
<u>AB</u>	ALTANA	<u>0.1%</u>	<u>A077061</u> <u>001</u>	Mar 28, 2005
<u>AB</u>	G AND W LABS	<u>0.1%</u>	<u>A077401</u> <u>001</u>	Jun 20, 2006
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A078571</u> <u>001</u>	May 28, 2008
<u>AB</u>	PERRIGO NEW YORK	<u>0.1%</u>	<u>A076067</u> <u>001</u>	Mar 18, 2002
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076624</u> <u>001</u>	Dec 03, 2004
<u>AB</u>	TOLMAR	<u>0.1%</u>	<u>A076481</u> <u>001</u>	Nov 14, 2003

POWDER; INHALATION

ASMANEX TWISTHALER

SCHERING

0.11MG/INH

+ 0.22MG/INH

N021067 002 Feb 01, 2008
N021067 001 Mar 30, 2005MOMETASONE FUROATE MONOHYDRATE

SPRAY, METERED; NASAL

NASONEX

+ SCHERING PLOUGH

EQ 0.05MG BASE/SPRAY

N020762 001 Oct 01, 1997

MONTELUKAST SODIUM

GRANULE; ORAL

SINGULAIR

+ MERCK

EQ 4MG BASE/PACKET

N021409 001 Jul 26, 2002

TABLET; ORAL

SINGULAIR

+ MERCK

EQ 10MG BASE

N020829 002 Feb 20, 1998

TABLET, CHEWABLE; ORAL

SINGULAIR

MERCK

EQ 4MG BASE

N020830 002 Mar 03, 2000

+ EQ 5MG BASE

N020830 001 Feb 20, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 297 (of 424)

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

KADIAN

<u>AB</u>	ACTAVIS ELIZABETH	<u>20MG</u>	<u>N020616</u>	<u>001</u>	Jul 03, 1996
<u>AB</u>		<u>30MG</u>	<u>N020616</u>	<u>004</u>	Mar 09, 2001
<u>AB</u>		<u>50MG</u>	<u>N020616</u>	<u>002</u>	Jul 03, 1996
<u>AB</u>		<u>60MG</u>	<u>N020616</u>	<u>005</u>	Mar 09, 2001
<u>AB</u>		<u>80MG</u>	<u>N020616</u>	<u>006</u>	Oct 27, 2006
<u>AB</u>	<u>+</u>	<u>100MG</u>	<u>N020616</u>	<u>003</u>	Jul 03, 1996
	<u>MORPHINE SULFATE</u>				
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>A200812</u>	<u>001</u>	Nov 10, 2011
<u>AB</u>		<u>30MG</u>	<u>A200812</u>	<u>002</u>	Nov 10, 2011
<u>AB</u>		<u>50MG</u>	<u>A200812</u>	<u>003</u>	Nov 10, 2011
<u>AB</u>		<u>60MG</u>	<u>A200812</u>	<u>004</u>	Nov 10, 2011
<u>AB</u>		<u>80MG</u>	<u>A200812</u>	<u>005</u>	Nov 10, 2011
<u>AB</u>		<u>100MG</u>	<u>A200812</u>	<u>006</u>	Nov 10, 2011
	AVINZA				
<u>BX</u>	KING PHARMS	30MG	N021260	001	Mar 20, 2002
<u>BX</u>		60MG	N021260	002	Mar 20, 2002
	AVINZA				
	KING PHARMS	45MG	N021260	005	Dec 18, 2008
		75MG	N021260	006	Dec 18, 2008
		90MG	N021260	003	Mar 20, 2002
	<u>+</u>	120MG	N021260	004	Mar 20, 2002
	KADIAN				
	<u>+</u> ACTAVIS ELIZABETH	10MG	N020616	008	Apr 20, 2007
	<u>+</u>	200MG	N020616	007	Feb 27, 2007

INJECTABLE; INJECTION

ASTRAMORPH PF

<u>AP</u>	APP PHARMS	<u>0.5MG/ML</u>	<u>A071050</u>	<u>001</u>	Oct 07, 1986
<u>AP</u>		<u>0.5MG/ML</u>	<u>A071051</u>	<u>001</u>	Oct 07, 1986
<u>AP</u>		<u>1MG/ML</u>	<u>A071052</u>	<u>001</u>	Oct 07, 1986
<u>AP</u>		<u>1MG/ML</u>	<u>A071053</u>	<u>001</u>	Oct 07, 1986
	<u>DURAMORPH PF</u>				
<u>AP</u>	<u>+</u> BAXTER HLTHCARE	<u>0.5MG/ML</u>	N018565	001	Sep 18, 1984
<u>AP</u>	<u>+</u>	<u>1MG/ML</u>	N018565	002	Sep 18, 1984
	<u>MORPHINE SULFATE</u>				
<u>AP</u>	HOSPIRA	<u>0.5MG/ML</u>	<u>A071849</u>	<u>001</u>	May 11, 1988
<u>AP</u>		<u>0.5MG/ML</u>	<u>A073509</u>	<u>001</u>	Sep 30, 1992
<u>AP</u>		<u>1MG/ML</u>	<u>A071850</u>	<u>001</u>	May 11, 1988
<u>AP</u>		<u>1MG/ML</u>	<u>A073510</u>	<u>001</u>	Sep 30, 1992
<u>AP</u>	<u>+</u>	<u>1MG/ML</u>	<u>N019916</u>	<u>001</u>	Oct 30, 1992
<u>AP</u>	WATSON LABS	<u>0.5MG/ML</u>	<u>A073373</u>	<u>001</u>	Sep 30, 1991
<u>AP</u>		<u>0.5MG/ML</u>	<u>A073375</u>	<u>001</u>	Sep 30, 1991
<u>AP</u>		<u>1MG/ML</u>	<u>A073374</u>	<u>001</u>	Sep 30, 1991
<u>AP</u>		<u>1MG/ML</u>	<u>A073376</u>	<u>001</u>	Sep 30, 1991
	INFUMORPH				
	<u>+</u> BAXTER HLTHCARE	10MG/ML	N018565	003	Jul 19, 1991
	<u>+</u>	25MG/ML	N018565	004	Jul 19, 1991
	MORPHINE SULFATE				
	<u>+</u> HOSPIRA	5MG/ML	N019916	002	Oct 27, 2006
	<u>+</u> HOSPIRA INC	2MG/ML	N202515	001	Nov 14, 2011
	<u>+</u>	4MG/ML	N202515	002	Nov 14, 2011
	<u>+</u>	8MG/ML	N202515	003	Nov 14, 2011
	<u>+</u>	10MG/ML	N202515	004	Nov 14, 2011
	<u>+</u>	15MG/ML	N202515	005	Nov 14, 2011
	<u>+</u> MERIDIAN MEDCL TECHN	15MG/ML	N019999	001	Jul 12, 1990

INJECTABLE, LIPOSOMAL; EPIDURAL

DEPODUR

<u>+</u> EKR THERAP	10MG/ML (10MG/ML)	N021671	001	May 18, 2004
---------------------	-------------------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 298 (of 424)

MORPHINE SULFATE

SOLUTION; ORAL

MORPHINE SULFATE

<u>AA</u>	MALLINCKRODT INC	<u>100MG/5ML</u>	<u>A202348 001</u>	Jul 15, 2011
<u>AA</u>	ROXANE	<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
<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
	MORPHINE SULFATE			
	LANNETT HOLDINGS INC	20MG/ML	N201517 001	Jun 23, 2011
	TABLET; ORAL			
	MORPHINE SULFATE			
	ROXANE	15MG	N022207 001	Mar 17, 2008
	+	30MG	N022207 002	Mar 17, 2008
	TABLET, EXTENDED RELEASE; ORAL			
	MORPHINE SULFATE			
<u>AB</u>	CLONMEL HLTHCARE	<u>15MG</u>	<u>A075407 001</u>	Jan 28, 2000
<u>AB</u>	ENDO PHARMS	<u>15MG</u>	<u>A075295 001</u>	Oct 28, 1998
<u>AB</u>		<u>30MG</u>	<u>A075295 002</u>	Oct 28, 1998
<u>AB</u>		<u>60MG</u>	<u>A075295 003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>A075295 004</u>	Sep 15, 2000
<u>AB</u>		<u>200MG</u>	<u>A075295 005</u>	Sep 15, 2000
<u>AB</u>	MALLINCKRODT	<u>15MG</u>	<u>A076412 001</u>	Jul 31, 2003
<u>AB</u>		<u>30MG</u>	<u>A076412 002</u>	Jul 31, 2003
<u>AB</u>		<u>60MG</u>	<u>A076412 003</u>	Jul 31, 2003
<u>AB</u>		<u>100MG</u>	<u>A076438 001</u>	Jul 03, 2003
<u>AB</u>		<u>200MG</u>	<u>A076438 002</u>	Jul 03, 2003
<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
<u>AB</u>		<u>100MG</u>	<u>A077855 001</u>	Sep 27, 2007
<u>AB</u>		<u>200MG</u>	<u>A077855 002</u>	Sep 27, 2007
<u>AB</u>	RHODES PHARMS	<u>15MG</u>	<u>A074862 001</u>	Jul 07, 1998
<u>AB</u>		<u>30MG</u>	<u>A074862 002</u>	Jul 07, 1998
<u>AB</u>		<u>60MG</u>	<u>A074862 003</u>	Jul 07, 1998
<u>AB</u>		<u>100MG</u>	<u>A074769 001</u>	Jul 02, 1998
<u>AB</u>		<u>200MG</u>	<u>A074769 002</u>	Jul 02, 1998
	<u>MS CONTIN</u>			
<u>AB</u>	PURDUE PHARMA LP	<u>15MG</u>	<u>N019516 003</u>	Sep 12, 1989
<u>AB</u>		<u>30MG</u>	<u>N019516 001</u>	May 29, 1987
<u>AB</u>		<u>60MG</u>	<u>N019516 002</u>	Apr 08, 1988
<u>AB</u> +		<u>100MG</u>	<u>N019516 004</u>	Jan 16, 1990
<u>AB</u>		<u>200MG</u>	<u>N019516 005</u>	Nov 08, 1993
	ORAMORPH SR			
BC	XANODYNE PHARM	15MG	N019977 004	Nov 23, 1994
BC		30MG	N019977 001	Aug 15, 1991
BC		60MG	N019977 002	Aug 15, 1991
BC +		100MG	N019977 003	Aug 15, 1991

MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EMBEDA

ALPHARMA KING	20MG;0.8MG	N022321 001	Aug 13, 2009
	30MG;1.2MG	N022321 002	Aug 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 299 (of 424)

MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
EMBEDA

ALPHARMA KING	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

MOXIFLOXACIN HYDROCHLORIDE

INJECTABLE; IV (INFUSION)
AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER
+ BAYER HLTHCARE 160MG/100ML

N021277 001 Nov 30, 2001

SOLUTION/DROPS; OPHTHALMIC

+ ALCON PHARMS LTD	EQ 0.5% BASE	N022428 001	Nov 19, 2010
VIGAMOX			
+ ALCON PHARMS LTD	EQ 0.5% BASE	N021598 001	Apr 15, 2003
TABLET; ORAL			
AVELOX			
+ BAYER HLTHCARE	EQ 400MG BASE	N021085 001	Dec 10, 1999

MUPIROCIN

OINTMENT; TOPICAL

BACTROBAN

<u>AB</u> + GLAXOSMITHKLINE	<u>2%</u>	<u>N050591 001</u>	Dec 31, 1987
<u>MUPIROCIN</u>			
<u>AB</u> ALTANA	<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			
<u>BX</u> PERRIGO NEW YORK	<u>2%</u>	<u>N050788 001</u>	Dec 04, 2002

MUPIROCIN CALCIUM

CREAM; TOPICAL

BACTROBAN

+ GLAXOSMITHKLINE	EQ 2% BASE	N050746 001	Dec 11, 1997
-------------------	------------	-------------	--------------

OINTMENT; NASAL

BACTROBAN

+ GLAXOSMITHKLINE	EQ 2% BASE	N050703 001	Sep 18, 1995
-------------------	------------	-------------	--------------

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 INC	<u>250MG</u>	<u>A090253 001</u>	May 04, 2009
<u>AB</u> APOTEX CORP	<u>250MG</u>	<u>A090419 001</u>	Apr 22, 2009
<u>AB</u> DR REDDYS LABS LTD	<u>250MG</u>	<u>A091315 001</u>	Oct 27, 2011
<u>AB</u> ENDO PHARMS	<u>250MG</u>	<u>A090111 001</u>	Dec 22, 2009
<u>AB</u> MYLAN	<u>250MG</u>	<u>A065520 001</u>	May 04, 2009
<u>AB</u> ROXANE	<u>250MG</u>	<u>A065410 001</u>	Jul 29, 2008
<u>AB</u> SANDOZ	<u>250MG</u>	<u>A065379 001</u>	Oct 15, 2008
<u>AB</u> STRIDES ARCOLAB LTD	<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> ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A065433 001</u>	May 04, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 300 (of 424)

MYCOPHENOLATE MOFETIL

SUSPENSION; ORAL CELLCEPT	+ ROCHE PALO	200MG/ML	N050759 001	Oct 01, 1998
TABLET; ORAL <u>CELLCEPT</u>				
<u>AB</u> + ROCHE PALO	<u>500MG</u>		<u>N050723 001</u>	Jun 19, 1997
<u>MYCOPHENOLATE MOFETIL</u>				
<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> ENDO PHARMS	<u>500MG</u>		<u>A090606 001</u>	Jul 16, 2010
<u>AB</u> MYLAN	<u>500MG</u>		<u>A065521 001</u>	May 04, 2009
<u>AB</u> ROXANE	<u>500MG</u>		<u>A065413 001</u>	Jul 29, 2008
<u>AB</u> SANDOZ	<u>500MG</u>		<u>A065451 001</u>	Oct 15, 2008
<u>AB</u> STRIDES ARCOLAB LTD	<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> ZYDUS PHARMS USA INC	<u>500MG</u>		<u>A065477 001</u>	May 04, 2009

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION CELLCEPT	+ ROCHE PALO	500MG/VIAL	N050758 001	Aug 12, 1998
-----------------------------------	--------------	------------	-------------	--------------

MYCOPHENOLIC ACID

TABLET, DELAYED RELEASE; ORAL MYFORTIC	NOVARTIS	180MG	N050791 001	Feb 27, 2004
	+	360MG	N050791 002	Feb 27, 2004

NABILONE

CAPSULE; ORAL CESAMET	+ MEDA PHARMS	1MG	N018677 001	Dec 26, 1985
--------------------------	---------------	-----	-------------	--------------

NABUMETONE

TABLET; ORAL <u>NABUMETONE</u>				
<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> DR REDDYS LABS LTD	<u>500MG</u>		<u>A078420 001</u>	Sep 24, 2008
<u>AB</u>	<u>750MG</u>		<u>A078420 002</u>	Sep 24, 2008
<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> MATRIX LABS LTD	<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> PAR PHARM	<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> 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> TEVA	<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> 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

PRESCRIPTION DRUG PRODUCT LIST

3 - 301 (of 424)

NADOLOL

TABLET; ORAL

CORGARD

<u>AB</u>	KING PHARMS	<u>20MG</u>	<u>N018063</u>	<u>005</u>	Oct 28, 1986
<u>AB</u>		<u>40MG</u>	<u>N018063</u>	<u>001</u>	
<u>AB</u>	+	<u>80MG</u>	<u>N018063</u>	<u>002</u>	
<u>NADOLOL</u>					
<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>	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

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>	ACIC FINE CHEMS	<u>EQ 1GM BASE</u>	<u>A090560</u>	<u>001</u>	Oct 03, 2011
<u>AP</u>		<u>EQ 2GM BASE</u>	<u>A090560</u>	<u>002</u>	Oct 03, 2011
<u>AP</u>	IBI	<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>	INSTITUTO BIOCHEMICO	<u>EQ 10GM BASE/VIAL</u>	<u>A090005</u>	<u>001</u>	Apr 20, 2011
<u>AP</u>	+	<u>SANDOZ</u>	<u>A062527</u>	<u>002</u>	Aug 02, 1984
<u>AP</u>	+		<u>A062732</u>	<u>001</u>	Dec 23, 1986
<u>AP</u>	+		<u>A062527</u>	<u>003</u>	Aug 02, 1984
<u>AP</u>	+		<u>A062732</u>	<u>002</u>	Dec 23, 1986
<u>AP</u>	+		<u>A062527</u>	<u>004</u>	Aug 02, 1984
NULLPEN IN PLASTIC CONTAINER					
+ BAXTER HLTHCARE	EQ 20MG BASE/ML	N050655	001	Oct 31, 1989	
+	EQ 2GM BASE/100ML	N050655	002	Oct 31, 1989	

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

+ MERZ PHARMS	1%	N019599	001	Feb 29, 1988
---------------	----	---------	-----	--------------

GEL; TOPICAL

NAFTIN

+ MERZ PHARMS	1%	N019356	001	Jun 18, 1990
---------------	----	---------	-----	--------------

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

<u>AP</u>	+	<u>HOSPIRA</u>	<u>10MG/ML</u>	<u>A070914</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>	+		<u>10MG/ML</u>	<u>A070915</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>	+		<u>20MG/ML</u>	<u>A070916</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>	+		<u>20MG/ML</u>	<u>A070918</u>	<u>001</u>	Feb 03, 1989

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HYDROCHLORIDE

<u>AP</u>	+	<u>HOSPIRA</u>	<u>0.4MG/ML</u>	<u>A070172</u>	<u>001</u>	Sep 24, 1986
-----------	---	----------------	-----------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 302 (of 424)

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HYDROCHLORIDE

<u>AP</u> + HOSPIRA	<u>0.4MG/ML</u>	<u>A070254</u> <u>001</u>	Jan 07, 1987
<u>AP</u> +	<u>0.4MG/ML</u>	<u>A070256</u> <u>001</u>	Jan 07, 1987
<u>AP</u> +	<u>0.4MG/ML</u>	<u>A070257</u> <u>001</u>	Jan 07, 1987
<u>AP</u> INTL MEDICATION	<u>0.4MG/ML</u>	<u>A070639</u> <u>001</u>	Sep 24, 1986
<u>AP</u> +	<u>1MG/ML</u>	<u>A072076</u> <u>001</u>	Mar 24, 1988

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

PENTAZOCINE AND NALOXONE HYDROCHLORIDE

<u>AB</u> GAVIS PHARMS	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>A075735</u> <u>001</u>	Jul 11, 2001
	<u>PENTAZOCINE AND NALOXONE HYDROCHLORIDES</u>		
<u>AB</u> RANBAXY	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>A075523</u> <u>001</u>	Mar 17, 2000

AB + WATSON LABSEQ 0.5MG BASE;EQ 50MG BASEA074736 001

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</u> <u>001</u>	Aug 17, 2011
<u>AB</u> BARR	<u>50MG</u>	<u>A074918</u> <u>001</u>	May 08, 1998
<u>AB</u> ELITE LABS	<u>50MG</u>	<u>A075274</u> <u>001</u>	May 26, 1999
<u>AB</u> MALLINCKRODT	<u>50MG</u>	<u>A076264</u> <u>002</u>	Mar 22, 2002
<u>AB</u> SANDOZ	<u>50MG</u>	<u>A075434</u> <u>001</u>	Mar 08, 2000
	<u>REVIA</u>		
<u>AB</u> + DURAMED	<u>50MG</u>	<u>N018932</u> <u>001</u>	Nov 20, 1984
	<u>NALTREXONE HYDROCHLORIDE</u>		
	MALLINCKRODT	25MG	A076264 001 Mar 22, 2002
		100MG	A076264 003 Mar 22, 2002

NANDROLONE DECANOATE

INJECTABLE; INJECTION

NANDROLONE DECANOATE

+ PHARMAFORCE

200MG/ML

A091252 001 Aug 30, 2010

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALBALONAT + ALLERGAN 0.1% A080248 001NAPHAZOLINE HYDROCHLORIDEAT TAYLOR 0.1% A083590 001NAPROXEN

SUSPENSION; ORAL

NAPROSYNAB + ROCHE PALO 25MG/ML N018965 001 Mar 23, 1987NAPROXENAB ROXANE 25MG/ML A074190 001 Mar 30, 1994

TABLET; ORAL

NAPROSYNAB ROCHE PALO 250MG N017581 002375MGN017581 003AB + 500MGN017581 004

Apr 15, 1982

PRESCRIPTION DRUG PRODUCT LIST

3 - 303 (of 424)

NAPROXEN

TABLET; ORAL

NAPROXEN

<u>AB</u>	AMNEAL PHARMS NY	<u>250MG</u>	<u>A075927</u>	<u>001</u>	Dec 18, 2001
<u>AB</u>		<u>375MG</u>	<u>A075927</u>	<u>002</u>	Dec 18, 2001
<u>AB</u>		<u>500MG</u>	<u>A075927</u>	<u>003</u>	Dec 18, 2001
<u>AB</u>	AUROBINDO PHARMA USA	<u>250MG</u>	<u>A200429</u>	<u>001</u>	Nov 08, 2011
<u>AB</u>		<u>375MG</u>	<u>A200429</u>	<u>002</u>	Nov 08, 2011
<u>AB</u>		<u>500MG</u>	<u>A200429</u>	<u>003</u>	Nov 08, 2011
<u>AB</u>	DAVA PHARMS INC	<u>250MG</u>	<u>A074410</u>	<u>001</u>	Apr 28, 1995
<u>AB</u>		<u>375MG</u>	<u>A074410</u>	<u>002</u>	Apr 28, 1995
<u>AB</u>		<u>500MG</u>	<u>A074410</u>	<u>003</u>	Apr 28, 1995
<u>AB</u>	GLENMARK GENERICS	<u>250MG</u>	<u>A078250</u>	<u>001</u>	Mar 28, 2007
<u>AB</u>		<u>375MG</u>	<u>A078250</u>	<u>002</u>	Mar 28, 2007
<u>AB</u>		<u>500MG</u>	<u>A078250</u>	<u>003</u>	Mar 28, 2007
<u>AB</u>	INVAGEN PHARMS	<u>250MG</u>	<u>A091305</u>	<u>001</u>	Aug 24, 2011
<u>AB</u>		<u>375MG</u>	<u>A091305</u>	<u>002</u>	Aug 24, 2011
<u>AB</u>		<u>500MG</u>	<u>A091305</u>	<u>003</u>	Aug 24, 2011
<u>AB</u>	MARKSANS PHARMA	<u>250MG</u>	<u>A091416</u>	<u>001</u>	Feb 14, 2011
<u>AB</u>		<u>375MG</u>	<u>A091416</u>	<u>002</u>	Feb 14, 2011
<u>AB</u>		<u>500MG</u>	<u>A091416</u>	<u>003</u>	Feb 14, 2011
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A074121</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>A074121</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>A074121</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>	PERRIGO R AND D	<u>250MG</u>	<u>A077339</u>	<u>001</u>	Apr 27, 2005
<u>AB</u>		<u>375MG</u>	<u>A077339</u>	<u>002</u>	Apr 27, 2005
<u>AB</u>		<u>500MG</u>	<u>A077339</u>	<u>003</u>	Apr 27, 2005
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A074140</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>A074140</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>A074140</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>	TEVA	<u>250MG</u>	<u>A074201</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>A074201</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>A074201</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A074457</u>	<u>001</u>	May 31, 1995
<u>AB</u>		<u>375MG</u>	<u>A074457</u>	<u>002</u>	May 31, 1995
<u>AB</u>		<u>500MG</u>	<u>A074457</u>	<u>003</u>	May 31, 1995
<u>AB</u>	WESTWARD	<u>250MG</u>	<u>A076494</u>	<u>001</u>	Jan 14, 2004
<u>AB</u>		<u>375MG</u>	<u>A076494</u>	<u>002</u>	Jan 14, 2004
<u>AB</u>		<u>500MG</u>	<u>A076494</u>	<u>003</u>	Jan 14, 2004
<u>AB</u>	ZYDUS PHARMS USA	<u>250MG</u>	<u>A078620</u>	<u>001</u>	Jun 07, 2007
<u>AB</u>		<u>375MG</u>	<u>A078620</u>	<u>002</u>	Jun 07, 2007
<u>AB</u>		<u>500MG</u>	<u>A078620</u>	<u>003</u>	Jun 07, 2007

TABLET, DELAYED RELEASE; ORAL

EC-NAPROSYN

<u>AB</u>	+ ROCHE PALO	<u>375MG</u>	<u>N020067</u>	<u>002</u>	Oct 14, 1994
<u>AB</u>	+	<u>500MG</u>	<u>N020067</u>	<u>003</u>	Oct 14, 1994

NAPROXEN

<u>AB</u>	ALPHAPHARM	<u>375MG</u>	<u>A075390</u>	<u>001</u>	Apr 19, 2001
<u>AB</u>		<u>500MG</u>	<u>A075390</u>	<u>002</u>	Apr 19, 2001
<u>AB</u>	INVAGEN PHARMS	<u>375MG</u>	<u>A091432</u>	<u>001</u>	Sep 19, 2011
<u>AB</u>		<u>500MG</u>	<u>A091432</u>	<u>002</u>	Sep 19, 2011
<u>AB</u>	PLIVA	<u>375MG</u>	<u>A075337</u>	<u>001</u>	May 26, 1999
<u>AB</u>		<u>500MG</u>	<u>A075337</u>	<u>002</u>	May 26, 1999
<u>AB</u>	SANDOZ	<u>375MG</u>	<u>A075061</u>	<u>001</u>	Feb 18, 1998
<u>AB</u>		<u>500MG</u>	<u>A075061</u>	<u>002</u>	Feb 18, 1998
<u>AB</u>	TEVA	<u>375MG</u>	<u>A075227</u>	<u>001</u>	Jun 30, 1998
<u>AB</u>		<u>500MG</u>	<u>A075227</u>	<u>002</u>	Jun 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 304 (of 424)

NAPROXEN SODIUM

TABLET; ORAL

ANAPROX

<u>AB</u>	ROCHE PALO	<u>EQ 250MG BASE</u>	<u>N018164 001</u>	
<u>AB</u>	<u>ANAPROX DS</u>			
<u>AB</u>	+ ROCHE PALO	<u>EQ 500MG BASE</u>	<u>N018164 003</u>	Sep 30, 1987
<u>AB</u>	<u>NAPROXEN SODIUM</u>			
<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 250MG BASE</u>	<u>A078432 001</u>	Apr 25, 2007
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A078432 002</u>	Apr 25, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A200629 001</u>	Oct 31, 2011
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A200629 002</u>	Oct 31, 2011
<u>AB</u>	DR REDDYS LABS LTD	<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 GENERICS	<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>	HIKMA	<u>EQ 250MG BASE</u>	<u>A074480 002</u>	Feb 18, 1998
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A074480 001</u>	May 14, 1996
<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
<u>AB</u>	WATSON LABS	<u>EQ 250MG BASE</u>	<u>A074455 001</u>	May 31, 1995
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A074455 002</u>	May 31, 1995

TABLET, EXTENDED RELEASE; ORAL

NAPRELAN

	STAT TRADE	<u>EQ 375MG BASE</u>	<u>N020353 001</u>	Jan 05, 1996
		<u>EQ 500MG BASE</u>	<u>N020353 002</u>	Jan 05, 1996
+		<u>EQ 750MG BASE</u>	<u>N020353 003</u>	Jan 05, 1996

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET; ORAL

TREXIMET

+	GLAXOSMITHKLINE	500MG;EQ 85MG BASE	<u>N021926 001</u>	Apr 15, 2008
---	-----------------	--------------------	--------------------	--------------

NARATRIPTAN

TABLET; ORAL

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>	SUN PHARM INDs LTD	<u>EQ 2.5MG BASE</u>	<u>A091552 001</u>	Feb 14, 2011

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

AMERGE

<u>AB</u>	GLAXOSMITHKLINE	<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>	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>	PADDICK 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>	ROXANE	<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
<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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 305 (of 424)

NATAMYCIN

SUSPENSION; OPHTHALMIC

NATACYN

+ ALCON

5%

N050514 001

NATEGLINIDE

TABLET; ORAL

NATEGLINIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>60MG</u>	<u>A077461</u> <u>001</u>	Sep 09, 2009
<u>AB</u>		<u>120MG</u>	<u>A077461</u> <u>002</u>	Sep 09, 2009
<u>AB</u>	PAR PHARM	<u>60MG</u>	<u>A077463</u> <u>001</u>	Sep 09, 2009
<u>AB</u>		<u>120MG</u>	<u>A077463</u> <u>002</u>	Sep 09, 2009
<u>AB</u>	TEVA PHARMS	<u>60MG</u>	<u>A077467</u> <u>001</u>	Sep 09, 2009
<u>AB</u>		<u>120MG</u>	<u>A077467</u> <u>002</u>	Sep 09, 2009
<u>AB</u>	WATSON LABS	<u>60MG</u>	<u>A077462</u> <u>001</u>	Mar 30, 2011
<u>AB</u>		<u>120MG</u>	<u>A077462</u> <u>002</u>	Mar 30, 2011
		<u>STARLIX</u>		
<u>AB</u>	NOVARTIS	<u>60MG</u>	<u>N021204</u> <u>001</u>	Dec 22, 2000
<u>AB</u> +		<u>120MG</u>	<u>N021204</u> <u>002</u>	Dec 22, 2000

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

NEDOCROMIL SODIUM

SOLUTION/DROPS; OPHTHALMIC

ALOCRIL

+ ALLERGAN

2%

N021009 001 Dec 08, 1999

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

<u>AB</u>	RANBAXY	<u>50MG</u>	<u>A076409</u> <u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>A076409</u> <u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>A076409</u> <u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>A076409</u> <u>004</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>A076409</u> <u>005</u>	Sep 16, 2003
<u>AB</u>	TEVA	<u>50MG</u>	<u>A076037</u> <u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>A076037</u> <u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>A076037</u> <u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>A076037</u> <u>004</u>	Sep 16, 2003
<u>AB</u> +		<u>250MG</u>	<u>A076037</u> <u>005</u>	Sep 16, 2003
		NEFAZODONE HYDROCHLORIDE		
BX	DR REDDYS LABS INC	50MG	A076309 001	Sep 16, 2003
BX		100MG	A076309 002	Sep 16, 2003
BX		150MG	A076309 003	Sep 16, 2003
BX		200MG	A076309 004	Sep 16, 2003
BX		250MG	A076309 005	Sep 16, 2003

NELARABINE

INJECTABLE; IV (INFUSION)

+ SMITHKLINE BEECHAM 250MG/50ML (5MG/ML)

N021877 001 Oct 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 306 (of 424)

NELFINAVIR MESYLATE

POWDER; ORAL VIRACEPT + AGOURON	EQ 50MG BASE/SCOOPFUL	N020778 001	Mar 14, 1997
TABLET; ORAL VIRACEPT + AGOURON	EQ 250MG BASE	N020779 001	Mar 14, 1997
	EQ 625MG BASE	N021503 001	Apr 30, 2003

NEOMYCIN SULFATE

POWDER; FOR RX COMPOUNDING NEO-RX X GEN PHARMS	100%	A061579 001	
SOLUTION; ORAL NEO-FRADIN + X GEN PHARMS	EQ 87.5MG BASE/5ML	A065010 001	May 23, 2002
TABLET; ORAL NEOMYCIN SULFATE <u>AA</u> OMAN PHARM PRODUCTS 500MG		<u>A065468 001</u>	Mar 29, 2010
<u>AA</u> + TEVA 500MG		<u>A060304 001</u>	
<u>AA</u> X GEN PHARMS 500MG		<u>A065220 001</u>	Jul 28, 2006

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION NEOMYCIN AND POLYMYXIN B SULFATE <u>AT</u> WATSON LABS EQ 40MG BASE/ML;200,000 UNITS/ML		<u>A062664 001</u>	Apr 08, 1986
<u>AT</u> X GEN PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML		<u>A065106 001</u>	Jan 31, 2006
<u>AT</u> EQ 40MG BASE/ML;200,000 UNITS/ML		<u>A065108 001</u>	Jan 31, 2006
NEOSPORIN G.U. IRRIGANT			
<u>AT</u> + MONARCH PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML		<u>A060707 001</u>	

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	
---	------------------------------------	-------------	--

NEPAFENAC

SUSPENSION/DROPS; OPHTHALMIC + ALCON PHARMS LTD	0.1%	N021862 001	Aug 19, 2005
--	------	-------------	--------------

NESIRITIDE RECOMBINANT

FOR SOLUTION; INTRAVENOUS NATRECOR + SCIOS	1.5MG/VIAL	N020920 001	Aug 10, 2001
--	------------	-------------	--------------

NEVIRAPINE

SUSPENSION; ORAL VIRAMUNE + BOEHRINGER INGELHEIM	50MG/5ML	N020933 001	Sep 11, 1998
TABLET; ORAL VIRAMUNE + BOEHRINGER INGELHEIM	200MG	N020636 001	Jun 21, 1996
TABLET, EXTENDED RELEASE; ORAL VIRAMUNE XR + BOEHRINGER INGELHEIM	400MG	N201152 001	Mar 25, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 307 (of 424)

NIACIN

TABLET; ORAL

NIACIN

<u>AA</u>	WOCKHARDT	<u>500MG</u>	<u>A081134 001</u>	Apr 28, 1992
-----------	-----------	--------------	--------------------	--------------

NIACOR

<u>AA</u>	+ UPSHER SMITH	<u>500MG</u>	<u>A040378 001</u>	May 03, 2000
-----------	----------------	--------------	--------------------	--------------

TABLET, EXTENDED RELEASE; ORAL

NIASPAN

ABBOTT	500MG	N020381 002	Jul 28, 1997
+	750MG	N020381 003	Jul 28, 1997
+	1GM	N020381 004	Jul 28, 1997

NIACIN; SIMVASTATIN

TABLET, EXTENDED RELEASE; ORAL

SIMCOR

+ ABBOTT	500MG;20MG	N022078 001	Feb 15, 2008
+	500MG;40MG	N022078 004	Jul 28, 2010
+	750MG;20MG	N022078 002	Feb 15, 2008
+	1GM;20MG	N022078 003	Feb 15, 2008
+	1GM;40MG	N022078 005	Jul 28, 2010

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

CARDENE

<u>AB</u>	EKR THERAP	<u>20MG</u>	<u>N019488 001</u>	Dec 21, 1988
<u>AB</u>	+	<u>30MG</u>	<u>N019488 002</u>	Dec 21, 1988

NICARDIPINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>20MG</u>	<u>A074439 001</u>	Dec 10, 1996
<u>AB</u>		<u>30MG</u>	<u>A074439 002</u>	Dec 10, 1996
<u>AB</u>	EPIC PHARMA	<u>20MG</u>	<u>A074928 001</u>	Mar 19, 1998
<u>AB</u>		<u>30MG</u>	<u>A074928 002</u>	Mar 19, 1998
<u>AB</u>	MYLAN	<u>20MG</u>	<u>A074642 001</u>	Jul 18, 1996
<u>AB</u>		<u>30MG</u>	<u>A074642 002</u>	Jul 18, 1996
<u>AB</u>	TEVA	<u>20MG</u>	<u>A074540 001</u>	Oct 28, 1996
<u>AB</u>		<u>30MG</u>	<u>A074540 002</u>	Oct 28, 1996
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>A074670 001</u>	Oct 28, 1996
<u>AB</u>		<u>30MG</u>	<u>A074670 002</u>	Oct 28, 1996

CAPSULE, EXTENDED RELEASE; ORAL

CARDENE SR

+	EKR THERAP	30MG	N020005 001	Feb 21, 1992
+		60MG	N020005 003	Feb 21, 1992

INJECTABLE; INJECTION

CARDENE

<u>AP</u>	+	EKR THERAP	<u>25MG/10ML (2.5MG/ML)</u>	<u>N019734 001</u>	Jan 30, 1992
-----------	---	------------	-----------------------------	--------------------	--------------

NICARDIPINE HYDROCHLORIDE

<u>AP</u>	BEDFORD	<u>25MG/10ML (2.5MG/ML)</u>	<u>A078714 001</u>	Dec 28, 2009
<u>AP</u>	BIONICHE PHARMA USA	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090664 001</u>	Nov 17, 2009
<u>AP</u>	EXELA PHARMA SCIENCE	<u>25MG/10ML (2.5MG/ML)</u>	<u>N022276 001</u>	Jul 24, 2008
<u>AP</u>	NAVINTA LLC	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090125 001</u>	Nov 17, 2009
<u>AP</u>	PHARMAFORCE	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090534 001</u>	Nov 17, 2009
<u>AP</u>	SUN PHARMA GLOBAL	<u>25MG/10ML (2.5MG/ML)</u>	<u>N078405 001</u>	Nov 17, 2009
<u>AP</u>	WOCKHARDT	<u>25MG/10ML (2.5MG/ML)</u>	<u>A090671 001</u>	Nov 17, 2009

INJECTABLE; INTRAVENOUS

CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER

+	EKR THERAP	40MG/200ML (0.2MG/ML)	N019734 004	Nov 07, 2008
---	------------	-----------------------	-------------	--------------

CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER

+	EKR THERAP	20MG/200ML (0.1MG/ML)	N019734 003	Jul 31, 2008
---	------------	-----------------------	-------------	--------------

CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER

+	EKR THERAP	20MG/200ML (0.1MG/ML)	N019734 002	Jul 31, 2008
---	------------	-----------------------	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 308 (of 424)

NICARDIPINE HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER
+ EKR THERAP 40MG/200ML (0.2MG/ML)

N019734 005 Nov 07, 2008

NICOTINE

INHALANT; ORAL

NICOTROL
+ PHARMACIA AND UPJOHN 4MG/CARTRIDGE

N020714 001 May 02, 1997

SPRAY, METERED; NASAL

NICOTROL
+ PFIZER INC 0.5MG/SPRAY

N020385 001 Mar 22, 1996

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072579</u> <u>001</u>	Jan 08, 1991
<u>AB</u>	CATALENT	<u>10MG</u>	<u>A073250</u> <u>001</u>	Oct 08, 1991
<u>AB</u>	INTERGEL PHARM	<u>10MG</u>	<u>A072781</u> <u>001</u>	Jul 30, 1993
<u>AB</u>	<u>PROCARDIA</u>			
<u>AB</u> +	PFIZER	<u>10MG</u>	<u>N018482</u> <u>001</u>	
	NIFEDIPINE			
	ACTAVIS ELIZABETH	20MG	A072556 001	Sep 20, 1990

TABLET, EXTENDED RELEASE; ORAL

ADALAT CC

<u>AB1</u>	BAYER HLTHCARE	<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

AFEDITAB CR

<u>AB1</u>	WATSON LABS	<u>30MG</u>	<u>A075128</u> <u>001</u>	Mar 10, 2000
<u>AB1</u>		<u>60MG</u>	<u>A075659</u> <u>001</u>	Oct 26, 2001

NIFEDIPINE

<u>AB1</u>	ACTAVIS	<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>	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>	VALEANT INTL	<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

NIFEDIPINE

<u>AB2</u>	MATRIX LABS LTD	<u>30MG</u>	<u>A090602</u> <u>001</u>	Sep 13, 2010
<u>AB2</u>		<u>60MG</u>	<u>A090602</u> <u>002</u>	Sep 13, 2010
<u>AB2</u>		<u>90MG</u>	<u>A090602</u> <u>003</u>	Sep 13, 2010
<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	<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>	VALEANT INTL	<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

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

PRESCRIPTION DRUG PRODUCT LIST

3 - 309 (of 424)

NILOTINIB HYDROCHLORIDE MONOHYDRATE

CAPSULE; ORAL TASIGNA NOVARTIS +	EQ 150MG BASE EQ 200MG BASE	N022068 002 N022068 001	Jun 17, 2010 Oct 29, 2007
--	--------------------------------	----------------------------	------------------------------

NILUTAMIDE

TABLET; ORAL NILANDRON + SANOFI AVENTIS US	150MG	N020169 002	Apr 30, 1999
--	-------	-------------	--------------

NIMODIPINE

CAPSULE; ORAL NIMODIPINE AB BANNER PHARMACAPS AB BARR LABS INC AB + SUN PHARM INDs INC	<u>30MG</u> <u>30MG</u> <u>30MG</u>	<u>A076740 001</u> <u>A077811 001</u> <u>A077067 001</u>	Jan 17, 2008 May 02, 2007 Apr 17, 2007
--	---	--	--

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL NISOLDIPINE AB MYLAN AB AB AB SULAR AB + SHIONOGI INC AB + AB AB + NISOLDIPINE + MYLAN + +	<u>8 .5MG</u> <u>17MG</u> <u>25 .5MG</u> <u>34MG</u> <u>8 .5MG</u> <u>17MG</u> <u>25 .5MG</u> <u>34MG</u> 20MG 30MG 40MG	<u>A091001 001</u> <u>A091001 002</u> <u>A091001 003</u> <u>A091001 004</u> <u>N020356 008</u> <u>N020356 007</u> <u>N020356 006</u> <u>N020356 005</u> A079051 001 A079051 002 A079051 003	Jan 26, 2011 Jan 26, 2011 Jan 26, 2011 Jan 26, 2011 Jan 02, 2008 Jan 02, 2008 Jan 02, 2008 Jan 02, 2008 Jul 25, 2008 Jul 25, 2008 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 RARE DIS	2MG 5MG 10MG	N021232 001 N021232 002 N021232 003	Jan 18, 2002 Jan 18, 2002 Jan 18, 2002
--------------------------------------	--------------------	---	--

NITRIC OXIDE

GAS; INHALATION INOMAX INO +	100PPM 800PPM	N020845 002 N020845 003	Dec 23, 1999 Dec 23, 1999
------------------------------------	------------------	----------------------------	------------------------------

PREScription DRUG PRODUCT LIST

3 - 310 (of 424)

NITROFURANTOIN

SUSPENSION; ORAL

<u>AB</u>	<u>FURADANTIN</u> + SHIONOGI INC	<u>25MG/5ML</u>	<u>N009175</u>	<u>001</u>	
<u>AB</u>	<u>NITROFURANTOIN</u> AMNEAL PHARMS	<u>25MG/5ML</u>	<u>A201679</u>	<u>001</u>	May 11, 2011

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

<u>AB</u>	ALVOGEN	<u>25MG</u>	<u>N016620</u>	<u>003</u>	
<u>AB</u>		<u>50MG</u>	<u>N016620</u>	<u>001</u>	
<u>AB</u>	+	<u>100MG</u>	<u>N016620</u>	<u>002</u>	
	<u>NITROFURANTOIN</u>				
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>50MG</u>	<u>A073671</u>	<u>001</u>	Jan 28, 1993
<u>AB</u>		<u>100MG</u>	<u>A073652</u>	<u>001</u>	Jan 28, 1993
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A074967</u>	<u>001</u>	Jul 09, 1997
<u>AB</u>		<u>100MG</u>	<u>A077025</u>	<u>001</u>	Aug 18, 2004
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A073696</u>	<u>001</u>	Dec 31, 1992
<u>AB</u>		<u>50MG</u>	<u>A073696</u>	<u>002</u>	Dec 31, 1992
<u>AB</u>		<u>100MG</u>	<u>A073696</u>	<u>003</u>	Dec 31, 1992

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

<u>AB</u>	<u>+</u>	ALVOGEN	<u>75MG;25MG</u>	<u>N020064</u>	<u>001</u>	Dec 24, 1991
<u>NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)</u>						
<u>AB</u>		MYLAN	<u>75MG;25MG</u>	<u>A076648</u>	<u>001</u>	Mar 22, 2004
<u>AB</u>		RANBAXY	<u>75MG;25MG</u>	<u>A076951</u>	<u>001</u>	Mar 30, 2005
<u>AB</u>		SANDOZ	<u>75MG;25MG</u>	<u>A077066</u>	<u>001</u>	Apr 05, 2005

NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST

+ NOVADEL 0.4MG/SPRAY

N021780 001 Nov 02, 2006

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

<u>AB1</u>	GRACEWAY	<u>0.1MG/HR</u>	<u>A089771</u>	<u>001</u>	Aug 30, 1996
<u>AB1</u>		<u>0.2MG/HR</u>	<u>A089772</u>	<u>001</u>	Aug 30, 1996
<u>AB1</u>		<u>0.4MG/HR</u>	<u>A089773</u>	<u>001</u>	Aug 30, 1996
<u>AB1</u>		<u>0.6MG/HR</u>	<u>A089774</u>	<u>001</u>	Aug 30, 1996

NITRO-DUR

<u>AB1</u> + KEY PHARMS	<u>0.1MG/HR</u>	<u>N020145</u> <u>001</u>	Apr 04, 1995
<u>AB1</u> +	<u>0.2MG/HR</u>	<u>N020145</u> <u>002</u>	Apr 04, 1995
<u>AB1</u> +	<u>0.4MG/HR</u>	<u>N020145</u> <u>004</u>	Apr 04, 1995
<u>AB1</u> +	<u>0.6MG/HR</u>	<u>N020145</u> <u>005</u>	Apr 04, 1995

NITROGLYCERTIN

AB1 KREMERS URBAN PHARMS 0 .2MG/HR A075115 001 Aug 10, 2004
AB1 0 .4MG/HR A075115 002 Aug 10, 2004

NITROGLYCERIN

<u>AB2</u>	HERCON LABS	<u>0 . 2MG / HR</u>	<u>A089884</u>	<u>001</u>	Oct 30, 1998
<u>AB2</u>		<u>0 . 4MG / HR</u>	<u>A089885</u>	<u>001</u>	Oct 30, 1998
<u>AB2</u>		<u>0 . 6MG / HR</u>	<u>A089886</u>	<u>001</u>	Oct 30, 1998
<u>AB2 +</u>	MYLAN TECHNOLOGIES	<u>0 . 2MG / HR</u>	<u>A074559</u>	<u>003</u>	Aug 30, 1996

2 +

<u>2</u> +	<u>0.6MG/HR</u>	<u>A074559</u>	<u>001</u>	Aug 30, 1996
NITRO-DUR				
+ KEY PHARMS	0.3MG/HR	N020145	003	Apr 04, 1995
	0.8MG/HR	N020145	006	Apr 04, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 311 (of 424)

NITROGLYCERIN

FILM, EXTENDED RELEASE; TRANSDERMAL			
NITROGLYCERIN			
+ MYLAN TECHNOLOGIES	0.1MG/HR	A074559 004	Feb 06, 1998
INJECTABLE; INJECTION			
<u>NITROGLYCERIN</u>			
AP + HOSPIRA	<u>5MG/ML</u>	<u>N018531 001</u>	
AP + LUITPOLD	<u>5MG/ML</u>	<u>A072034 001</u>	May 24, 1988
<u>NITROGLYCERIN IN DEXTROSE 5%</u>			
AP + BAXTER HLTHCARE	<u>10MG/100ML</u>	<u>N019970 001</u>	Dec 29, 1989
AP +	<u>20MG/100ML</u>	<u>N019970 002</u>	Dec 29, 1989
AP +	<u>40MG/100ML</u>	<u>N019970 003</u>	Dec 29, 1989
AP HOSPIRA	<u>10MG/100ML</u>	<u>A071846 001</u>	Aug 31, 1990
AP	<u>20MG/100ML</u>	<u>A071847 001</u>	Aug 31, 1990
AP	<u>40MG/100ML</u>	<u>A071848 001</u>	Aug 31, 1990
OINTMENT; INTRA-ANAL			
RECTIV			
+ PROSTRAKAN INC	0.4%	N021359 001	Jun 21, 2011
OINTMENT; TRANSDERMAL			
NITROGLYCERIN			
+ FOUGERA	2%	A087355 001	Jul 08, 1988
SPRAY, METERED; SUBLINGUAL			
NITROLINGUAL PUMPSPRAY			
+ POHL BOSKAMP	0.4MG/SPRAY	N018705 002	Jan 10, 1997
TABLET; SUBLINGUAL			
NITROSTAT			
PFIZER PHARMS	0.3MG	N021134 001	May 01, 2000
	0.4MG	N021134 002	May 01, 2000
+	0.6MG	N021134 003	May 01, 2000

NIZATIDINE

CAPSULE; ORAL			
<u>AXID</u>			
AB SMITHKLINE BEECHAM	<u>150MG</u>	<u>N019508 001</u>	Apr 12, 1988
AB +	<u>300MG</u>	<u>N019508 002</u>	Apr 12, 1988
<u>NIZATIDINE</u>			
AB APOTEX	<u>150MG</u>	<u>A076383 001</u>	Jan 23, 2003
AB	<u>300MG</u>	<u>A076383 002</u>	Jan 23, 2003
AB DR REDDYS LABS LTD	<u>150MG</u>	<u>A077314 001</u>	Sep 15, 2005
AB	<u>300MG</u>	<u>A077314 002</u>	Sep 15, 2005
AB GLENMARK GENERICS	<u>150MG</u>	<u>A090618 001</u>	Jul 15, 2011
AB	<u>300MG</u>	<u>A090618 002</u>	Jul 15, 2011
AB MYLAN	<u>150MG</u>	<u>A075806 001</u>	Jul 05, 2002
AB	<u>150MG</u>	<u>A075934 001</u>	Jul 09, 2002
AB	<u>300MG</u>	<u>A075806 002</u>	Jul 05, 2002
AB	<u>300MG</u>	<u>A075934 002</u>	Jul 09, 2002
AB SANDOZ	<u>150MG</u>	<u>A076178 001</u>	Jul 05, 2002
AB	<u>300MG</u>	<u>A076178 002</u>	Jul 05, 2002
AB TEVA	<u>150MG</u>	<u>A075668 001</u>	Sep 12, 2002
AB	<u>300MG</u>	<u>A075668 002</u>	Sep 12, 2002
AB WATSON LABS	<u>150MG</u>	<u>A075616 001</u>	Jul 09, 2002
AB	<u>300MG</u>	<u>A075616 002</u>	Jul 09, 2002
SOLUTION; ORAL			
<u>AXID</u>			
AA + BRAINTREE	<u>15MG/ML</u>	<u>N021494 001</u>	May 25, 2004
<u>NIZATIDINE</u>			
AA AMNEAL PHARMS	<u>15MG/ML</u>	<u>A090576 001</u>	Nov 18, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 312 (of 424)

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

<u>LEVOPHED</u>			
AP + HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N007513 001</u>	
	<u>NOREPINEPHRINE BITARTRATE</u>		
AP BEDFORD	<u>EQ 1MG BASE/ML</u>	<u>A040462 001</u>	Oct 31, 2003
AP TEVA PARENTERAL	<u>EQ 1MG BASE/ML</u>	<u>A040455 001</u>	Mar 03, 2003

NORETHINDRONE

TABLET; ORAL-28

<u>CAMILA</u>			
AB1 BARR	<u>0.35MG</u>	<u>A076177 001</u>	Oct 21, 2002
<u>HEATHER</u>			
AB1 GLENMARK GENERICS	<u>0.35MG</u>	<u>A090454 001</u>	Apr 23, 2010
<u>NORETHINDRONE</u>			
AB1 LUPIN LTD	<u>0.35MG</u>	<u>A091325 001</u>	Sep 19, 2011
<u>NOR-QD</u>			
AB1 + WATSON LABS (UTAH)	<u>0.35MG</u>	<u>N017060 001</u>	
<u>ERRIN</u>			
AB2 BARR	<u>0.35MG</u>	<u>A076225 001</u>	Oct 21, 2002
<u>MICRONOR</u>			
AB2 + JANSSEN PHARMS	<u>0.35MG</u>	<u>N016954 001</u>	
<u>NORETHIDRONE</u>			
AB2 GLENMARK GENERICS	<u>0.35MG</u>	<u>A091209 001</u>	Jul 21, 2010

NORETHINDRONE ACETATE

TABLET; ORAL

<u>AYGESTIN</u>			
AB + DURAMED RES	<u>5MG</u>	<u>N018405 001</u>	Apr 21, 1982
<u>NORETHINDRONE ACETATE</u>			
AB BARR	<u>5MG</u>	<u>A075951 001</u>	May 25, 2001
AB GLENMARK GENERICS	<u>5MG</u>	<u>A091090 001</u>	Jul 21, 2010

NORFLOXACIN

TABLET; ORAL

NOROXIN

+ MERCK

400MG

N019384 002 Oct 31, 1986

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

<u>NORTRIPTYLINE HYDROCHLORIDE</u>			
AB MYLAN	<u>EQ 10MG BASE</u>	<u>A074234 001</u>	Jul 26, 1993
	<u>EQ 25MG BASE</u>	<u>A074234 002</u>	Jul 26, 1993
	<u>EQ 50MG BASE</u>	<u>A074234 003</u>	Jul 26, 1993
	<u>EQ 75MG BASE</u>	<u>A074234 004</u>	Jul 26, 1993
AB TARO	<u>EQ 10MG BASE</u>	<u>A075520 004</u>	May 08, 2000
	<u>EQ 25MG BASE</u>	<u>A075520 003</u>	May 08, 2000
	<u>EQ 50MG BASE</u>	<u>A075520 001</u>	May 08, 2000
	<u>EQ 75MG BASE</u>	<u>A075520 002</u>	May 08, 2000
AB TEVA	<u>EQ 10MG BASE</u>	<u>A074132 001</u>	Mar 27, 1995
	<u>EQ 25MG BASE</u>	<u>A074132 002</u>	Mar 27, 1995
	<u>EQ 50MG BASE</u>	<u>A074132 003</u>	Mar 27, 1995
	<u>EQ 75MG BASE</u>	<u>A074132 004</u>	Mar 27, 1995
AB WATSON LABS	<u>EQ 10MG BASE</u>	<u>A073553 001</u>	Mar 30, 1992
	<u>EQ 25MG BASE</u>	<u>A073554 001</u>	Mar 30, 1992
	<u>EQ 50MG BASE</u>	<u>A073555 001</u>	Mar 30, 1992
	<u>EQ 75MG BASE</u>	<u>A073556 001</u>	Mar 30, 1992
<u>PAMELOR</u>			
AB MALLINCKRODT LLC	<u>EQ 10MG BASE</u>	<u>N018013 001</u>	
	<u>EQ 25MG BASE</u>	<u>N018013 002</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 313 (of 424)

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

PAMELOR

<u>AB</u>	MALLINCKRODT LLC	<u>EQ 50MG BASE</u>	<u>N018013 004</u>
<u>AB</u>	+	<u>EQ 75MG BASE</u>	<u>N018013 003</u>

SOLUTION; ORAL

AVENTYL HYDROCHLORIDE

<u>AA</u>	+	RANBAXY	<u>EQ 10MG BASE/5ML</u>	<u>N014685 001</u>
-----------	---	---------	-------------------------	--------------------

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

NYSTATIN

CREAM; TOPICAL

NYSTATIN

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062949 001</u>	Jun 13, 1988
<u>AT</u>	ALTANA	<u>100,000 UNITS/GM</u>	<u>A062129 001</u>	
<u>AT</u>	PERRIGO NEW YORK	<u>100,000 UNITS/GM</u>	<u>A062225 001</u>	
<u>AT</u>	TARO	<u>100,000 UNITS/GM</u>	<u>A064022 001</u>	Jan 29, 1993
<u>AT</u>	VINTAGE	<u>100,000 UNITS/GM</u>	<u>A065315 001</u>	May 31, 2006

OINTMENT; TOPICAL

NYSTATIN

<u>AT</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<u>A062840 001</u>	Nov 13, 1987
<u>AT</u>	+	ALTANA	<u>100,000 UNITS/GM</u>	<u>A062124 002</u>
<u>AT</u>	PERRIGO NEW YORK	<u>100,000 UNITS/GM</u>	<u>A062472 001</u>	Feb 13, 1984

POWDER; TOPICAL

NYSTATIN

<u>AT</u>	+	COASTAL PHARMS	<u>100,000 UNITS/GM</u>	<u>A065203 001</u>	Jul 15, 2004
<u>AT</u>		NESHER PHARMS	<u>100,000 UNITS/GM</u>	<u>A065321 001</u>	Aug 18, 2006
<u>AT</u>		PAR PHARM	<u>100,000 UNITS/GM</u>	<u>A065138 001</u>	Jul 23, 2004
<u>AT</u>		UPSHER SMITH	<u>100,000 UNITS/GM</u>	<u>A065183 001</u>	May 03, 2005
<u>AT</u>		X GEN PHARMS	<u>100,000 UNITS/GM</u>	<u>A065175 001</u>	Dec 17, 2004

NYSTOP

<u>AT</u>	PADDOCK LLC	<u>100,000 UNITS/GM</u>	<u>A064118 001</u>	Aug 16, 1996
-----------	-------------	-------------------------	--------------------	--------------

SUSPENSION; ORAL

NILSTAT

<u>AA</u>	+	GLENMARK GENERICS	<u>100,000 UNITS/ML</u>	<u>N050299 001</u>
-----------	---	-------------------	-------------------------	--------------------

NYSTATIN

<u>AA</u>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/ML</u>	<u>A062349 001</u>	Jul 14, 1982	
<u>AA</u>	FOUGERA	<u>100,000 UNITS/ML</u>	<u>A062517 001</u>	Jun 07, 1984	
<u>AA</u>	HI TECH PHARMA	<u>100,000 UNITS/ML</u>	<u>A064042 001</u>	Feb 28, 1994	
<u>AA</u>	TARO	<u>100,000 UNITS/ML</u>	<u>A062876 001</u>	Feb 29, 1988	
<u>AA</u>	VINTAGE PHARMS	<u>100,000 UNITS/ML</u>	<u>A065148 001</u>	Jun 28, 2005	
<u>AA</u>	VISTAPHARM	<u>100,000 UNITS/ML</u>	<u>A064142 001</u>	Jun 25, 1998	
<u>AA</u>		<u>100,000 UNITS/ML</u>	<u>A065422 001</u>	Mar 07, 2011	
<u>AA</u>	+	WOCKHARDT	<u>100,000 UNITS/ML</u>	<u>A062512 001</u>	Oct 29, 1984

TABLET; ORAL

NYSTATIN

<u>AA</u>	MUTUAL PHARM	<u>500,000 UNITS</u>	<u>A062838 001</u>	Dec 22, 1988
<u>AA</u>	+	TEVA	<u>500,000 UNITS</u>	<u>A062506 001</u>

TABLET; VAGINAL

NYSTATIN

+	ODYSSEY PHARMS	100,000 UNITS	A062615 001	Oct 17, 1985
---	----------------	---------------	-------------	--------------

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>100,000 UNITS/GM;0.1%</u>	<u>A062599 001</u>	Oct 08, 1985
<u>AT</u>	+	TARO	<u>100,000 UNITS/GM;0.1%</u>	<u>A062364 001</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 314 (of 424)

NYSTATIN; TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

NYSTATIN AND TRIAMCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>100,000 UNITS/GM; 0.1%</u>	<u>A062602</u>	<u>001</u>	Oct 09, 1985	
<u>AT</u>	+	<u>TARO</u>	<u>100,000 UNITS/GM; 0.1%</u>	<u>A063305</u>	<u>001</u>	Mar 29, 1993

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

<u>AP</u>	APP PHARMS	<u>EQ 0.2MG BASE/ML</u>	<u>A077450</u>	<u>001</u>	Feb 10, 2006	
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A077450</u>	<u>002</u>	Feb 10, 2006	
<u>AP</u>	+	<u>BEDFORD</u>	<u>A076330</u>	<u>001</u>	Apr 08, 2005	
<u>AP</u>	+		<u>A076330</u>	<u>002</u>	Apr 08, 2005	
<u>AP</u>	SUN PHARM IND'S	<u>EQ 0.05MG BASE/ML</u>	<u>A077329</u>	<u>001</u>	Mar 04, 2008	
<u>AP</u>		<u>EQ 0.05MG BASE/ML</u>	<u>A077372</u>	<u>001</u>	Aug 14, 2007	
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A077329</u>	<u>002</u>	Mar 04, 2008	
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A077372</u>	<u>002</u>	Aug 14, 2007	
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A077330</u>	<u>001</u>	Mar 04, 2008	
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A077373</u>	<u>001</u>	Aug 14, 2007	
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A077329</u>	<u>003</u>	Mar 04, 2008	
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A077372</u>	<u>003</u>	Aug 14, 2007	
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A077331</u>	<u>001</u>	Mar 04, 2008	
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A077373</u>	<u>002</u>	Aug 14, 2007	
<u>AP</u>	TEVA PARENTERAL	<u>EQ 0.05MG BASE/ML</u>	<u>A075957</u>	<u>001</u>	Oct 03, 2005	
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A075957</u>	<u>002</u>	Oct 03, 2005	
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>A075959</u>	<u>001</u>	Nov 21, 2005	
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A075957</u>	<u>003</u>	Oct 03, 2005	
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075959</u>	<u>002</u>	Nov 21, 2005	
<u>AP</u>	WOCHARDT USA	<u>EQ 0.2MG BASE/ML</u>	<u>A090986</u>	<u>001</u>	May 11, 2011	
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A090986</u>	<u>002</u>	May 11, 2011	
		<u>OCTREOTIDE ACETATE (PRESERVATIVE FREE)</u>				
<u>AP</u>	APP PHARMS	<u>EQ 0.05MG BASE/ML</u>	<u>A077457</u>	<u>001</u>	Feb 10, 2006	
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A077457</u>	<u>002</u>	Feb 10, 2006	
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A077457</u>	<u>003</u>	Feb 10, 2006	
<u>AP</u>	+	<u>BEDFORD</u>	<u>A076313</u>	<u>001</u>	Mar 28, 2005	
<u>AP</u>	+		<u>A076313</u>	<u>003</u>	Mar 28, 2005	
<u>AP</u>	+		<u>A076313</u>	<u>002</u>	Mar 28, 2005	
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 0.05MG BASE/ML</u>	<u>A079198</u>	<u>001</u>	Feb 10, 2011	
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A079198</u>	<u>002</u>	Feb 10, 2011	
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A079198</u>	<u>003</u>	Feb 10, 2011	
<u>AP</u>	WOCHARDT USA	<u>EQ 0.05MG BASE/ML</u>	<u>A090985</u>	<u>001</u>	May 11, 2011	
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>A090985</u>	<u>002</u>	May 11, 2011	
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>A090985</u>	<u>003</u>	May 11, 2011	
		<u>SANDOSTATIN</u>				
<u>AP</u>	+	<u>NOVARTIS</u>	<u>EQ 0.05MG BASE/ML</u>	<u>N019667</u>	<u>001</u>	Oct 21, 1988
<u>AP</u>	+		<u>EQ 0.1MG BASE/ML</u>	<u>N019667</u>	<u>002</u>	Oct 21, 1988
<u>AP</u>	+		<u>EQ 0.2MG BASE/ML</u>	<u>N019667</u>	<u>004</u>	Jun 12, 1991
<u>AP</u>	+		<u>EQ 0.5MG BASE/ML</u>	<u>N019667</u>	<u>003</u>	Oct 21, 1988
<u>AP</u>	+		<u>EQ 1MG BASE/ML</u>	<u>N019667</u>	<u>005</u>	Jun 12, 1991
		<u>SANDOSTATIN LAR</u>				
	NOVARTIS	<u>EQ 10MG BASE/VIAL</u>	<u>N021008</u>	<u>001</u>	Nov 25, 1998	
		<u>EQ 20MG BASE/VIAL</u>	<u>N021008</u>	<u>002</u>	Nov 25, 1998	
	+	<u>EQ 30MG BASE/VIAL</u>	<u>N021008</u>	<u>003</u>	Nov 25, 1998	

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OCUFLOX

<u>AT</u>	+	<u>ALLERGAN</u>	<u>0.3%</u>	<u>N019921</u>	<u>001</u>	Jul 30, 1993
<u>AT</u>		<u>AKORN</u>	<u>0.3%</u>	<u>A076407</u>	<u>001</u>	Apr 15, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 315 (of 424)

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OFLOXACIN

<u>AT</u>	ALCON PHARMS LTD	<u>0.3%</u>	<u>A076231</u>	<u>001</u>	May 14, 2004
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076622</u>	<u>001</u>	May 14, 2004
<u>AT</u>	FDC LTD	<u>0.3%</u>	<u>A078559</u>	<u>001</u>	Feb 25, 2009
<u>AT</u>	FERA PHARMS	<u>0.3%</u>	<u>A076830</u>	<u>001</u>	Aug 31, 2004
<u>AT</u>	HI TECH PHARMA	<u>0.3%</u>	<u>A076615</u>	<u>001</u>	May 14, 2004
<u>AT</u>	NOVEX	<u>0.3%</u>	<u>A076513</u>	<u>001</u>	May 14, 2004
<u>AT</u>	SANDOZ	<u>0.3%</u>	<u>A076848</u>	<u>001</u>	Nov 25, 2008

SOLUTION/DROPS; OTIC

FLOXIN OTIC

<u>AT</u> + DAIICHI		<u>0.3%</u>	<u>N020799</u>	<u>001</u>	Dec 16, 1997
	<u>OFLOXACIN</u>				
<u>AT</u>	ALCON PHARMS LTD	<u>0.3%</u>	<u>A078222</u>	<u>001</u>	Mar 17, 2008
<u>AT</u>	APOTEX INC	<u>0.3%</u>	<u>A076527</u>	<u>001</u>	Sep 28, 2007
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A076128</u>	<u>001</u>	Mar 17, 2008
<u>AT</u>	FERA PHARMS	<u>0.3%</u>	<u>A090395</u>	<u>001</u>	Aug 11, 2009
<u>AT</u>	HI TECH PHARMA	<u>0.3%</u>	<u>A076616</u>	<u>001</u>	Mar 17, 2008

TABLET; ORAL

OFLOXACIN

<u>AB</u>	DR REDDYS LABS LTD	<u>200MG</u>	<u>A077098</u>	<u>001</u>	Feb 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077098</u>	<u>002</u>	Feb 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077098</u>	<u>003</u>	Feb 10, 2006
<u>AB</u>	RANBAXY	<u>200MG</u>	<u>A076220</u>	<u>001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>A076220</u>	<u>002</u>	Sep 02, 2003
<u>AB</u>		<u>400MG</u>	<u>A076220</u>	<u>003</u>	Sep 02, 2003
<u>AB</u>	TEVA	<u>200MG</u>	<u>A076182</u>	<u>001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>A076182</u>	<u>002</u>	Sep 02, 2003
<u>AB</u> +		<u>400MG</u>	<u>A076182</u>	<u>003</u>	Sep 02, 2003

OLANZAPINE

INJECTABLE; INTRAMUSCULAR

OLANZAPINE

<u>AP</u>	INNOPHARMA LLC	<u>10MG/VIAL</u>	<u>A201588</u>	<u>001</u>	Oct 24, 2011
<u>AP</u> +	LILLY	<u>10MG/VIAL</u>	<u>N021253</u>	<u>001</u>	Mar 29, 2004

TABLET; ORAL

OLANZAPINE

<u>AB</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A076133</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>	TEVA PHARMS	<u>2.5MG</u>	<u>A076000</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>5MG</u>	<u>A076000</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A076000</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A076000</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076000</u>	<u>005</u>	Oct 24, 2011

ZYPREXA

<u>AB</u>	LILLY	<u>2.5MG</u>	<u>N020592</u>	<u>001</u>	Sep 30, 1996
<u>AB</u> +		<u>5MG</u>	<u>N020592</u>	<u>002</u>	Sep 30, 1996
<u>AB</u>		<u>7.5MG</u>	<u>N020592</u>	<u>003</u>	Sep 30, 1996
<u>AB</u>		<u>10MG</u>	<u>N020592</u>	<u>004</u>	Sep 30, 1996
<u>AB</u>		<u>15MG</u>	<u>N020592</u>	<u>005</u>	Sep 09, 1997
<u>AB</u>		<u>20MG</u>	<u>N020592</u>	<u>006</u>	Sep 09, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A091265</u>	<u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A091265</u>	<u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A091265</u>	<u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A091265</u>	<u>004</u>	Oct 24, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076534</u>	<u>001</u>	Oct 24, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 316 (of 424)

OLANZAPINE

TABLET, ORALLY DISINTEGRATING; ORAL

OLANZAPINE

<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A076534</u> <u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076534</u> <u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A076534</u> <u>004</u>	Oct 24, 2011
<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>A078109</u> <u>001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A078109</u> <u>002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A078109</u> <u>003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A078109</u> <u>004</u>	Oct 24, 2011
<u>AB</u>	TORRENT PHARMS LLC	<u>5MG</u>	<u>A091415</u> <u>001</u>	Oct 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A091415</u> <u>002</u>	Oct 25, 2011
<u>AB</u>		<u>15MG</u>	<u>A091415</u> <u>003</u>	Oct 25, 2011
<u>AB</u>		<u>20MG</u>	<u>A091415</u> <u>004</u>	Oct 25, 2011
		<u>ZYPREXA ZYDIS</u>		
<u>AB</u>	+ LILLY	<u>5MG</u>	<u>N021086</u> <u>001</u>	Apr 06, 2000
<u>AB</u>		<u>10MG</u>	<u>N021086</u> <u>002</u>	Apr 06, 2000
<u>AB</u>		<u>15MG</u>	<u>N021086</u> <u>003</u>	Apr 06, 2000
<u>AB</u>		<u>20MG</u>	<u>N021086</u> <u>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

OIMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR

DAIICHI SANKYO	5MG	N021286 001	Apr 25, 2002
	20MG	N021286 003	Apr 25, 2002
+	40MG	N021286 004	Apr 25, 2002

OILOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

+ ALCON PHARMS LTD	EQ 0.2% BASE	N021545 001	Dec 22, 2004
PATANOL			
+ ALCON	EQ 0.1% BASE	N020688 001	Dec 18, 1996
SPRAY, METERED; NASAL PATANASE			
+ ALCON PHARMS LTD	0.665MG/SPRAY	N021861 001	Apr 15, 2008

OISALAZINE SODIUM

CAPSULE; ORAL

DIPENTUM

+ MEDA PHARMS	250MG	N019715 001	Jul 31, 1990
---------------	-------	-------------	--------------

OMEGA-3-ACID ETHYL ESTERS

CAPSULE; ORAL

LOVAZA

+ SMITHKLINE BEECHAM	1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS	N021654 001	Nov 10, 2004
----------------------	---	-------------	--------------

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE

<u>AB</u>	APOTEX	<u>10MG</u>	<u>A076048</u> <u>001</u>	Oct 22, 2007
-----------	--------	-------------	---------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 317 (of 424)

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE

<u>AB</u>	APOTEX	<u>20MG</u>	<u>A076048</u>	<u>002</u>	Oct 22, 2007
<u>AB</u>		<u>40MG</u>	<u>A076048</u>	<u>003</u>	Jan 21, 2009
<u>AB</u>	DR REDDYS LABS	<u>40MG</u>	<u>A078490</u>	<u>001</u>	Apr 17, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A075576</u>	<u>003</u>	Oct 22, 2007
<u>AB</u>		<u>10MG</u>	<u>A078693</u>	<u>001</u>	Mar 16, 2009
<u>AB</u>		<u>20MG</u>	<u>A075576</u>	<u>002</u>	Oct 22, 2007
<u>AB</u>		<u>20MG</u>	<u>A078693</u>	<u>002</u>	Mar 16, 2009
<u>AB</u>		<u>40MG</u>	<u>A075576</u>	<u>001</u>	Jan 21, 2009
<u>AB</u>	IMPAK LABS	<u>10MG</u>	<u>A075785</u>	<u>001</u>	Oct 22, 2007
<u>AB</u>		<u>20MG</u>	<u>A075785</u>	<u>002</u>	Oct 22, 2007
<u>AB</u>		<u>40MG</u>	<u>A075785</u>	<u>003</u>	Jan 21, 2009
<u>AB</u>	KREMERS URBAN PHARMS	<u>10MG</u>	<u>A075410</u>	<u>001</u>	Nov 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075410</u>	<u>002</u>	Nov 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075410</u>	<u>003</u>	Jan 23, 2009
<u>AB</u>	LEK PHARMS	<u>10MG</u>	<u>A075757</u>	<u>001</u>	Jan 28, 2003
<u>AB</u>		<u>20MG</u>	<u>A075757</u>	<u>002</u>	Jan 28, 2003
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A075876</u>	<u>001</u>	May 29, 2003
<u>AB</u>		<u>20MG</u>	<u>A075876</u>	<u>002</u>	May 29, 2003
<u>AB</u>		<u>40MG</u>	<u>A075876</u>	<u>003</u>	Jan 21, 2009
<u>AB</u>	SANDOZ	<u>40MG</u>	<u>A076515</u>	<u>001</u>	Jan 21, 2009
<u>AB</u>	WATSON LABS FLORIDA	<u>10MG</u>	<u>A075347</u>	<u>001</u>	May 30, 2008
<u>AB</u>		<u>20MG</u>	<u>A075347</u>	<u>002</u>	May 30, 2008
<u>AB</u>		<u>40MG</u>	<u>A075347</u>	<u>003</u>	May 30, 2008
	<u>PRILOSEC</u>				
<u>AB</u>	ASTRAZENECA	<u>10MG</u>	<u>N019810</u>	<u>003</u>	Oct 05, 1995
<u>AB</u>	+	<u>20MG</u>	<u>N019810</u>	<u>001</u>	Sep 14, 1989
<u>AB</u>	+	<u>40MG</u>	<u>N019810</u>	<u>002</u>	Jan 15, 1998

OMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL

PRILOSEC

ASTRAZENECA	EQ 2.5MG BASE/PACKET	N022056	001	Mar 20, 2008
+	EQ 10MG BASE/PACKET	N022056	002	Mar 20, 2008

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE; ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	PAR PHARM	<u>20MG;1.1GM</u>	<u>A078966</u>	<u>001</u>	May 25, 2010
<u>AB</u>		<u>40MG;1.1GM</u>	<u>A078966</u>	<u>002</u>	May 25, 2010
	<u>ZEGERID</u>				
<u>AB</u>	SANTARUS	<u>20MG;1.1GM</u>	<u>N021849</u>	<u>001</u>	Feb 27, 2006

FOR SUSPENSION; ORAL

ZEGERID

SANTARUS	20MG/PACKET;1.68GM/PACKET	N021636	001	Jun 15, 2004
+	40MG/PACKET;1.68GM/PACKET	N021636	002	Dec 21, 2004

ONDANSETRON

FILM; ORAL

ZUPLENZ

MONOSOL RX LLC	4MG	N022524	001	Jul 02, 2010
+	8MG	N022524	002	Jul 02, 2010

TABLET, ORALLY DISINTEGRATING; ORAL

ONDANSETRON

<u>AB</u>	AUROBINDO PHARMA	<u>4MG</u>	<u>A090469</u>	<u>001</u>	Apr 12, 2010
<u>AB</u>		<u>8MG</u>	<u>A090469</u>	<u>002</u>	Apr 12, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 318 (of 424)

ONDANSETRON

TABLET, ORALLY DISINTEGRATING; ORAL

ONDANSETRON

<u>AB</u>	BARR	<u>4MG</u>	<u>A076693</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A076693</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	GLENMARK GENERICS	<u>4MG</u>	<u>A078152</u>	<u>001</u>	Jun 27, 2007
<u>AB</u>		<u>8MG</u>	<u>A078152</u>	<u>002</u>	Jun 27, 2007
<u>AB</u>	MYLAN	<u>4MG</u>	<u>A078139</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A078139</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	NESHER PHARMS	<u>4MG</u>	<u>A077717</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A077717</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	PAR PHARM	<u>4MG</u>	<u>A076506</u>	<u>001</u>	Dec 26, 2006
<u>AB</u>		<u>8MG</u>	<u>A076506</u>	<u>002</u>	Dec 26, 2006
<u>AB</u>	RANBAXY	<u>4MG</u>	<u>A078602</u>	<u>001</u>	Feb 24, 2011
<u>AB</u>		<u>8MG</u>	<u>A078602</u>	<u>002</u>	Feb 24, 2011
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>A078050</u>	<u>001</u>	Aug 13, 2007
<u>AB</u>		<u>8MG</u>	<u>A078050</u>	<u>002</u>	Aug 13, 2007
<u>AB</u>	SUN PHARM INDUS	<u>4MG</u>	<u>A077557</u>	<u>001</u>	Aug 02, 2007
<u>AB</u>		<u>8MG</u>	<u>A077557</u>	<u>002</u>	Aug 02, 2007
<u>AB</u>	TEVA	<u>4MG</u>	<u>A076810</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>8MG</u>	<u>A076810</u>	<u>002</u>	Jun 25, 2007
<u>ZOFRAN ODT</u>					
<u>AB</u>	GLAXOSMITHKLINE	<u>4MG</u>	<u>N020781</u>	<u>001</u>	Jan 27, 1999
<u>AB</u>	+	<u>8MG</u>	<u>N020781</u>	<u>002</u>	Jan 27, 1999
<u>AB</u>	ONDANSETRON				
	PAR PHARM	<u>16MG</u>	<u>A077406</u>	<u>001</u>	Dec 26, 2006
		<u>24MG</u>	<u>A077406</u>	<u>002</u>	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

<u>AP</u>	APOTEX	<u>EQ 2MG BASE/ML</u>	<u>A077368</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	APP PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A076974</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A077365</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	BEDFORD	<u>EQ 2MG BASE/ML</u>	<u>A076967</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	EMCURE PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A090424</u>	<u>001</u>	Apr 16, 2010
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A079224</u>	<u>001</u>	Sep 25, 2009
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>	<u>A076781</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077473</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077840</u>	<u>001</u>	Jan 19, 2007
<u>AP</u>	LANNETT	<u>EQ 2MG BASE/ML</u>	<u>A090116</u>	<u>001</u>	Apr 14, 2010
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A090883</u>	<u>001</u>	Aug 05, 2010
<u>AP</u>	LUITPOLD	<u>EQ 2MG BASE/ML</u>	<u>A077582</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A079039</u>	<u>001</u>	Nov 18, 2008
<u>AP</u>	PFIZER	<u>EQ 2MG BASE/ML</u>	<u>A078257</u>	<u>001</u>	Apr 23, 2008
<u>AP</u>	PLIVA HRVATSKA DOO	<u>EQ 2MG BASE/ML</u>	<u>A077544</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	SANDOZ	<u>EQ 2MG BASE/ML</u>	<u>A077430</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>	SPECTRUM PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A078180</u>	<u>001</u>	Mar 26, 2007
<u>AP</u>	SUN PHARM INDUS (IN)	<u>EQ 2MG BASE/ML</u>	<u>A077172</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	TEVA	<u>EQ 2MG BASE/ML</u>	<u>A076876</u>	<u>001</u>	Nov 22, 2006
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077577</u>	<u>001</u>	Dec 26, 2006

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

<u>AP</u>	BEDFORD LABS	<u>EQ 0.64MG BASE/ML</u>	<u>A078291</u>	<u>001</u>	Apr 13, 2009
<u>AP</u>	CLARIS LIFESCIENCES	<u>EQ 0.64MG BASE/ML</u>	<u>A078308</u>	<u>001</u>	Mar 17, 2008
<u>AP</u>	+	<u>EQ 0.64MG BASE/ML</u>	<u>A077348</u>	<u>001</u>	Feb 01, 2007
<u>AP</u>	HOSPIRA	<u>EQ 0.64MG BASE/ML</u>	<u>A077480</u>	<u>001</u>	Nov 22, 2006

ONDANSETRON HYDROCHLORIDE AND SODIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	+ BAXTER HLTHCARE	<u>EQ 0.64MG BASE/ML</u>	<u>N021915</u>	<u>002</u>	Dec 27, 2006
-----------	-------------------	--------------------------	----------------	------------	--------------

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	APOTEX INC	<u>EQ 2MG BASE/ML</u>	<u>A077343</u>	<u>001</u>	Dec 26, 2006
-----------	------------	-----------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 319 (of 424)

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	APP PHARMS	<u>EQ 2MG BASE/ML</u>	<u>A076972</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A077541</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	BEDFORD LABS	<u>EQ 2MG BASE/ML</u>	<u>A077011</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>	<u>A076780</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077548</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	LUITPOLD	<u>EQ 2MG BASE/ML</u>	<u>A077387</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A079032</u>	<u>001</u>	Nov 18, 2008
<u>AP</u>	PFIZER	<u>EQ 2MG BASE/ML</u>	<u>A078244</u>	<u>001</u>	Apr 23, 2008
<u>AP</u>	SANDOZ	<u>EQ 2MG BASE/ML</u>	<u>A077551</u>	<u>001</u>	Jun 27, 2007
<u>AP</u>	SUN PHARM INDs LTD	<u>EQ 2MG BASE/ML</u>	<u>A077173</u>	<u>001</u>	Dec 26, 2006
<u>AP</u>	TARO PHARMS IRELAND	<u>EQ 2MG BASE/ML</u>	<u>A078014</u>	<u>001</u>	Mar 21, 2008
<u>AP</u>	TEVA	<u>EQ 2MG BASE/ML</u>	<u>A076759</u>	<u>001</u>	Nov 22, 2006
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077716</u>	<u>001</u>	Dec 26, 2006
	<u>ZOFRAN</u>				
<u>AP</u>	+ GLAXOSMITHKLINE	<u>EQ 2MG BASE/ML</u>	<u>N020007</u>	<u>001</u>	Jan 04, 1991
	<u>ZOFRAN PRESERVATIVE FREE</u>				
<u>AP</u>	+ GLAXOSMITHKLINE	<u>EQ 2MG BASE/ML</u>	<u>N020007</u>	<u>003</u>	Dec 10, 1993

SOLUTION; ORAL

ONDANSETRON HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>EQ 4MG BASE/5ML</u>	<u>A091483</u>	<u>001</u>	Jan 31, 2011
<u>AA</u>	APOTEX	<u>EQ 4MG BASE/5ML</u>	<u>A078127</u>	<u>001</u>	Jun 25, 2007
<u>AA</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE/5ML</u>	<u>A078776</u>	<u>001</u>	Nov 28, 2007
<u>AA</u>	ROXANE	<u>EQ 4MG BASE/5ML</u>	<u>A076960</u>	<u>001</u>	Dec 26, 2006
<u>AA</u>	SILARX	<u>EQ 4MG BASE/5ML</u>	<u>A091342</u>	<u>001</u>	Jan 27, 2011
<u>AA</u>	TARO	<u>EQ 4MG BASE/5ML</u>	<u>A077009</u>	<u>001</u>	Nov 30, 2007
	<u>ZOFRAN</u>				
<u>AA</u>	+ GLAXOSMITHKLINE	<u>EQ 4MG BASE/5ML</u>	<u>N020605</u>	<u>001</u>	Jan 24, 1997

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 4MG BASE</u>	<u>A077306</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077306</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE</u>	<u>A078539</u>	<u>001</u>	Jul 31, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078539</u>	<u>002</u>	Jul 31, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A078539</u>	<u>003</u>	Jul 31, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE</u>	<u>A076183</u>	<u>003</u>	Dec 26, 2006
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076183</u>	<u>002</u>	Dec 26, 2006
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076183</u>	<u>001</u>	Dec 26, 2006
<u>AB</u>	GLENMARK GENERICS	<u>EQ 4MG BASE</u>	<u>A077535</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077535</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077535</u>	<u>003</u>	Jun 25, 2007
<u>AB</u>	MYLAN	<u>EQ 4MG BASE</u>	<u>A076930</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076930</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076930</u>	<u>004</u>	Jun 25, 2007
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 4MG BASE</u>	<u>A077851</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077851</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	PAR PHARM	<u>EQ 4MG BASE</u>	<u>A077303</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077303</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077303</u>	<u>004</u>	Jun 25, 2007
<u>AB</u>	PLIVA HRVATSKA DOO	<u>EQ 4MG BASE</u>	<u>A077112</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077112</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077112</u>	<u>003</u>	Jun 25, 2007
<u>AB</u>	SANDOZ	<u>EQ 4MG BASE</u>	<u>A077517</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077517</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077517</u>	<u>003</u>	Jun 25, 2007
<u>AB</u>	SUN PHARM INDs (IN)	<u>EQ 4MG BASE</u>	<u>A077050</u>	<u>001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077050</u>	<u>002</u>	Jun 25, 2007
<u>AB</u>	TARO	<u>EQ 4MG BASE</u>	<u>A077729</u>	<u>001</u>	Mar 28, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 320 (of 424)

ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>	TARO	<u>EQ 8MG BASE</u>	<u>A077729 002</u>	Mar 28, 2011
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077729 003</u>	Mar 28, 2011
<u>AB</u>	TEVA	<u>EQ 4MG BASE</u>	<u>A076252 001</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A076252 002</u>	Jun 25, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076252 003</u>	Jun 25, 2007
<u>AB</u>	WEST WARD	<u>EQ 4MG BASE</u>	<u>A077545 001</u>	Sep 06, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077545 002</u>	Sep 06, 2007
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A077545 003</u>	Sep 06, 2007
<u>ZOFRAN</u>				
<u>AB</u>	GLAXOSMITHKLINE	<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> +		<u>EQ 24MG BASE</u>	<u>N020103 003</u>	Aug 27, 1999
ONDANSETRON HYDROCHLORIDE				
	DR REDDYS LABS LTD	EQ 16MG BASE	A076559 001	Dec 26, 2006

ORLISTAT

CAPSULE; ORAL

XENICAL

+ HOFFMANN LA ROCHE 120MG

N020766 001 Apr 23, 1999

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NOREFLEX

<u>AP</u> +	MEDICIS	<u>30MG/ML</u>	<u>N013055 001</u>	
<u>ORPHENADRINE CITRATE</u>				
<u>AP</u>	AKORN	<u>30MG/ML</u>	<u>A040484 001</u>	May 24, 2006
<u>AP</u>	BEDFORD LABS	<u>30MG/ML</u>	<u>A040463 001</u>	Mar 04, 2003
<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>		<u>30MG/ML</u>	<u>A087062 001</u>	

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>	KIEL	<u>100MG</u>	<u>A040249 001</u>	Jan 29, 1999
<u>AB</u> +	SANDOZ	<u>100MG</u>	<u>A040327 001</u>	Feb 15, 2000

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

TAMIFLU

ROCHE

EQ 30MG BASE

N021087 003 Jul 02, 2007

EQ 45MG BASE

N021087 002 Jul 02, 2007

+

EQ 75MG BASE

N021087 001 Oct 27, 1999

FOR SUSPENSION; ORAL

TAMIFLU

ROCHE

EQ 6MG BASE/ML

N021246 002 Mar 21, 2011

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

<u>AP</u> +	SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A061490 003</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062737 001</u>	Dec 23, 1986
<u>AP</u> +		<u>EQ 2GM BASE/VIAL</u>	<u>A061490 004</u>	
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062737 002</u>	Dec 23, 1986
<u>AP</u> +		<u>EQ 10GM BASE/VIAL</u>	<u>A061490 006</u>	May 09, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 321 (of 424)

OXACILLIN SODIUM

INJECTABLE; INJECTION

BACTOCILL IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	EQ 20MG BASE/ML	N050640 001	Oct 26, 1989
+	EQ 40MG BASE/ML	N050640 002	Oct 26, 1989

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

AP + SANOFI AVENTIS US	<u>50MG/10ML (5MG/ML)</u>	<u>N021759 001</u>	Jan 31, 2005
AP +	<u>100MG/20ML (5MG/ML)</u>	<u>N021759 002</u>	Jan 31, 2005

OXALIPLATIN

AP APP PHARMS	<u>50MG/VIAL</u>	<u>A078819 001</u>	Jun 02, 2010
AP EBEWE PHARMA	<u>100MG/VIAL</u>	<u>A078819 002</u>	Jun 02, 2010
AP FRESENIUS KABI ONCOL	<u>50MG/10ML (5MG/ML)</u>	<u>A078812 001</u>	Aug 07, 2009
AP	<u>100MG/20ML (5MG/ML)</u>	<u>A078812 002</u>	Aug 07, 2009
AP HOSPIRA INC	<u>50MG/10ML (5MG/ML)</u>	<u>A078811 001</u>	Jun 10, 2010
AP	<u>100MG/20ML (5MG/ML)</u>	<u>A078811 002</u>	Jun 10, 2010
AP HOSPIRA WORLDWIDE	<u>50MG/VIAL</u>	<u>A078810 001</u>	Aug 07, 2009
AP	<u>100MG/VIAL</u>	<u>A078810 002</u>	Aug 07, 2009
AP SANDOZ	<u>50MG/10ML (5MG/ML)</u>	<u>A078815 001</u>	Sep 30, 2009
AP	<u>100MG/20ML (5MG/ML)</u>	<u>A078815 002</u>	Sep 30, 2009
AP + SUN PHARMA GLOBAL	<u>50MG/VIAL</u>	<u>A078813 001</u>	Aug 07, 2009
AP +	<u>100MG/VIAL</u>	<u>A078813 002</u>	Aug 07, 2009
AP TEVA PARENTERAL	<u>50MG/10ML (5MG/ML)</u>	<u>N022160 001</u>	Aug 07, 2009
AP	<u>100MG/20ML (5MG/ML)</u>	<u>N022160 002</u>	Aug 07, 2009
AP ELOXATIN			
+ SANOFI AVENTIS US	<u>200MG/40ML (5MG/ML)</u>	<u>N021759 003</u>	Nov 17, 2006

OXANDROLONE

TABLET; ORAL

OXANDRIN

AB SAVIENT PHARMS	<u>2.5MG</u>	<u>N013718 001</u>	
AB +	<u>10MG</u>	<u>N013718 002</u>	Nov 05, 2001

OXANDROLONE

AB PAR PHARM	<u>2.5MG</u>	<u>A077827 001</u>	Jun 22, 2007
AB	<u>10MG</u>	<u>A077827 002</u>	Jun 22, 2007
AB SANDOZ	<u>2.5MG</u>	<u>A076897 001</u>	Dec 01, 2006
AB	<u>10MG</u>	<u>A076897 002</u>	Dec 01, 2006
AB UPSHER SMITH	<u>2.5MG</u>	<u>A076761 001</u>	Dec 01, 2006
AB	<u>10MG</u>	<u>A078033 001</u>	Mar 22, 2007

OXAPROZIN

TABLET; ORAL

DAYPRO

AB + GD SEARLE	<u>600MG</u>	<u>N018841 004</u>	Oct 29, 1992
AB OXAPROZIN			

AB APOTEX INC	<u>600MG</u>	<u>A075987 001</u>	Sep 02, 2004
AB CARACO	<u>600MG</u>	<u>A075844 001</u>	Jan 03, 2002
AB DR REDDYS LABS LTD	<u>600MG</u>	<u>A075855 001</u>	Jan 31, 2001
AB IVAX SUB TEVA PHARMS	<u>600MG</u>	<u>A075846 001</u>	May 13, 2002
AB MYLAN	<u>600MG</u>	<u>A075847 001</u>	Feb 28, 2001
AB SANDOZ	<u>600MG</u>	<u>A075845 001</u>	Jan 31, 2001
AB TEVA	<u>600MG</u>	<u>A075849 001</u>	Jul 03, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 322 (of 424)

OXAPROZIN

TABLET; ORAL

OXAPROZIN

<u>AB</u>	WATSON LABS	<u>600MG</u>	<u>A075848 001</u>	Feb 09, 2001
-----------	-------------	--------------	--------------------	--------------

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072251 001</u>	Apr 14, 1988
<u>AB</u>		<u>15MG</u>	<u>A072252 001</u>	Apr 14, 1988
<u>AB</u>		<u>30MG</u>	<u>A072253 001</u>	Apr 14, 1988
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>10MG</u>	<u>A070943 001</u>	Aug 03, 1987
<u>AB</u>		<u>15MG</u>	<u>A070944 001</u>	Aug 03, 1987
<u>AB</u>	+	<u>30MG</u>	<u>A070945 001</u>	Aug 03, 1987
<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
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A072952 001</u>	Sep 28, 1990
<u>AB</u>		<u>15MG</u>	<u>A072953 001</u>	Sep 28, 1990
<u>AB</u>		<u>30MG</u>	<u>A072954 001</u>	Sep 28, 1990

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

<u>AB</u>	RANBAXY	<u>300MG/5ML</u>	<u>A078734 001</u>	Jun 26, 2009
<u>AB</u>	<u>TRILEPTAL</u>			
<u>AB</u>	+	NOVARTIS	<u>N021285 001</u>	May 25, 2001

TABLET; ORAL

OXCARBAZEPINE

<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
<u>AB</u>	CADISTA PHARMS	<u>150MG</u>	<u>A090239 001</u>	Jan 25, 2010
<u>AB</u>		<u>300MG</u>	<u>A090239 002</u>	Jan 25, 2010
<u>AB</u>		<u>600MG</u>	<u>A090239 003</u>	Jan 25, 2010
<u>AB</u>	GLENMARK GENERICS	<u>150MG</u>	<u>A077802 001</u>	Oct 09, 2007
<u>AB</u>		<u>300MG</u>	<u>A077802 002</u>	Oct 09, 2007
<u>AB</u>		<u>600MG</u>	<u>A077802 003</u>	Oct 09, 2007
<u>AB</u>	ROXANE	<u>150MG</u>	<u>A077795 001</u>	Oct 09, 2007
<u>AB</u>		<u>300MG</u>	<u>A077795 002</u>	Oct 09, 2007
<u>AB</u>		<u>600MG</u>	<u>A077795 003</u>	Oct 09, 2007
<u>AB</u>	SUN PHARM IND'S	<u>150MG</u>	<u>A077794 001</u>	Oct 09, 2007
<u>AB</u>		<u>300MG</u>	<u>A077794 002</u>	Oct 09, 2007
<u>AB</u>		<u>600MG</u>	<u>A077794 003</u>	Oct 09, 2007
<u>AB</u>	TARO	<u>150MG</u>	<u>A077801 001</u>	Nov 15, 2007
<u>AB</u>		<u>300MG</u>	<u>A077801 002</u>	Nov 15, 2007
<u>AB</u>		<u>600MG</u>	<u>A077801 003</u>	Nov 15, 2007
<u>AB</u>	TEVA PHARMS	<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>	<u>TRILEPTAL</u>			
<u>AB</u>	NOVARTIS	<u>150MG</u>	<u>N021014 001</u>	Jan 14, 2000
<u>AB</u>		<u>300MG</u>	<u>N021014 002</u>	Jan 14, 2000
<u>AB</u>	+	<u>600MG</u>	<u>N021014 003</u>	Jan 14, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 323 (of 424)

OXICONAZOLE NITRATE

CREAM; TOPICAL OXISTAT + ALTANA	EQ 1% BASE	N019828 001	Dec 30, 1988
LOTION; TOPICAL OXISTAT + FOUGERA PHARMS	EQ 1% BASE	N020209 001	Sep 30, 1992

OXTRIPHYLLINE

TABLET, EXTENDED RELEASE; ORAL CHOLEDYL SA + WARNER CHILCOTT	400MG	A087863 001	May 24, 1983
	600MG	A086742 001	

OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL OXYTROL + WATSON LABS (UTAH)	3.9MG/24HR	N021351 002	Feb 26, 2003
GEL, METERED; TRANSDERMAL ANTUROL + ANTARES PHARMA INC	3%	N202513 001	Dec 07, 2011

OXYBUTYNIN CHLORIDE

GEL; TRANSDERMAL GELNIQUE + WATSON LABS	10%(100MG/PACKET)	N022204 001	Jan 27, 2009
---	-------------------	-------------	--------------

SYRUP; ORAL

OXYBUTYNIN CHLORIDE

<u>AA</u> MIKART	<u>5MG/5ML</u>	<u>A075039 001</u>	Jan 29, 1999
<u>AA</u> NOVEX	<u>5MG/5ML</u>	<u>A074997 001</u>	Oct 15, 1997
<u>AA</u> PHARM ASSOC	<u>5MG/5ML</u>	<u>A075137 001</u>	Dec 18, 1998
<u>AA</u> SILARX	<u>5MG/5ML</u>	<u>A074520 001</u>	Mar 29, 1996
<u>AA</u> VINTAGE PHARMS	<u>5MG/5ML</u>	<u>A076682 001</u>	Dec 28, 2004
<u>AA</u> + WOCKHARDT	<u>5MG/5ML</u>	<u>A074868 001</u>	Feb 12, 1997

TABLET; ORAL

DITROPAN

<u>AB</u> + JANSSEN PHARMS	<u>5MG</u>	<u>N017577 001</u>
----------------------------	------------	--------------------

OXYBUTYNIN CHLORIDE

<u>AB</u> PLIVA	<u>5MG</u>	<u>A071655 001</u>	Nov 14, 1988
<u>AB</u> USL PHARMA	<u>5MG</u>	<u>A074625 001</u>	Jul 31, 1996
<u>AB</u> VINTAGE PHARMS	<u>5MG</u>	<u>A075079 001</u>	Oct 31, 1997

TABLET, EXTENDED RELEASE; ORAL

DITROPAN XL

<u>AB</u> JANSSEN PHARMS	<u>5MG</u>	<u>N020897 001</u>	Dec 16, 1998
<u>AB</u>	<u>10MG</u>	<u>N020897 002</u>	Dec 16, 1998
<u>AB</u> +	<u>15MG</u>	<u>N020897 003</u>	Jun 22, 1999

OXYBUTYNIN CHLORIDE

<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>	<u>10MG</u>	<u>A076644 001</u>	Nov 09, 2006
<u>AB</u>	<u>15MG</u>	<u>A078293 001</u>	May 10, 2007
<u>AB</u> OSMOTICA PHARM	<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

PRESCRIPTION DRUG PRODUCT LIST

3 - 324 (of 424)

OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL			
OXYCODONE HYDROCHLORIDE			
+ LEHIGH VALLEY	5MG	N200534 001	Oct 20, 2010
SOLUTION; ORAL			
OXYCODONE HYDROCHLORIDE			
+ LEHIGH VALLEY	100MG/5ML	N200535 001	Oct 20, 2010
TABLET; ORAL			
<u>OXYCODONE HYDROCHLORIDE</u>			
AB ACTAVIS TOTOWA	<u>15MG</u>	A076636 001	Feb 06, 2004
AB	<u>30MG</u>	A076636 002	Feb 06, 2004
AB ALVOGEN INC	<u>5MG</u>	A202116 001	Dec 30, 2011
AB	<u>15MG</u>	A202116 002	Dec 30, 2011
AB	<u>30MG</u>	A202116 003	Dec 30, 2011
AB AVANTHI INC	<u>5MG</u>	A091393 001	Aug 31, 2009
AB	<u>10MG</u>	A091393 002	Aug 31, 2009
AB	<u>15MG</u>	A091393 003	Aug 31, 2009
AB	<u>20MG</u>	A091393 004	Aug 31, 2009
AB	<u>30MG</u>	A091393 005	Aug 31, 2009
AB COASTAL PHARMS	<u>5MG</u>	A091313 001	Feb 18, 2011
AB	<u>15MG</u>	A091313 002	Feb 18, 2011
AB	<u>30MG</u>	A091313 003	Feb 18, 2011
AB COREPHARMA	<u>5MG</u>	A090895 001	Aug 24, 2009
AB	<u>15MG</u>	A090895 002	Aug 24, 2009
AB	<u>30MG</u>	A090895 003	Aug 24, 2009
AB MALLINCKRODT	<u>15MG</u>	A076758 001	Jun 30, 2004
AB	<u>30MG</u>	A076758 002	Jun 30, 2004
AB MALLINCKRODT INC	<u>5MG</u>	A078206 001	Mar 19, 2007
AB NESHER PHARMS	<u>5MG</u>	A077290 001	Dec 08, 2005
AB	<u>10MG</u>	A077290 002	Dec 08, 2005
AB	<u>15MG</u>	A077290 003	Dec 08, 2005
AB	<u>20MG</u>	A077290 004	Dec 08, 2005
AB	<u>30MG</u>	A077290 005	Dec 08, 2005
AB RHODES PHARMS	<u>5MG</u>	A091490 001	Mar 09, 2011
AB	<u>10MG</u>	A091490 002	Mar 09, 2011
AB	<u>15MG</u>	A091490 003	Mar 09, 2011
AB	<u>20MG</u>	A091490 004	Mar 09, 2011
AB	<u>30MG</u>	A091490 005	Mar 09, 2011
AB SUN PHARM IND S INC	<u>5MG</u>	A090659 001	Apr 10, 2009
AB	<u>15MG</u>	A090659 002	Apr 10, 2009
AB	<u>30MG</u>	A090659 003	Apr 10, 2009
AB VINTAGE PHARMS	<u>5MG</u>	A077712 003	Mar 02, 2009
AB	<u>15MG</u>	A077712 001	Jan 31, 2007
AB	<u>30MG</u>	A077712 002	Jan 31, 2007
<u>ROXICODONE</u>			
AB XANODYNE PHARMS	<u>5MG</u>	N021011 003	May 15, 2009
AB +	<u>15MG</u>	N021011 001	Aug 31, 2000
AB	<u>30MG</u>	N021011 002	Aug 31, 2000
OXECTA			
KING PHARMS R AND D	5MG	N202080 001	Jun 17, 2011
	7.5MG	N202080 002	Jun 17, 2011
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

PRESCRIPTION DRUG PRODUCT LIST

3 - 325 (of 424)

OXYMETHOLONE

TABLET; ORAL
 ANADROL-50
 + MEDA PHARMS 50MG N016848 001

OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION
 OPANA
 + ENDO PHARMS 1MG/ML N011707 002

TABLET; ORAL
OPANA
AB ENDO PHARMS 5MG N021611 001 Jun 22, 2006
AB + 10MG N021611 002 Jun 22, 2006
OXYMORPHONE HYDROCHLORIDE
AB ROXANE 5MG A090964 001 Sep 27, 2010
AB 10MG A090964 002 Sep 27, 2010
AB TEVA 5MG A091443 002 Feb 15, 2011
AB 10MG A091443 001 Feb 15, 2011

TABLET, EXTENDED RELEASE; ORAL

OPANA ER
AB ENDO PHARMS 5MG N021610 001 Jun 22, 2006
AB 10MG N021610 002 Jun 22, 2006
AB 20MG N021610 003 Jun 22, 2006
AB 30MG N021610 007 Feb 29, 2008
AB + 40MG N021610 004 Jun 22, 2006

OXYMORPHONE HYDROCHLORIDE
AB ACTAVIS 7.5MG A079046 001 Dec 13, 2010
AB 15MG A079046 002 Dec 13, 2010
AB IMPAX LABS 5MG A079087 001 Jun 14, 2010
AB 7.5MG A079087 002 Dec 21, 2010
AB 10MG A079087 003 Jun 14, 2010
AB 15MG A079087 004 Dec 21, 2010
AB 20MG A079087 005 Jun 14, 2010
AB 30MG A079087 006 Jul 22, 2010
AB 40MG A079087 007 Jun 14, 2010

OPANA ER
 ENDO PHARMS 5MG N201655 001 Dec 09, 2011
 7.5MG N201655 002 Dec 09, 2011
 10MG N201655 003 Dec 09, 2011
 15MG N201655 004 Dec 09, 2011
 20MG N201655 005 Dec 09, 2011
 30MG N201655 006 Dec 09, 2011
 40MG N201655 007 Dec 09, 2011

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC
 TERRAMYCIN W/ POLYMYXIN B SULFATE
 + PFIZER EQ 5MG BASE/GM;10,000 UNITS/GM A061015 001

OXYTOCIN

INJECTABLE; INJECTION
OXYTOCIN
AP + APP PHARMS 10USP UNITS/ML (10USP UNITS/ML) N018248 001
AP + 100USP UNITS/10ML (10USP UNITS/ML) N018248 002
AP + BAXTER HLTHCARE CORP 10USP UNITS/ML (10USP UNITS/ML) N018243 001
AP + 100USP UNITS/10ML (10USP UNITS/ML) N018243 002 Jan 10, 2007
AP TEVA PARENTERAL 10USP UNITS/ML (10USP UNITS/ML) A077453 001 Jan 24, 2008
AP 100USP UNITS/10ML (10USP UNITS/ML) A077453 002 Jan 24, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 326 (of 424)

OXYTOCIN

INJECTABLE; INJECTION

PITOCIN

<u>AP</u>	+ JHP PHARMS	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018261 001</u>	
<u>AP</u>		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018261 002</u>	Jul 27, 2007
	OXYTOCIN			
	+ APP PHARMS	300USP UNITS/30ML (10USP UNITS/ML)	N018248 003	Jul 27, 2007

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

+ ABRAXIS BIOSCIENCE 100MG/VIAL

N021660 001 Jan 07, 2005

INJECTABLE; INJECTION

PACLITAXEL

<u>AP</u>	ACCORD HLTHCARE INC	<u>6MG/ML</u>	<u>A075436 001</u>	Nov 12, 2004
<u>AP</u>	ACTAVIS TOTOWA	<u>6MG/ML</u>	<u>A090130 001</u>	Dec 09, 2009
<u>AP</u>	BEDFORD	<u>6MG/ML</u>	<u>A075190 001</u>	Jan 28, 2002
<u>AP</u>	EBEWE PHARMA	<u>6MG/ML</u>	<u>A078167 001</u>	Dec 26, 2007
<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	<u>6MG/ML</u>	<u>A075278 001</u>	Jan 25, 2002
<u>AP</u>	ONCO THERAPIES LTD	<u>6MG/ML</u>	<u>A091540 001</u>	Sep 29, 2011
<u>AP</u>	TEVA PARENTERAL	<u>6MG/ML</u>	<u>A075184 001</u>	Jan 25, 2002

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA

JANSSEN PHARMS	1.5MG	N021999 006	Aug 26, 2008
	3MG	N021999 001	Dec 19, 2006
+	6MG	N021999 002	Dec 19, 2006
	9MG	N021999 003	Dec 19, 2006

PALIPERIDONE 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

PALONOSETRON HYDROCHLORIDE

INJECTABLE; INTRAVENOUS

ALOXI

+ HELSINN HLTHCARE	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	N021372 002	Feb 29, 2008
+	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N021372 001	Jul 25, 2003

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

AREDIA

<u>AP</u>	+ NOVARTIS	<u>30MG/VIAL</u>	<u>N020036 001</u>	Oct 31, 1991
	<u>PAMIDRONATE DISODIUM</u>			
<u>AP</u>	APP PHARMS	<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>	AREVA PHARMS	<u>30MG/VIAL</u>	<u>A077433 001</u>	Nov 26, 2008
<u>AP</u>		<u>60MG/VIAL</u>	<u>A077433 002</u>	Nov 26, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 327 (of 424)

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

<u>AP</u>	AREVA PHARMS	<u>90MG/VIAL</u>	<u>A077433</u>	<u>003</u>	Nov 26, 2008
<u>AP</u>	BEDFORD	<u>30MG/VIAL</u>	<u>A075290</u>	<u>001</u>	Apr 30, 2001
<u>AP</u> +		<u>30MG/10ML (3MG/ML)</u>	<u>N021113</u>	<u>001</u>	Mar 04, 2002
<u>AP</u>		<u>90MG/VIAL</u>	<u>A075290</u>	<u>003</u>	Apr 30, 2001
<u>AP</u> +		<u>90MG/10ML (9MG/ML)</u>	<u>N021113</u>	<u>002</u>	Mar 04, 2002
<u>AP</u> +	HOSPIRA	<u>30MG/10ML (3MG/ML)</u>	<u>A075841</u>	<u>001</u>	Jun 27, 2002
<u>AP</u> +		<u>60MG/10ML (6MG/ML)</u>	<u>A075841</u>	<u>002</u>	Jun 27, 2002
<u>AP</u> +		<u>90MG/10ML (9MG/ML)</u>	<u>A075841</u>	<u>003</u>	Jun 27, 2002
<u>AP</u>	LUITPOLD	<u>30MG/10ML (3MG/ML)</u>	<u>A078942</u>	<u>001</u>	Jul 25, 2008
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078942</u>	<u>002</u>	Jul 25, 2008
<u>AP</u>	MN PHARMS	<u>30MG/VIAL</u>	<u>A078300</u>	<u>001</u>	Mar 10, 2009
<u>AP</u>		<u>90MG/VIAL</u>	<u>A078300</u>	<u>002</u>	Mar 10, 2009
<u>AP</u>	MUSTAFA NEVZAT	<u>30MG/10ML (3MG/ML)</u>	<u>A078373</u>	<u>001</u>	Dec 23, 2008
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078373</u>	<u>002</u>	Dec 23, 2008
<u>AP</u>	PFIZER	<u>30MG/10ML (3MG/ML)</u>	<u>A078520</u>	<u>001</u>	Oct 31, 2008
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078520</u>	<u>002</u>	Oct 31, 2008
<u>AP</u>	PLIVA LACHEMA	<u>30MG/10ML (3MG/ML)</u>	<u>A078156</u>	<u>001</u>	Aug 19, 2008
<u>AP</u>		<u>60MG/10ML (6MG/ML)</u>	<u>A078156</u>	<u>002</u>	Aug 19, 2008
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A078156</u>	<u>003</u>	Aug 19, 2008
<u>AP</u>	SUN PHARMA GLOBAL	<u>30MG/VIAL</u>	<u>A077703</u>	<u>001</u>	Dec 24, 2008
<u>AP</u>		<u>90MG/VIAL</u>	<u>A077703</u>	<u>002</u>	Dec 24, 2008
<u>AP</u>	TEVA PARENTERAL	<u>30MG/10ML (3MG/ML)</u>	<u>A076153</u>	<u>001</u>	Mar 27, 2002
<u>AP</u>		<u>90MG/10ML (9MG/ML)</u>	<u>A076153</u>	<u>002</u>	Mar 27, 2002

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE, DELAYED RELEASE; ORAL

CREON

	ABBOTT PRODS	30,000USP UNITS;6,000USP UNITS;19,000USP UNITS	N020725	001	Apr 30, 2009
		60,000USP UNITS;12,000USP UNITS;38,000USP UNITS	N020725	002	Apr 30, 2009
+		120,000USP UNITS;24,000USP UNITS;76,000USP UNITS	N020725	003	Apr 30, 2009
	PANCREAZE				
	JANSSEN PHARMS	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
		70,000USP/ UNITS;16,800USP/ UNITS;40,000USP/ UNITS	N022523	004	Apr 12, 2010
+		61,000USP/ UNITS;21,000USP/ UNITS;37,000USP/ UNITS	N022523	003	Apr 12, 2010
	ZENPEP				
	APTALIS PHARMA US	27,000USP UNITS;5,000USP UNITS;17,000USP UNITS	N022210	001	Aug 27, 2009
		55,000USP UNITS;10,000USP UNITS;34,000USP UNITS	N022210	002	Aug 27, 2009
		82,000USP UNITS;15,000USP UNITS;51,000USP UNITS	N022210	003	Aug 27, 2009
+		109,000USP UNITS;20,000USP UNITS;68,000USP UNITS	N022210	004	Aug 27, 2009

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

<u>AP</u>	HOSPIRA	<u>1Mg/ML</u>	<u>A072320</u>	<u>001</u>	Jan 19, 1989
<u>AP</u> +	TEVA PARENTERAL	<u>1Mg/ML</u>	<u>A072759</u>	<u>001</u>	Jul 31, 1990
<u>AP</u> +		<u>2Mg/ML</u>	<u>A072760</u>	<u>001</u>	Jul 31, 1990

PRESCRIPTION DRUG PRODUCT LIST

3 - 328 (of 424)

PANTOPRAZOLE SODIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL
PROTONIX

+ WYETH PHARMS INC EQ 40MG BASE N022020 001 Nov 14, 2007
 INJECTABLE; IV (INFUSION)

+ WYETH PHARMS INC EQ 40MG BASE/VIAL N020988 001 Mar 22, 2001

TABLET, DELAYED RELEASE; ORAL

PANTOPRAZOLE SODIUM

<u>AB</u>	ACTAVIS TOTOWA	<u>EQ 20MG BASE</u>	<u>A090797</u> <u>001</u>	Feb 07, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090797</u> <u>002</u>	Feb 07, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A077619</u> <u>001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077619</u> <u>002</u>	Jan 19, 2011
<u>AB</u>	JUBILANT ORGANOSYS	<u>EQ 20MG BASE</u>	<u>A090901</u> <u>001</u>	Aug 30, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090901</u> <u>002</u>	Aug 30, 2011
<u>AB</u>	KUDCO IRELAND	<u>EQ 20MG BASE</u>	<u>A078281</u> <u>001</u>	Jan 20, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078281</u> <u>002</u>	Jan 20, 2011
<u>AB</u>	MATRIX LABS LTD	<u>EQ 20MG BASE</u>	<u>A090970</u> <u>001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090970</u> <u>002</u>	Jan 19, 2011
<u>AB</u>	SUN PHARMA GLOBAL	<u>EQ 20MG BASE</u>	<u>A077058</u> <u>001</u>	Sep 10, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077058</u> <u>002</u>	Sep 10, 2007
<u>AB</u>	TEVA	<u>EQ 20MG BASE</u>	<u>A077056</u> <u>001</u>	Aug 02, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077056</u> <u>002</u>	Aug 02, 2007
<u>AB</u>	TORRENT PHARMS	<u>EQ 20MG BASE</u>	<u>A090074</u> <u>001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A090074</u> <u>002</u>	Jan 19, 2011
<u>AB</u>	WOCKHARDT	<u>EQ 20MG BASE</u>	<u>A091231</u> <u>001</u>	Jan 19, 2011
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A091231</u> <u>002</u>	Jan 19, 2011
	<u>PROTONIX</u>			
<u>AB</u>	WYETH PHARMS INC	<u>EQ 20MG BASE</u>	<u>N020987</u> <u>002</u>	Jun 12, 2001
<u>AB</u>	+	<u>EQ 40MG BASE</u>	<u>N020987</u> <u>001</u>	Feb 02, 2000

PARICALCITOL

CAPSULE; ORAL
ZEMPLAR

ABBOTT 1MCG N021606 001 May 26, 2005
 2MCG N021606 002 May 26, 2005
 + 4MCG N021606 003 May 26, 2005

INJECTABLE; INJECTION

PARICALCITOL

<u>AP</u>	SANDOZ CANADA INC	<u>0.002MG/ML</u>	<u>A091108</u> <u>001</u>	Jul 27, 2011
<u>AP</u>		<u>0.005MG/ML</u>	<u>A091108</u> <u>002</u>	Jul 27, 2011
	<u>ZEMPLAR</u>			
<u>AP</u>	+	<u>ABBOTT</u>	<u>0.002MG/ML</u>	Feb 01, 2000
<u>AP</u>	+		<u>0.005MG/ML</u>	Apr 17, 1998

PAROMOMYCIN SULFATE

CAPSULE; ORAL
PAROMOMYCIN SULFATE

<u>AA</u>	+	CARACO	<u>EQ 250MG BASE</u>	<u>A064171</u> <u>001</u>	Jun 30, 1997
<u>AA</u>		HERITAGE PHARMS INC	<u>EQ 250MG BASE</u>	<u>A065173</u> <u>001</u>	Dec 14, 2007

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL
PAROXETINE HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>EQ 10MG BASE/5ML</u>	<u>A077395</u> <u>001</u>	Dec 05, 2006
<u>AB</u>	+	<u>PAXIL</u>	<u>N020710</u> <u>001</u>	Jun 25, 1997
		<u>GLAXOSMITHKLINE</u>	<u>EQ 10MG BASE/5ML</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 329 (of 424)

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

AB	ALPHAPHARM	EQ 10MG BASE	A075716 001	Mar 08, 2004
AB		EQ 20MG BASE	A075716 002	Mar 08, 2004
AB		EQ 30MG BASE	A075716 003	Mar 08, 2004
AB		EQ 40MG BASE	A075716 004	Mar 08, 2004
AB	APOTEX	EQ 10MG BASE	A075356 001	Jul 30, 2003
AB		EQ 20MG BASE	A075356 002	Jul 30, 2003
AB		EQ 30MG BASE	A075356 003	Jul 30, 2003
AB		EQ 40MG BASE	A075356 004	Jul 30, 2003
AB	AUROBINDO PHARMA	EQ 10MG BASE	A078406 001	Jul 25, 2007
AB		EQ 20MG BASE	A078406 002	Jul 25, 2007
AB		EQ 30MG BASE	A078406 003	Jul 25, 2007
AB		EQ 40MG BASE	A078406 004	Jul 25, 2007
AB	CARACO	EQ 10MG BASE	A078194 001	Jun 29, 2007
AB		EQ 20MG BASE	A078194 002	Jun 29, 2007
AB		EQ 30MG BASE	A078194 003	Jun 29, 2007
AB		EQ 40MG BASE	A078194 004	Jun 29, 2007
AB	MYLAN	EQ 10MG BASE	A078902 001	Mar 13, 2008
AB		EQ 20MG BASE	A078902 002	Mar 13, 2008
AB		EQ 30MG BASE	A078902 003	Mar 13, 2008
AB		EQ 40MG BASE	A078902 004	Mar 13, 2008
AB	TEVA	EQ 10MG BASE	A076618 001	Aug 15, 2005
AB		EQ 20MG BASE	A076618 002	Aug 15, 2005
AB		EQ 30MG BASE	A076618 003	Aug 15, 2005
AB		EQ 40MG BASE	A076618 004	Aug 15, 2005
AB	ZYDUS PHARMS USA	EQ 10MG BASE	A077584 001	Mar 07, 2007
AB		EQ 20MG BASE	A077584 002	Mar 07, 2007
AB		EQ 30MG BASE	A077584 003	Mar 07, 2007
AB		EQ 40MG BASE	A077584 004	Mar 07, 2007
	PAXIL			
AB	GLAXOSMITHKLINE	EQ 10MG BASE	N020031 001	Dec 29, 1992
AB		EQ 20MG BASE	N020031 002	Dec 29, 1992
AB		EQ 30MG BASE	N020031 003	Dec 29, 1992
AB	+	EQ 40MG BASE	N020031 005	Dec 29, 1992

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

AB	MYLAN	EQ 12.5MG BASE	A077873 001	Jun 29, 2007
AB		EQ 25MG BASE	A077873 002	Jun 29, 2007
AB		EQ 37.5MG BASE	A091427 001	Apr 14, 2011
	PAXIL CR			
AB	GLAXOSMITHKLINE	EQ 12.5MG BASE	N020936 001	Feb 16, 1999
AB		EQ 25MG BASE	N020936 002	Feb 16, 1999
AB	+	EQ 37.5MG BASE	N020936 003	Dec 06, 2000

PAROXETINE MESYLATE

TABLET; ORAL

PEXEVA

NOVEN THERAP

		EQ 10MG BASE	N021299 001	Jul 03, 2003
		EQ 20MG BASE	N021299 002	Jul 03, 2003
		EQ 30MG BASE	N021299 003	Jul 03, 2003
		EQ 40MG BASE	N021299 004	Jul 03, 2003

PAZOPANIB HYDROCHLORIDE

TABLET; ORAL

VOTRIENT

GLAXOSMITHKLINE

		EQ 200MG BASE	N022465 001	Oct 19, 2009
	+	EQ 400MG BASE	N022465 002	Oct 19, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 330 (of 424)

PEGADEMASE BOVINE

INJECTABLE; INJECTION
 ADAGEN
 + SIGMA TAU 250 UNITS/ML N019818 001 Mar 21, 1990

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL
 MACUGEN
 + EYETECH INC 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

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

PEMIROLAST POTASSIUM

SOLUTION/DROPS; OPHTHALMIC
 + SANTEN 0.1% N021079 001 Sep 24, 1999

PENBUTOLOL SULFATE

TABLET; ORAL
 LEVATOL
 + SCHWARZ PHARMA 20MG N018976 004 Jan 05, 1989

PENCICLOVIR SODIUM

CREAM; TOPICAL
 DENAVIR
 + Denco Asset 1% N020629 001 Sep 24, 1996

PENICILLAMINE

CAPSULE; ORAL
 CUPRIMINE
 + ATON 250MG N019853 001

TABLET; ORAL
 DEPEN
 + MEDA PHARMS 250MG N019854 001

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION
 BICILLIN L-A
 BC + KING PHARMS 600,000 UNITS/ML N050141 001
 PERMAPEN
 BC PFIZER 600,000 UNITS/ML A060014 001
 BICILLIN L-A
 + KING PHARMS 300,000 UNITS/ML N050141 003

PRESCRIPTION DRUG PRODUCT LIST

3 - 331 (of 424)

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION

BICILLIN C-R

+ KING PHARMS	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	900,000 UNITS/2ML;300,000 UNITS/2ML	N050138 003

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

<u>AP</u>	APP PHARMS	<u>5,000,000 UNITS/VIAL</u>	<u>A065448 001</u>	Aug 18, 2009
<u>AP</u>		<u>20,000,000 UNITS/VIAL</u>	<u>A065448 002</u>	Aug 18, 2009
<u>AP</u>	HANFORD GC	<u>5,000,000 UNITS/VIAL</u>	<u>A065149 002</u>	Jul 23, 2009
<u>AP</u>		<u>20,000,000 UNITS/VIAL</u>	<u>A065149 003</u>	Jul 23, 2009
<u>AP</u>	SANDOZ	<u>5,000,000 UNITS/VIAL</u>	<u>A065079 002</u>	Aug 30, 2002
<u>AP</u>		<u>20,000,000 UNITS/VIAL</u>	<u>A065079 003</u>	Aug 30, 2002
	<u>PFIZERPEN</u>			
<u>AP</u>	+ PFIZER	<u>5,000,000 UNITS/VIAL</u>	<u>A060657 002</u>	
<u>AP</u>	+	<u>20,000,000 UNITS/VIAL</u>	<u>A060657 003</u>	
	PENICILLIN G POTASSIUM			
	HANFORD GC	1,000,000 UNITS/VIAL	A065149 001	Jul 23, 2009
	PENICILLIN G POTASSIUM IN PLASTIC CONTAINER			
	+ BAXTER HLTHCARE	20,000 UNITS/ML	N050638 001	Jun 25, 1990
	+	40,000 UNITS/ML	N050638 002	Jun 25, 1990
	+	60,000 UNITS/ML	N050638 003	Jun 25, 1990

PENICILLIN G PROCAINE

INJECTABLE; INJECTION

PENICILLIN G PROCAINE

+ KING PHARMS	300,000 UNITS/ML	A060101 002
+	600,000 UNITS/ML	A060101 001

PENICILLIN G SODIUM

INJECTABLE; IM-IV

PENICILLIN G SODIUM

+ SANDOZ	5,000,000 UNITS/VIAL	A065068 001	Feb 26, 2001
----------	----------------------	-------------	--------------

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN V POTASSIUM

<u>AA</u>	DAVA PHARMS INC	<u>EQ 125MG BASE/5ML</u>	<u>A062981 001</u>	Feb 10, 1989
<u>AA</u>		<u>EQ 250MG BASE/5ML</u>	<u>A062981 002</u>	Feb 10, 1989
	<u>PENICILLIN-VK</u>			
<u>AA</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>A060456 001</u>	
<u>AA</u>	+	<u>EQ 250MG BASE/5ML</u>	<u>A060456 002</u>	

TABLET; ORAL

PENICILLIN V POTASSIUM

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A065435 001</u>	Apr 29, 2008
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065435 002</u>	Apr 29, 2008
<u>AB</u>	DAVA PHARMS INC	<u>EQ 250MG BASE</u>	<u>A062936 001</u>	Nov 25, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062935 001</u>	Nov 23, 1988
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>A064071 001</u>	Nov 30, 1995
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>A064071 002</u>	Nov 30, 1995
	<u>PENICILLIN-VK</u>			
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A060711 002</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A060711 003</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 332 (of 424)

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION NEBUPENT			
+ APP PHARMS	300MG/VIAL	N019887 001	Jun 15, 1989
INJECTABLE; INJECTION <u>PENTAM</u>			
<u>AP</u> + APP PHARMS	<u>300MG/VIAL</u>	<u>N019264 001</u>	Oct 16, 1984
<u>PENTAMIDINE ISETHIONATE</u>			
<u>AP</u> WATSON LABS	<u>300MG/VIAL</u>	<u>A074303 001</u>	Aug 17, 1995

PENTAZOCINE LACTATE

INJECTABLE; INJECTION TALWIN			
+ HOSPIRA	EQ 30MG BASE/ML	N016194 001	

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION NEMBUTAL SODIUM			
+ LUNDBECK INC	50MG/ML	A083246 001	

PENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL ELMIRON			
+ JANSSEN PHARMS	100MG	N020193 001	Sep 26, 1996

PENTOSTATIN

INJECTABLE; INJECTION <u>NIPENT</u>			
<u>AP</u> + HOSPIRA INC	<u>10MG/VIAL</u>	<u>N020122 001</u>	Oct 11, 1991
<u>PENTOSTATIN</u>			
<u>AP</u> BEDFORD LABS	<u>10MG/VIAL</u>	<u>A077841 001</u>	Aug 07, 2007

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL <u>PENTOXIFYLLINE</u>			
<u>AB</u> APOTEX	<u>400MG</u>	<u>A075191 001</u>	Jun 09, 1999
<u>AB</u> IMPAX LABS	<u>400MG</u>	<u>A075093 001</u>	Aug 10, 1999
<u>AB</u> MYLAN	<u>400MG</u>	<u>A074425 001</u>	Jul 08, 1997
<u>AB</u> PLIVA	<u>400MG</u>	<u>A074874 001</u>	May 25, 1999
<u>AB</u> VALEANT INTL	<u>400MG</u>	<u>A075028 001</u>	Jul 20, 1998
<u>AB</u> WATSON LABS	<u>400MG</u>	<u>A075107 001</u>	Sep 04, 1998
<u>PENTOXIL</u>			
<u>AB</u> UPSHER SMITH	<u>400MG</u>	<u>A074962 001</u>	Mar 31, 1999
<u>TRENTAL</u>			
<u>AB</u> + SANOFI AVENTIS US	<u>400MG</u>	<u>N018631 001</u>	Aug 30, 1984

PERFLUTREN

INJECTABLE; INTRAVENOUS DEFINITY			
+ LANTHEUS MEDCL	6.52MG/ML	N021064 001	Jul 31, 2001

PERINDOPRIL ERBUMINE

TABLET; ORAL <u>ACEON</u>			
<u>AB</u> ABBOTT PRODS	<u>2MG</u>	<u>N020184 001</u>	Dec 30, 1993
<u>AB</u>	<u>4MG</u>	<u>N020184 002</u>	Dec 30, 1993
<u>AB</u> +	<u>8MG</u>	<u>N020184 003</u>	Dec 30, 1993
<u>PERINDOPRIL ERBUMINE</u>			
<u>AB</u> APOTEX	<u>2MG</u>	<u>A090463 001</u>	Aug 30, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 333 (of 424)

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

<u>AB</u>	APOTEX	<u>4MG</u>	<u>A090463</u> <u>002</u>	Aug 30, 2010
<u>AB</u>		<u>8MG</u>	<u>A090463</u> <u>003</u>	Aug 30, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>2MG</u>	<u>A079070</u> <u>001</u>	Nov 10, 2009
<u>AB</u>		<u>4MG</u>	<u>A079070</u> <u>002</u>	Nov 10, 2009
<u>AB</u>		<u>8MG</u>	<u>A079070</u> <u>003</u>	Nov 10, 2009
<u>AB</u>	IVAX PHARMS	<u>2MG</u>	<u>A078138</u> <u>001</u>	Nov 10, 2009
<u>AB</u>		<u>4MG</u>	<u>A078138</u> <u>002</u>	Nov 10, 2009
<u>AB</u>		<u>8MG</u>	<u>A078138</u> <u>003</u>	Nov 10, 2009
<u>AB</u>	LUPIN LTD	<u>2MG</u>	<u>A078263</u> <u>001</u>	Jan 27, 2010
<u>AB</u>		<u>4MG</u>	<u>A078263</u> <u>002</u>	Jan 27, 2010
<u>AB</u>		<u>8MG</u>	<u>A078263</u> <u>003</u>	Jan 27, 2010
<u>AB</u>	ROXANE	<u>2MG</u>	<u>A090072</u> <u>001</u>	Nov 10, 2009
<u>AB</u>		<u>4MG</u>	<u>A090072</u> <u>002</u>	Nov 10, 2009
<u>AB</u>		<u>8MG</u>	<u>A090072</u> <u>003</u>	Nov 10, 2009

PERMETHRIN

CREAM; TOPICAL

ELIMITE

<u>AB</u>	+ ALLERGAN	<u>5%</u>	<u>N019855</u> <u>001</u>	Aug 25, 1989
	<u>PERMETHRIN</u>			
<u>AB</u>	ACTAVIS MID ATLANTIC	<u>5%</u>	<u>A074806</u> <u>001</u>	Jan 23, 1998
<u>AB</u>	PERRIGO NEW YORK	<u>5%</u>	<u>A076369</u> <u>001</u>	Apr 21, 2003

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

<u>AB</u>	SANDOZ	<u>2MG</u>	<u>A089683</u> <u>001</u>	Dec 08, 1988
<u>AB</u>		<u>4MG</u>	<u>A089684</u> <u>001</u>	Dec 08, 1988
<u>AB</u>		<u>8MG</u>	<u>A089685</u> <u>001</u>	Dec 08, 1988
<u>AB</u>	+	<u>16MG</u>	<u>A089686</u> <u>001</u>	Dec 08, 1988
<u>AB</u>	VINTAGE PHARMS	<u>2MG</u>	<u>A040226</u> <u>001</u>	Dec 31, 1998
<u>AB</u>		<u>4MG</u>	<u>A040226</u> <u>002</u>	Dec 31, 1998
<u>AB</u>		<u>8MG</u>	<u>A040226</u> <u>003</u>	Dec 31, 1998
<u>AB</u>		<u>16MG</u>	<u>A040226</u> <u>004</u>	Dec 31, 1998

PHENDIMETRAZINE TARTRATECAPSULE, EXTENDED RELEASE; ORAL
BONTRIL

<u>BC</u>	VALEANT	105MG	<u>A088021</u> <u>001</u>	Sep 21, 1982
	PHENDIMETRAZINE TARTRATE			
<u>BC</u>	+	SANDOZ	<u>N018074</u> <u>001</u>	

TABLET; ORAL

BONTRIL PDM

<u>AA</u>	+	VALEANT	<u>35MG</u>	<u>A085272</u> <u>001</u>
		<u>PHENDIMETRAZINE TARTRATE</u>		
<u>AA</u>		KVK TECH	<u>35MG</u>	<u>A091042</u> <u>001</u>
<u>AA</u>		MIKAH PHARMA	<u>35MG</u>	<u>A040762</u> <u>001</u>
<u>AA</u>		MIKART	<u>35MG</u>	<u>A089452</u> <u>001</u>
<u>AA</u>		SANDOZ	<u>35MG</u>	<u>A085588</u> <u>001</u>

PHENELZINE SULFATE

TABLET; ORAL

NARDIL

<u>AB</u>	+	PARKE DAVIS	<u>EQ 15MG BASE</u>	<u>N011909</u> <u>002</u>
		<u>PHENELZINE SULFATE</u>		
<u>AB</u>		NOVEL LABS INC	<u>EQ 15MG BASE</u>	<u>A200181</u> <u>001</u>

Dec 08, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 334 (of 424)

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL
 DIBENZYLINE
 + WELLSPRING PHARM 10MG N008708 001

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

ADIPEX-P

<u>AA</u>	+ TEVA	<u>37.5MG</u>	<u>A088023 001</u>	Aug 02, 1983
<u>PHENTERMINE HYDROCHLORIDE</u>				
<u>AA</u>	BARR	<u>15MG</u>	<u>A090591 001</u>	Mar 18, 2010
<u>AA</u>		<u>30MG</u>	<u>A090591 002</u>	Mar 18, 2010
<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>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>	MUTUAL PHARM	<u>30MG</u>	<u>A040525 001</u>	Oct 23, 2003
<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>	

TABLET; ORAL

ADIPEX-P

<u>AA</u>	+ TEVA	<u>37.5MG</u>	<u>A085128 001</u>	
<u>PHENTERMINE HYDROCHLORIDE</u>				
<u>AA</u>	ACTAVIS ELIZABETH	<u>37.5MG</u>	<u>A040276 001</u>	Nov 25, 1998
<u>AA</u>	BARR	<u>37.5MG</u>	<u>A090470 001</u>	Aug 31, 2009
<u>AA</u>	CARACO	<u>37.5MG</u>	<u>A040790 001</u>	Aug 21, 2007
<u>AA</u>	ELITE LABS	<u>37.5MG</u>	<u>A200272 001</u>	Jan 31, 2011
<u>AA</u>	KVK TECH	<u>37.5MG</u>	<u>A040876 001</u>	Mar 31, 2008
<u>AA</u>	LANNETT	<u>37.5MG</u>	<u>A040555 001</u>	Apr 15, 2005
<u>AA</u>	MUTUAL PHARM	<u>37.5MG</u>	<u>A040526 001</u>	Oct 23, 2003
<u>AA</u>	VINTAGE PHARMS	<u>37.5MG</u>	<u>A040377 001</u>	Jan 04, 2002
<u>PHENTERMINE HYDROCHLORIDE</u>				
+ SANDOZ	30MG		A088605 001	Sep 28, 1987

TABLET, ORALLY DISINTEGRATING; ORAL

SUPRENZA

CITIUS PHARMS	15MG	N202088 001	Jun 13, 2011
+	30MG	N202088 002	Jun 13, 2011

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

PHENTERMINE RESIN COMPLEX

LANNETT HOLDINGS INC	EQ 15MG BASE	A040872 001	Jul 28, 2011
	EQ 30MG BASE	A040872 002	Jul 28, 2011

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

<u>AP</u>	BEDFORD	<u>5MG/VIAL</u>	<u>A040235 001</u>	Mar 11, 1998
<u>REGITINE</u>				
<u>AP</u>	+ NOVARTIS	<u>5MG/VIAL</u>	<u>N008278 003</u>	
ORaverse				
+ SEPTODONT HOLDING	0.4MG/1.7ML		N022159 001	May 09, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 335 (of 424)

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

<u>AA</u>	VINTAGE	<u>5MG/5ML; 6.25MG/5ML</u>	<u>A040654</u> <u>001</u>	Dec 07, 2006
<u>AA</u>	<u>PROMETH VC PLAIN</u>			
<u>AA</u>	+ ACTAVIS MID ATLANTIC	<u>5MG/5ML; 6.25MG/5ML</u>	<u>A088761</u> <u>001</u>	Nov 08, 1984
<u>AA</u>	<u>PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE</u>			
<u>AA</u>	AMNEAL PHARMS	<u>5MG/5ML; 6.25MG/5ML</u>	<u>A040902</u> <u>001</u>	Aug 25, 2009

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-125

<u>AB</u>	+ PARKE DAVIS	<u>125MG/5ML</u>	<u>N008762</u> <u>001</u>	
<u>AB</u>	<u>PHENYTOIN</u>			
<u>AB</u>	TARO	<u>125MG/5ML</u>	<u>A040521</u> <u>001</u>	Mar 08, 2004
<u>AB</u>	VISTAPHARM	<u>125MG/5ML</u>	<u>A040342</u> <u>001</u>	Jan 31, 2001
<u>AB</u>		<u>125MG/5ML</u>	<u>A040610</u> <u>001</u>	Aug 18, 2005
<u>AB</u>	WOCKHARDT	<u>125MG/5ML</u>	<u>A040420</u> <u>001</u>	Apr 19, 2002

TABLET, CHEWABLE; ORAL

DILANTIN

+ PFIZER PHARMS

50MG

A084427 001

PHENYTOIN SODIUM

CAPSULE; ORAL

DILANTIN

<u>AB</u>	+ PARKE DAVIS	<u>30MG EXTENDED</u>	<u>A084349</u> <u>001</u>	
<u>AB</u>	+	<u>100MG EXTENDED</u>	<u>A084349</u> <u>002</u>	
<u>AB</u>	<u>EXTENDED PHENYTOIN SODIUM</u>			
<u>AB</u>	AMNEAL PHARMS NY	<u>100MG EXTENDED</u>	<u>A040765</u> <u>001</u>	Nov 12, 2008
<u>AB</u>	MYLAN	<u>100MG EXTENDED</u>	<u>A040298</u> <u>001</u>	Dec 28, 1998
<u>AB</u>	SUN PHARM INDNS	<u>200MG EXTENDED</u>	<u>A040731</u> <u>001</u>	Jun 30, 2008
<u>AB</u>		<u>300MG EXTENDED</u>	<u>A040731</u> <u>002</u>	Jun 30, 2008
<u>AB</u>	SUN PHARM INDNS (IN)	<u>100MG EXTENDED</u>	<u>A040621</u> <u>001</u>	Dec 11, 2006
<u>AB</u>	TARO	<u>100MG EXTENDED</u>	<u>A040684</u> <u>001</u>	Sep 05, 2006
<u>AB</u>	WOCKHARDT	<u>30MG EXTENDED</u>	<u>A040759</u> <u>001</u>	Dec 18, 2007
<u>AB</u>	WOCKHARDT USA	<u>100MG EXTENDED</u>	<u>A040732</u> <u>001</u>	Jan 30, 2008
<u>AB</u>	<u>PHENYTEK</u>			
<u>AB</u>	MYLAN	<u>200MG EXTENDED</u>	<u>A040298</u> <u>002</u>	Dec 06, 2001
<u>AB</u>	+	<u>300MG EXTENDED</u>	<u>A040298</u> <u>003</u>	Dec 06, 2001

INJECTABLE; INJECTION

PHENYTOIN SODIUM

<u>AP</u>	+ BAXTER HLTHCARE	<u>50MG/ML</u>	<u>A084307</u> <u>001</u>	
<u>AP</u>	HOSPIRA	<u>50MG/ML</u>	<u>A089521</u> <u>001</u>	Mar 17, 1987
<u>AP</u>		<u>50MG/ML</u>	<u>A089744</u> <u>001</u>	Dec 18, 1987
<u>AP</u>	LUITPOLD	<u>50MG/ML</u>	<u>A040781</u> <u>001</u>	Dec 04, 2007
<u>AP</u>	X-GEN PHARMS	<u>50MG/ML</u>	<u>A040573</u> <u>001</u>	Sep 13, 2006

PHYTONADIONE

INJECTABLE; INJECTION

PHYTONADIONE

BP	INTL MEDICATION	1MG/0.5ML	A083722 001	
	VITAMIN K1			
BP	+ HOSPIRA	1MG/0.5ML	A087954 001	Jul 25, 1983
	VITAMIN K1			
	+ HOSPIRA	10MG/ML	A087955 001	Jul 25, 1983
	TABLET; ORAL			
	MEPHYTON			
	+ BIOVAIL TECHNOLOGIES	5MG	N010104 003	

PRESCRIPTION DRUG PRODUCT LIST

3 - 336 (of 424)

PILOCARPINE HYDROCHLORIDE

GEL; OPHTHALMIC PILOPINE HS + ALCON	4%	N018796 001	Oct 01, 1984
SOLUTION; OPHTHALMIC ISOPTO CARPINE ALCON RES	1%	N200890 001	Jun 22, 2010
	2%	N200890 002	Jun 22, 2010
+	4%	N200890 003	Jun 22, 2010
TABLET; ORAL PILOCARPINE HYDROCHLORIDE			
<u>AB</u> COREPHARMA	<u>5MG</u>	<u>A076746</u> <u>001</u>	Nov 16, 2004
<u>AB</u> IMPAX LABS	<u>5MG</u>	<u>A077248</u> <u>001</u>	Mar 31, 2006
<u>AB</u>	<u>7.5MG</u>	<u>A077248</u> <u>002</u>	Mar 31, 2006
<u>AB</u> LANNETT	<u>5MG</u>	<u>A077220</u> <u>001</u>	Oct 14, 2005
<u>AB</u>	<u>7.5MG</u>	<u>A077220</u> <u>002</u>	May 06, 2009
<u>AB</u> ROXANE	<u>5MG</u>	<u>A076963</u> <u>001</u>	Dec 22, 2004
<u>AB</u>	<u>7.5MG</u>	<u>A076963</u> <u>002</u>	Feb 27, 2007
SALAGEN			
<u>AB</u> EISAI INC	<u>5MG</u>	<u>N020237</u> <u>001</u>	Mar 22, 1994
<u>AB</u> +	<u>7.5MG</u>	<u>N020237</u> <u>002</u>	Apr 18, 2003

PIMECROLIMUS

CREAM; TOPICAL ELIDEL + VALEANT INTL	1%	N021302 001	Dec 13, 2001
--	----	-------------	--------------

PIMOZIDE

TABLET; ORAL ORAP TEVA	1MG	N017473 003	Aug 27, 1997
+	2MG	N017473 001	Jul 31, 1984

PINDOLOL

TABLET; ORAL PINDOLOL			
<u>AB</u> MYLAN	<u>5MG</u>	<u>A074019</u> <u>001</u>	Sep 03, 1992
<u>AB</u> +	<u>10MG</u>	<u>A074019</u> <u>002</u>	Sep 03, 1992
<u>AB</u> WATSON LABS	<u>5MG</u>	<u>A074437</u> <u>001</u>	Feb 27, 1995
<u>AB</u>	<u>10MG</u>	<u>A074437</u> <u>002</u>	Feb 27, 1995

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL ACTOS TAKEDA PHARMS NA	EQ 15MG BASE	N021073 001	Jul 15, 1999
	EQ 30MG BASE	N021073 002	Jul 15, 1999
+	EQ 45MG BASE	N021073 003	Jul 15, 1999

PIPERACILLIN SODIUM

INJECTABLE; INJECTION PIPERACILLIN			
+ ISTITUTO BIOCHIMICO	EQ 2GM BASE/VIAL	A065114 001	Nov 14, 2003
+	EQ 3GM BASE/VIAL	A065114 002	Nov 14, 2003
+	EQ 4GM BASE/VIAL	A065114 003	Nov 14, 2003
+	EQ 40GM BASE/VIAL	A065157 001	Jul 12, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 337 (of 424)

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

PIPERACILLIN AND TAZOBACTAM

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065498 001</u>	May 23, 2011
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065498 002</u>	May 23, 2011
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065498 003</u>	May 23, 2011
<u>AP</u>	HOSPIRA INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065386 001</u>	Sep 15, 2009
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065386 002</u>	Sep 15, 2009
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065386 003</u>	Sep 15, 2009
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A065446 001</u>	Sep 15, 2009
<u>AP</u>	INSTITUTO BIOCHEMICO	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065523 001</u>	May 31, 2011
<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>	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>ZOSYN</u>				
<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

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

+ MEDICIS

EQ 0.2MG BASE/INH

N020014 001 Nov 30, 1992

PIROXICAM

CAPSULE; ORAL

FELDENNE

<u>AB</u>	PFIZER	<u>10MG</u>	<u>N018147 002</u>	Apr 06, 1982
<u>AB</u>	+	<u>20MG</u>	<u>N018147 003</u>	Apr 06, 1982
	<u>PIROXICAM</u>			
<u>AB</u>	MUTUAL PHARM	<u>10MG</u>	<u>A073535 001</u>	Mar 12, 1993
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A074102 001</u>	Jul 31, 1992
<u>AB</u>		<u>20MG</u>	<u>A074102 002</u>	Jul 31, 1992
<u>AB</u>	NOSTRUM LABS	<u>10MG</u>	<u>A074116 001</u>	Jun 15, 1993
<u>AB</u>		<u>20MG</u>	<u>A074118 001</u>	Jun 15, 1993
<u>AB</u>	TEVA	<u>10MG</u>	<u>A074131 001</u>	Dec 11, 1992
<u>AB</u>		<u>20MG</u>	<u>A074131 002</u>	Dec 11, 1992
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A074287 001</u>	May 16, 1996
<u>AB</u>		<u>10MG</u>	<u>A074460 001</u>	Sep 29, 1995
<u>AB</u>		<u>20MG</u>	<u>A074287 002</u>	May 16, 1996
<u>AB</u>		<u>20MG</u>	<u>A074460 002</u>	Sep 29, 1995

PITAVASTATIN CALCIUM

TABLET; ORAL

LIVALO

KOWA CO

EQ 1MG BASE

N022363 001 Aug 03, 2009

EQ 2MG BASE

N022363 002 Aug 03, 2009

+

EQ 4MG BASE

N022363 003 Aug 03, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 338 (of 424)

PLERIXAFOR

SOLUTION; SUBCUTANEOUS
 MOZOBIL
 + GENZYME 24MG/1.2ML (20MG/ML) N022311 001 Dec 15, 2008

PODOFILOX

GEL; TOPICAL CONDYLOX + WATSON PHARMS	0.5%	N020529 001	Mar 13, 1997
SOLUTION; TOPICAL <u>CONDYLOX</u> <u>PODOFILOX</u>	<u>0.5%</u>	<u>N019795 001</u>	Dec 13, 1990
AT + WATSON PHARMS	<u>0.5%</u>	<u>A075600 001</u>	Jan 29, 2002
AT PADDOCK LLC	<u>0.5%</u>	<u>A090184 001</u>	Jul 21, 2010
AT PRECISION DERMAT	<u>0.5%</u>		

POLIDOCANOL

SOLUTION; INTRAVENOUS
 ASCLERA
 CHEMISCH FBRK KRSSLR 10MG/2ML (5MG/ML)
 + 20MG/2ML (10MG/ML) N021201 001 Mar 30, 2010
 N021201 002 Mar 30, 2010

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL <u>GLYCOLAX</u>			
AA KREMERS URBAN PHARMS	<u>17GM/SCOOPFUL</u>	<u>A076652 001</u>	Jul 02, 2004
AA BRECKENRIDGE PHARM	<u>17GM/SCOOPFUL</u>	<u>A077736 001</u>	May 26, 2006
AA NEXGEN PHARMA INC	<u>17GM/SCOOPFUL</u>	<u>A077706 001</u>	Sep 27, 2006
AA PADDOCK LLC	<u>17GM/SCOOPFUL</u>	<u>A077893 001</u>	May 26, 2006

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

FOR SOLUTION; ORAL <u>LAX-LYTE WITH FLAVOR PACKS</u>			
AA PADDOCK LLC	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A079232 001</u>	Feb 25, 2010
AA + BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N019797 001</u>	Apr 22, 1991
AA + BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N019797 002</u>	Nov 18, 1994
AA MYLAN	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A090409 001</u>	Apr 02, 2010
AA NOVEL LABS INC	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A090019 001</u>	May 27, 2009
AA MEDA PHARMS	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>A076491 001</u>	Feb 05, 2004

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

FOR SOLUTION; ORAL <u>CLENZ-LYTE</u>			
AA PADDOCK LLC	<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>A090769 001</u>	Jun 07, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 339 (of 424)

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

COLYTE

<u>AA</u>	+ MEDA PHARMS	<u>227.1GM/BOT; 2.82GM/BOT; 6.36GM/BOT; 5.53G M/BOT; 21.5GM/BOT</u>	<u>N018983 010</u>	Jan 31, 1989
<u>AA</u>	+	<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	<u>N018983 007</u>	Jun 12, 1987
		<u>COLYTE WITH FLAVOR PACKS</u>		
<u>AA</u>	+ MEDA PHARMS	<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	<u>N018983 012</u>	Oct 08, 1998
		<u>COLYTE-FLAVORED</u>		
<u>AA</u>	+ MEDA PHARMS	<u>227.1GM/BOT; 2.82GM/BOT; 6.36GM/BOT; 5.53G M/BOT; 21.5GM/BOT</u>	<u>N018983 008</u>	Nov 14, 1991
<u>AA</u>	+	<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	<u>N018983 009</u>	Nov 14, 1991
		<u>GOLYTELY</u>		
<u>AA</u>	+ BRAINTREE	<u>236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT</u>	<u>N019011 001</u>	Jul 13, 1984
		<u>PEG 3350 AND ELECTROLYTES</u>		
<u>AA</u>	MYLAN	<u>236GM; 2.97GM; 6.74GM; 5.86GM; 22.74GM</u>	<u>A090928 001</u>	Jan 28, 2010
<u>AA</u>	NOVEL LABS INC	<u>236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT</u>	<u>A090231 001</u>	Jun 01, 2009
<u>AA</u>		<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	<u>A090186 001</u>	Jun 01, 2009
		<u>POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES</u>		
<u>AA</u>	PADDOCK LLC	<u>240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT</u>	<u>A090712 001</u>	Feb 25, 2010
		<u>GOLYTELY</u>		
	+ 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

<u>AP</u>	SAGENT STRIDES	<u>EQ 500,000 UNITS BASE/VIAL</u>	<u>A090110 001</u>	Jun 29, 2011
<u>AP</u>	POLYMYXIN B SULFATE	<u>EQ 500,000 U BASE/VIAL</u>	<u>A065372 001</u>	Jan 10, 2008
<u>AP</u>	+ BEDFORD	<u>EQ 500,000 U BASE/VIAL</u>	<u>A060716 001</u>	

X GEN PHARMSEQ 500,000 U BASE/VIALA063000 001

Sep 30, 1994

POWDER; FOR RX COMPOUNDING

POLY-RX

+ X GEN PHARMS

100,000,000 UNITS/BOT

A061578 001

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC

POLYTRIM

<u>AT</u>	+ ALLERGAN	<u>10,000 UNITS/ML; EQ 1MG BASE/ML</u>	<u>N050567 001</u>	Oct 20, 1988
		<u>TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE</u>		
<u>AT</u>	BAUSCH AND LOMB	<u>10,000 UNITS/ML; EQ 1MG BASE/ML</u>	<u>A064120 001</u>	Feb 14, 1997

FALCON PHARMS10,000 UNITS/ML; EQ 1MG BASE/MLA064211 001

Apr 13, 1998

TAYLOR PHARMA10,000 UNITS/ML; EQ 1MG BASE/MLA065006 001

Dec 17, 1998

PORACTANT ALFA

SUSPENSION; INTRATRACHEAL

CUROSURF

+ CORNERSTONE THERAP

80MG/ML

N020744 001

Nov 18, 1999

PORFIMER SODIUM

INJECTABLE; INJECTION

PHOTOFRIN

PINNACLE BIOLGS

75MG/VIAL

N020451 001

Dec 27, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 340 (of 424)

POSACONAZOLE

SUSPENSION; ORAL
 NOXAFL
 + SCHERING 40MG/ML N022003 001 Sep 15, 2006

POTASSIUM ACETATE

INJECTABLE; INJECTION
 POTASSIUM ACETATE IN PLASTIC CONTAINER
 + HOSPIRA 2MEQ/ML N018896 001 Jul 20, 1984

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL			
<u>MICRO-K</u>			
<u>AB</u>	NESHER PHARMS	<u>8MEQ</u>	<u>N018238 001</u>
<u>MICRO-K 10</u>			
<u>AB</u>	NESHER PHARMS	<u>10MEQ</u>	<u>N018238 002</u> May 14, 1984
<u>POTASSIUM CHLORIDE</u>			
<u>AB</u>	PADDOCK LLC	<u>8MEQ</u>	<u>A200185 001</u> May 18, 2011
<u>AB</u>		<u>10MEQ</u>	<u>A200185 002</u> May 18, 2011
<u>AB</u>	WATSON LABS FLORIDA	<u>8MEQ</u>	<u>A077419 001</u> Jun 02, 2008
<u>AB</u> +		<u>10MEQ</u>	<u>A077419 002</u> Jun 02, 2008
INJECTABLE; INJECTION			
<u>POTASSIUM CHLORIDE</u>			
<u>AP</u>	APP PHARMS	<u>2MEQ/ML</u>	<u>A080225 001</u>
<u>AP</u>	B BRAUN	<u>2MEQ/ML</u>	<u>A085870 001</u>
<u>AP</u>	BAXTER HLTHCARE	<u>2MEQ/ML</u>	<u>A085499 001</u>
<u>AP</u> +	HOSPIRA	<u>2MEQ/ML</u>	<u>A080205 001</u>
<u>AP</u>	INTL MEDICATION	<u>2MEQ/ML</u>	<u>A083163 001</u>
<u>POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER</u>			
<u>AP</u> +	BAXTER HLTHCARE	<u>14.9MG/ML</u>	<u>N019904 001</u> Dec 26, 1989
<u>AP</u> +		<u>746MG/100ML</u>	<u>N019904 005</u> Dec 17, 1990
<u>AP</u>	HOSPIRA	<u>14.9MG/ML</u>	<u>N020161 005</u> Nov 30, 1992
<u>AP</u>		<u>745MG/100ML</u>	<u>N020161 001</u> Nov 30, 1992
<u>POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER</u>			
<u>AP</u> +	BAXTER HLTHCARE	<u>29.8MG/ML</u>	<u>N019904 002</u> Dec 26, 1989
<u>AP</u> +		<u>1.49GM/100ML</u>	<u>N019904 006</u> Dec 17, 1990
<u>AP</u> +	HOSPIRA	<u>29.8MG/ML</u>	<u>N020161 006</u> Aug 11, 1998
<u>AP</u>		<u>1.49GM/100ML</u>	<u>N020161 002</u> Nov 30, 1992
<u>POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER</u>			
<u>AP</u> +	BAXTER HLTHCARE	<u>2.24GM/100ML</u>	<u>N019904 003</u> Dec 26, 1989
<u>AP</u> +	HOSPIRA	<u>2.24GM/100ML</u>	<u>N020161 003</u> Aug 11, 1998
<u>POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER</u>			
<u>AP</u> +	BAXTER HLTHCARE	<u>2.98GM/100ML</u>	<u>N019904 004</u> Dec 26, 1989
<u>AP</u> +	HOSPIRA	<u>2.98GM/100ML</u>	<u>N020161 004</u> Aug 11, 1998
<u>POTASSIUM CHLORIDE IN PLASTIC CONTAINER</u>			
<u>AP</u>	APP PHARMS	<u>2MEQ/ML</u>	<u>A088901 001</u> Jan 25, 1985
<u>AP</u>		<u>2MEQ/ML</u>	<u>A088908 001</u> Jan 25, 1985
POTASSIUM CHLORIDE			
+ APP PHARMS		3MEQ/ML	A080225 003
TABLET, EXTENDED RELEASE; ORAL			
<u>KLOR-CON M10</u>			
<u>AB</u>	UPSHER SMITH	<u>10MEQ</u>	<u>A074726 002</u> Aug 09, 2000
<u>KLOR-CON M20</u>			
<u>AB</u>	UPSHER SMITH	<u>20MEQ</u>	<u>A074726 001</u> Nov 20, 1998
<u>POTASSIUM CHLORIDE</u>			
<u>AB</u>	EURAND	<u>20MEQ</u>	<u>A076368 001</u> Aug 18, 2004
<u>AB</u>	NESHER PHARMS	<u>20MEQ</u>	<u>A076044 001</u> Apr 05, 2002
<u>AB</u>	SCHERING	<u>10MEQ</u>	<u>N019439 002</u> Jun 13, 1986
<u>AB</u> +		<u>20MEQ</u>	<u>N019439 001</u> Jun 13, 1986
<u>AB</u>	WATSON LABS FLORIDA	<u>10MEQ</u>	<u>A075604 001</u> Apr 10, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 341 (of 424)

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CHLORIDE

<u>AB</u>	WATSON LABS FLORIDA	<u>20MEQ</u>	<u>A075604 002</u>	Apr 10, 2002
	K+10			
BC	FUTURE PAK	10MEQ	A070999 001	Oct 22, 1987
	KAON CL-10			
BC	SAVAGE LABS	10MEQ	N017046 002	
	KLOR-CON			
BC	UPSHER-SMITH LABS	8MEQ	N019123 001	Apr 17, 1986
BC	+ KLOTRIX	10MEQ	N019123 002	Apr 17, 1986
BC	APOTHECON	10MEQ	N017850 001	
	K-TAB			
BC	ABBOTT	10MEQ	N018279 001	
	POTASSIUM CHLORIDE			
BC	ABBOTT	8MEQ	N018279 002	Aug 01, 1988
	KLOR-CON M15			
	UPSHER SMITH	15MEQ	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>	HOSPIRA	<u>149MG/100ML;450MG/100ML</u>	<u>A078446 001</u>	Sep 10, 2008
<u>AP</u>	BAXTER HLTHCARE	<u>150MG/100ML;450MG/100ML</u>	<u>N017648 005</u>	Nov 26, 2002
<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>	
<u>AP</u>	BAXTER HLTHCARE	<u>300MG/100ML;900MG/100ML</u>	<u>N017648 002</u>	
<u>AP</u>	POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
<u>AP</u>	HOSPIRA	<u>149MG/100ML;900MG/100ML</u>	<u>N019686 001</u>	Oct 17, 1988
<u>AP</u>	POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
<u>AP</u>	HOSPIRA	<u>298MG/100ML;900MG/100ML</u>	<u>N019686 002</u>	Oct 17, 1988
	POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9%			
	BAXTER HLTHCARE	<u>224MG/100ML;900MG/100ML</u>	<u>N017648 003</u>	

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CITRATE

<u>AB</u>	COREPHARMA	<u>5MEQ</u>	<u>A077440 001</u>	Jun 09, 2006
<u>AB</u>		<u>10MEQ</u>	<u>A077440 002</u>	Jun 09, 2006
<u>AB</u>	UROCIT-K			
<u>AB</u>	MISSION PHARMA	<u>5MEQ</u>	<u>N019071 001</u>	Aug 30, 1985
<u>AB</u>	+ MISSION PHARMA	<u>10MEQ</u>	<u>N019071 002</u>	Aug 31, 1992
	UROCIT-K			
	+ MISSION PHARMA	15MEQ	N019071 003	Dec 30, 2009

POVIDONE-IODINE

SOLUTION/DROPS; OPHTHALMIC

+ ALCON PHARMS LTD	5%	N018634 001	Dec 17, 1986
--------------------	----	-------------	--------------

PRALATREXATE

SOLUTION; INTRAVENOUS

FOLOTYN

ALLOS	20MG/ML (20MG/ML)	N022468 001	Sep 24, 2009
+ ALLOS	40MG/2ML (20MG/ML)	N022468 002	Sep 24, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 342 (of 424)

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
AB BOEHRINGER INGELHEIM 0.125MG N020667 001 Jul 01, 1997
AB + 0.25MG N020667 002 Jul 01, 1997
AB 0.5MG N020667 006 Feb 12, 1998
AB 0.75MG N020667 007 Jul 30, 2007
AB 1MG N020667 003 Jul 01, 1997
AB 1.5MG N020667 005 Jul 01, 1997
PRAMIPEXOLE DIHYDROCHLORIDE
AB ACTAVIS PHARMA 0.125MG A091254 001 Nov 30, 2010
AB 0.25MG A091254 002 Nov 30, 2010
AB 0.5MG A091254 003 Nov 30, 2010
AB 0.75MG A091254 004 Nov 30, 2010
AB 1MG A091254 005 Nov 30, 2010
AB 1.5MG A091254 006 Nov 30, 2010
AB ALEMBIC LTD 0.125MG A078894 001 Oct 08, 2010
AB 0.25MG A078894 002 Oct 08, 2010
AB 0.5MG A078894 003 Oct 08, 2010
AB 1MG A078894 004 Oct 08, 2010
AB 1.5MG A078894 005 Oct 08, 2010
AB BARR 0.125MG A077724 001 Feb 19, 2008
AB 0.25MG A077724 002 Feb 19, 2008
AB 0.5MG A077724 003 Feb 19, 2008
AB 1MG A077724 004 Feb 19, 2008
AB 1.5MG A077724 005 Feb 19, 2008
AB BRECKENRIDGE PHARM 0.125MG A091450 001 Oct 08, 2010
AB 0.25MG A091450 002 Oct 08, 2010
AB 0.5MG A091450 003 Oct 08, 2010
AB 1MG A091450 004 Oct 08, 2010
AB 1.5MG A091450 005 Oct 08, 2010
AB GLENMARK GENERICS 0.125MG A090781 001 Oct 08, 2010
AB 0.25MG A090781 002 Oct 08, 2010
AB 0.5MG A090781 003 Oct 08, 2010
AB 1MG A090781 004 Oct 08, 2010
AB 1.5MG A090781 005 Oct 08, 2010
AB MYLAN 0.125MG A077854 001 Oct 08, 2010
AB 0.25MG A077854 002 Oct 08, 2010
AB 0.5MG A077854 003 Oct 08, 2010
AB 0.75MG A090764 001 Apr 09, 2010
AB 1MG A077854 004 Oct 08, 2010
AB 1.5MG A077854 005 Oct 08, 2010
AB SANDOZ 0.125MG A090190 001 Jul 06, 2010
AB 0.25MG A090190 002 Jul 06, 2010
AB 0.5MG A090190 003 Jul 06, 2010
AB 0.75MG A090190 006 Oct 08, 2010
AB 1MG A090190 004 Jul 06, 2010
AB 1.5MG A090190 005 Jul 06, 2010
AB TEVA PHARMS 0.125MG A090241 001 Oct 08, 2010
AB 0.25MG A090241 002 Oct 08, 2010
AB 0.5MG A090241 003 Oct 08, 2010
AB 0.75MG A090241 004 Oct 08, 2010
AB 1MG A090241 005 Oct 08, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 343 (of 424)

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	TEVA PHARMS	<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>	WATSON LABS	<u>0.125MG</u>	<u>A078551</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A078551</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078551</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>1MG</u>	<u>A078551</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A078551</u>	<u>005</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

+ BOEHRINGER INGELHEIM	0.375MG	N022421	001	Feb 19, 2010
	0.75MG	N022421	002	Feb 19, 2010
	1.5MG	N022421	003	Feb 19, 2010
	2.25MG	N022421	006	Jun 17, 2011
	3MG	N022421	004	Feb 19, 2010
	3.75MG	N022421	007	Jun 17, 2011
	4.5MG	N022421	005	Feb 19, 2010

PRAMLINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN

AMYLIN	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
+	EQ 3MG BASE/5ML (EQ 600MCG BASE/ML)	N021332	001	Mar 16, 2005

PRASUGREL HYDROCHLORIDE

TABLET; ORAL

EFFIENT

ELI LILLY AND CO	EQ 5MG BASE	N022307	001	Jul 10, 2009
+	EQ 10MG BASE	N022307	002	Jul 10, 2009

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>10MG</u>	<u>N019898</u>	<u>002</u>	Oct 31, 1991
<u>AB</u>		<u>20MG</u>	<u>N019898</u>	<u>003</u>	Oct 31, 1991
<u>AB</u>		<u>40MG</u>	<u>N019898</u>	<u>004</u>	Mar 22, 1993
<u>AB</u>	+	<u>80MG</u>	<u>N019898</u>	<u>008</u>	Dec 18, 2001

PRAVASTATIN SODIUM

<u>AB</u>	APOTEX	<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>	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

PRESCRIPTION DRUG PRODUCT LIST

3 - 344 (of 424)

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVASTATIN SODIUM

AB	GLENMARK GENERICS	<u>20MG</u>	<u>A077987</u> <u>002</u>	May 11, 2007
AB		<u>40MG</u>	<u>A077987</u> <u>003</u>	May 11, 2007
AB		<u>80MG</u>	<u>A077987</u> <u>004</u>	Dec 28, 2007
AB	LEK PHARMS DD	<u>10MG</u>	<u>A076397</u> <u>003</u>	Oct 23, 2006
AB		<u>20MG</u>	<u>A076397</u> <u>002</u>	Oct 23, 2006
AB		<u>40MG</u>	<u>A076397</u> <u>001</u>	Oct 23, 2006
AB		<u>80MG</u>	<u>A077491</u> <u>001</u>	Feb 11, 2008
AB	LUPIN PHARMS	<u>10MG</u>	<u>A077917</u> <u>001</u>	Jan 08, 2008
AB		<u>20MG</u>	<u>A077917</u> <u>002</u>	Jan 08, 2008
AB		<u>40MG</u>	<u>A077917</u> <u>003</u>	Jan 08, 2008
AB		<u>80MG</u>	<u>A077917</u> <u>004</u>	Jan 08, 2008
AB	MATRIX LABS LTD	<u>10MG</u>	<u>A079187</u> <u>001</u>	May 27, 2010
AB		<u>20MG</u>	<u>A079187</u> <u>002</u>	May 27, 2010
AB		<u>40MG</u>	<u>A079187</u> <u>003</u>	May 27, 2010
AB		<u>80MG</u>	<u>A079187</u> <u>004</u>	May 27, 2010
AB	MYLAN	<u>10MG</u>	<u>A077013</u> <u>001</u>	Oct 23, 2006
AB		<u>20MG</u>	<u>A077013</u> <u>002</u>	Oct 23, 2006
AB		<u>40MG</u>	<u>A077013</u> <u>003</u>	Oct 23, 2006
AB		<u>80MG</u>	<u>A077013</u> <u>004</u>	Dec 28, 2007
AB	PLIVA HRVATSKA DOO	<u>10MG</u>	<u>A077730</u> <u>001</u>	Nov 21, 2006
AB		<u>20MG</u>	<u>A077730</u> <u>002</u>	Nov 21, 2006
AB		<u>40MG</u>	<u>A077730</u> <u>005</u>	Nov 21, 2006
AB	RANBAXY	<u>10MG</u>	<u>A076445</u> <u>001</u>	Apr 23, 2007
AB		<u>20MG</u>	<u>A076445</u> <u>002</u>	Apr 23, 2007
AB		<u>40MG</u>	<u>A076445</u> <u>003</u>	Apr 23, 2007
AB		<u>80MG</u>	<u>A076445</u> <u>004</u>	Apr 23, 2007
AB	TEVA	<u>10MG</u>	<u>A076056</u> <u>001</u>	Apr 24, 2006
AB		<u>20MG</u>	<u>A076056</u> <u>002</u>	Apr 24, 2006
AB		<u>40MG</u>	<u>A076056</u> <u>003</u>	Apr 24, 2006
AB	TEVA PHARMS	<u>80MG</u>	<u>A077793</u> <u>001</u>	Jan 15, 2008
AB	WATSON LABS	<u>10MG</u>	<u>A076939</u> <u>004</u>	Oct 23, 2006
AB		<u>20MG</u>	<u>A077904</u> <u>001</u>	Oct 23, 2006
AB		<u>40MG</u>	<u>A076939</u> <u>003</u>	Oct 23, 2006
AB		<u>80MG</u>	<u>A077904</u> <u>002</u>	Oct 23, 2006
AB		<u>40MG</u>	<u>A077904</u> <u>003</u>	Oct 23, 2006
AB		<u>80MG</u>	<u>A076939</u> <u>001</u>	Dec 28, 2007
AB	ZYDUS PHARMS USA	<u>10MG</u>	<u>A077751</u> <u>001</u>	Apr 30, 2008
AB		<u>20MG</u>	<u>A077751</u> <u>002</u>	Apr 30, 2008
AB		<u>40MG</u>	<u>A077751</u> <u>003</u>	Apr 30, 2008
AB		<u>80MG</u>	<u>A077751</u> <u>004</u>	Apr 30, 2008
	PRAVASTATIN SODIUM			
	PLIVA HRVATSKA DOO	<u>30MG</u>	<u>A077730</u> <u>003</u>	Nov 21, 2006

PRAZIQUANTEL

TABLET; ORAL

BILTRICIDE

+ BAYER PHARMA AG 600MG N018714 001 Dec 29, 1982

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIPRESS

AB	PFIZER	<u>EQ 1MG BASE</u>	<u>N017442</u> <u>002</u>
AB	+	<u>EQ 2MG BASE</u>	<u>N017442</u> <u>003</u>
AB		<u>EQ 5MG BASE</u>	<u>N017442</u> <u>001</u>
	PRAZOSIN HYDROCHLORIDE		
AB	IVAX SUB TEVA PHARMS	<u>EQ 1MG BASE</u>	<u>A071994</u> <u>001</u>
			Sep 12, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 345 (of 424)

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 2MG BASE</u>	<u>A071995 001</u>	Sep 12, 1988
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A071745 001</u>	Sep 12, 1988
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>A072575 003</u>	May 16, 1989
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A072575 002</u>	May 16, 1989
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A072575 001</u>	May 16, 1989

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLlient

<u>AB</u> + SANOFI AVENTIS US	<u>0.1%</u>	<u>N020279 001</u>	Oct 29, 1993
<u>AB</u> ALTANA	<u>0.1%</u>	<u>A077287 001</u>	Sep 19, 2006

OINTMENT; TOPICAL

DERMATOP

<u>AB</u> + SANOFI AVENTIS US	<u>0.1%</u>	<u>N019568 001</u>	Sep 23, 1991
<u>AB</u> ALTANA	<u>0.1%</u>	<u>A077236 001</u>	Mar 09, 2007

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

<u>AA</u> ALPHARMA	<u>15MG/5ML</u>	<u>A040323 001</u>	May 13, 1999
<u>AA</u> APOTEX INC	<u>5MG/5ML</u>	<u>A040570 001</u>	Aug 25, 2005
<u>AA</u>	<u>15MG/5ML</u>	<u>A040571 001</u>	Aug 25, 2005
<u>AA</u> HI TECH PHARMA	<u>15MG/5ML</u>	<u>A040401 001</u>	Feb 27, 2003
<u>AA</u> + NESHER PHARMS	<u>5MG/5ML</u>	<u>A040423 001</u>	Oct 22, 2001
<u>AA</u> +	<u>15MG/5ML</u>	<u>A040364 001</u>	Apr 10, 2002
<u>AA</u> PHARM ASSOC	<u>15MG/5ML</u>	<u>A040399 001</u>	Mar 05, 2003
<u>AA</u> VINTAGE	<u>15MG/5ML</u>	<u>A040775 001</u>	Sep 21, 2007
<u>AA</u> WOCKHARDT	<u>15MG/5ML</u>	<u>A040313 001</u>	Sep 10, 2003
<u>AA</u> PRELONE			
<u>AA</u> TEVA	<u>15MG/5ML</u>	<u>A089081 001</u>	Feb 04, 1986

TABLET; ORAL

PREDNISOLONE

+ WATSON LABS

5MG

A080354 001

PREDNISOLONE ACETATE

SUSPENSION; ORAL

FLO-PRED

+ TARO

EQ 15MG BASE/5ML

N022067 002 Jan 17, 2008

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED

<u>AB</u> ALCON	<u>1%</u>	<u>N017469 001</u>
-----------------	-----------	--------------------

PRED FORTE

<u>AB</u> + ALLERGAN	<u>1%</u>	<u>N017011 001</u>
----------------------	-----------	--------------------

PRED MILD

+ ALLERGAN

0.12%

N017100 001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUMOINTMENT; OPHTHALMIC
BLEPHAMIDE S.O.P.

+ ALLERGAN

0.2%;10%

A087748 001 Dec 03, 1986

SUSPENSION; OPHTHALMIC
BLEPHAMIDE

+ ALLERGAN

0.2%;10%

N012813 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 346 (of 424)

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

ORAPRED

<u>AA</u>	+ SHIONOGI INC	<u>EQ 15MG BASE/5ML</u>	<u>A075117 001</u>	Dec 14, 2000
		<u>PEDIAFRED</u>		
<u>AA</u>	+ UCB INC	<u>EQ 5MG BASE/5ML</u>	<u>N019157 001</u>	May 28, 1986
		<u>PREDNISOLONE SODIUM PHOSPHATE</u>		
<u>AA</u>	AMNEAL PHARMS	<u>EQ 15MG BASE/5ML</u>	<u>A078345 001</u>	Mar 10, 2009
<u>AA</u>	HI TECH PHARMA	<u>EQ 5MG BASE/5ML</u>	<u>A075183 001</u>	Mar 26, 2003
<u>AA</u>	NESHER PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A076982 001</u>	May 24, 2005
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>A076988 001</u>	May 24, 2005
<u>AA</u>	PADDOCK LLC	<u>EQ 5MG BASE/5ML</u>	<u>A075988 001</u>	May 25, 2004
<u>AA</u>	PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>A076123 001</u>	Dec 23, 2002
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>A076913 001</u>	Apr 25, 2005
<u>AA</u>	VINTAGE	<u>EQ 15MG BASE/5ML</u>	<u>A079010 001</u>	May 26, 2009
<u>AA</u>	WE PHARMS	<u>EQ 15MG BASE/5ML</u>	<u>A075250 001</u>	Jul 12, 2002
<u>AA</u>	WOCKHARDT	<u>EQ 5MG BASE/5ML</u>	<u>A075099 001</u>	Jun 28, 2002
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>A076895 001</u>	Oct 04, 2004
		<u>PREDNISOLONE SODIUM PHOSPHATE</u>		
+ PHARM ASSOC		<u>EQ 10MG BASE/5ML</u>	<u>A078465 001</u>	Mar 07, 2008
+		<u>EQ 20MG BASE/5ML</u>	<u>A078988 001</u>	Jun 09, 2008
+		<u>EQ 25MG BASE/5ML</u>	<u>A091396 001</u>	Sep 13, 2010

SOLUTION/DROPS; OPHTHALMIC

+ BAUSCH AND LOMB	<u>EQ 0.9% PHOSPHATE</u>	
		<u>A040070 001</u>

Jul 29, 1994

TABLET, ORALLY DISINTEGRATING; ORAL

ORAPRED ODT

SHIONOGI INC	<u>EQ 10MG BASE</u>	<u>N021959 001</u>	Jun 01, 2006
	<u>EQ 15MG BASE</u>	<u>N021959 002</u>	Jun 01, 2006
+	<u>EQ 30MG BASE</u>	<u>N021959 003</u>	Jun 01, 2006

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

<u>AT</u>	ALCON PHARMS LTD	<u>EQ 0.23% PHOSPHATE;10%</u>	<u>A073630 001</u>	May 27, 1993
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.23% PHOSPHATE;10%</u>	<u>A074449 001</u>	Dec 29, 1995
		<u>VASOCIDIN</u>		
<u>AT</u>	+ NOVARTIS	<u>EQ 0.23% PHOSPHATE;10%</u>	<u>N018988 001</u>	Aug 26, 1988

PREDNISONE

SOLUTION; ORAL

PREDNISONE

+ ROXANE	<u>5MG/5ML</u>	<u>A088703 001</u>	Nov 08, 1984
PREDNISONE INTENSOL			
+ ROXANE	<u>5MG/ML</u>	<u>A088810 001</u>	Feb 20, 1985

TABLET; ORAL

PREDNISONE

<u>AB</u>	CONTRACT PHARMACAL	<u>5MG</u>	<u>A080209 001</u>	
<u>AB</u>	JUBILANT CADISTA	<u>1MG</u>	<u>A040611 001</u>	Jun 06, 2005
<u>AB</u>		<u>5MG</u>	<u>A040362 002</u>	Aug 29, 2001
<u>AB</u>		<u>10MG</u>	<u>A040362 001</u>	Aug 29, 2001
<u>AB</u>		<u>20MG</u>	<u>A040362 003</u>	Jun 29, 2005
<u>AB</u>	MUTUAL PHARM	<u>5MG</u>	<u>A089245 001</u>	Dec 04, 1985
<u>AB</u>		<u>10MG</u>	<u>A089246 001</u>	Dec 04, 1985
<u>AB</u>		<u>20MG</u>	<u>A089247 001</u>	Dec 04, 1985
<u>AB</u>	+ ROXANE	<u>1MG</u>	<u>A087800 001</u>	Apr 22, 1982
<u>AB</u>	+	<u>2.5MG</u>	<u>A087801 001</u>	Apr 22, 1982
<u>AB</u>	+	<u>5MG</u>	<u>A080352 001</u>	
<u>AB</u>	+	<u>10MG</u>	<u>A084122 001</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 347 (of 424)

PREDNISONE

TABLET; ORAL

PREDNISONE

<u>AB</u>	+ ROXANE	<u>20MG</u>	<u>A087342</u> <u>001</u>
<u>AB</u>	+	<u>50MG</u>	<u>A084283</u> <u>001</u>
<u>AB</u>	VINTAGE PHARMS	<u>1MG</u>	<u>A040584</u> <u>001</u>
<u>AB</u>		<u>2.5MG</u>	<u>A040581</u> <u>001</u>
<u>AB</u>		<u>5MG</u>	<u>A040256</u> <u>001</u>
<u>AB</u>		<u>10MG</u>	<u>A040256</u> <u>002</u>
<u>AB</u>		<u>20MG</u>	<u>A040392</u> <u>001</u>
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>A080356</u> <u>001</u>
<u>AB</u>		<u>10MG</u>	<u>A085162</u> <u>001</u>
<u>AB</u>		<u>20MG</u>	<u>A085161</u> <u>001</u>
<u>AB</u>	WEST WARD	<u>2.5MG</u>	<u>A040538</u> <u>001</u>
<u>AB</u>		<u>5MG</u>	<u>A080292</u> <u>001</u>
<u>AB</u>		<u>10MG</u>	<u>A088832</u> <u>001</u>
<u>AB</u>		<u>20MG</u>	<u>A083677</u> <u>001</u>
<u>AB</u>	WEST WARD PHARM CORP	<u>1MG</u>	<u>A040890</u> <u>001</u>

PREGABALIN

CAPSULE; ORAL

LYRICA

CPPI CV	25MG	N021446 001	Dec 30, 2004
	50MG	N021446 002	Dec 30, 2004
	75MG	N021446 003	Dec 30, 2004
	100MG	N021446 004	Dec 30, 2004
	150MG	N021446 005	Dec 30, 2004
	200MG	N021446 006	Dec 30, 2004
	225MG	N021446 007	Dec 30, 2004
+	300MG	N021446 008	Dec 30, 2004

SOLUTION; ORAL

LYRICA

+	CPPI CV	20MG/ML	N022488 001	Jan 04, 2010
---	---------	---------	-------------	--------------

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST PLAIN DENTAL

<u>AP</u>	+ DENTSPLY PHARM	<u>4%</u>	<u>N021382</u> <u>001</u>
<u>AP</u>	PRILOCAINE HYDROCHLORIDE		
<u>AP</u>	SEPTODONT INC	<u>4%</u>	<u>A079235</u> <u>001</u>

Sep 29, 2010

PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE

+	SANOFI AVENTIS US	EQ 15MG BASE	N008316 001
---	-------------------	--------------	-------------

PRIMIDONE

TABLET; ORAL

MYSOLINE

<u>AB</u>	+ VALEANT	<u>50MG</u>	<u>N009170</u> <u>003</u>
<u>AB</u>		<u>250MG</u>	<u>N009170</u> <u>002</u>
<u>AB</u>	AMNEAL PHARM	<u>50MG</u>	<u>A040866</u> <u>001</u>
<u>AB</u>		<u>250MG</u>	<u>A040866</u> <u>002</u>
<u>AB</u>	DR REDDYS LABS LTD	<u>50MG</u>	<u>A040862</u> <u>001</u>
<u>AB</u>		<u>250MG</u>	<u>A040862</u> <u>002</u>
<u>AB</u>	IMPAK LABS	<u>50MG</u>	<u>A040717</u> <u>001</u>
<u>AB</u>		<u>250MG</u>	<u>A040717</u> <u>002</u>
<u>AB</u>	LANNETT	<u>50MG</u>	<u>A084903</u> <u>002</u>

Apr 23, 2008

Apr 23, 2008

Oct 03, 2008

Oct 03, 2008

Feb 12, 2008

Feb 12, 2008

May 24, 2001

PRESCRIPTION DRUG PRODUCT LIST

3 - 348 (of 424)

PRIMIDONE

TABLET; ORAL

PRIMIDONE

<u>AB</u>	LANNETT	<u>250MG</u>	<u>A084903</u> <u>001</u>	
<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>A040626</u> <u>001</u>	Sep 29, 2005
<u>AB</u>		<u>250MG</u>	<u>A040626</u> <u>002</u>	Sep 29, 2005
<u>AB</u>	VINTAGE PHARMS	<u>50MG</u>	<u>A040586</u> <u>001</u>	Feb 24, 2005
<u>AB</u>		<u>250MG</u>	<u>A040586</u> <u>002</u>	Feb 24, 2005
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A083551</u> <u>001</u>	
<u>AB</u>	WEST WARD	<u>50MG</u>	<u>A040667</u> <u>001</u>	Jul 27, 2006
<u>AB</u>		<u>250MG</u>	<u>A040667</u> <u>002</u>	Jul 27, 2006

PROBENECID

TABLET; ORAL

PROBALAN

<u>AB</u>	LANNETT	<u>500MG</u>	<u>A080966</u> <u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>A083740</u> <u>001</u>	May 09, 1984
<u>AB</u>	+ MYLAN	<u>500MG</u>	<u>A084211</u> <u>002</u>	
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A084442</u> <u>004</u>	Mar 29, 1983

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

<u>AP</u>	+ HOSPIRA	<u>100MG/ML</u>	<u>A089069</u> <u>001</u>	Feb 12, 1986
<u>AP</u>	+	<u>500MG/ML</u>	<u>A089070</u> <u>001</u>	Feb 12, 1986
<u>AP</u>	INTL MEDICATION	<u>100MG/ML</u>	<u>A088636</u> <u>001</u>	Jul 31, 1984
<u>AP</u>		<u>500MG/ML</u>	<u>A088637</u> <u>001</u>	Jul 31, 1984

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

+ SIGMA TAU EQ 50MG BASE

N016785 001

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO

<u>AB</u>	PADDOCK LLC	<u>25MG</u>	<u>A040246</u> <u>001</u>	Jun 28, 2000
<u>AB</u>	+ G AND W LABS	<u>25MG</u>	<u>A040058</u> <u>001</u>	Nov 24, 1993

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

<u>AP</u>	+ BAXTER HLTHCARE	<u>EQ 5MG BASE/ML</u>	<u>A089903</u> <u>001</u>	Aug 29, 1989
<u>AP</u>	BEDFORD	<u>EQ 5MG BASE/ML</u>	<u>A040540</u> <u>001</u>	May 28, 2004

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

<u>AB</u>	MYLAN	<u>EQ 5MG BASE</u>	<u>A040185</u> <u>002</u>	Oct 28, 1996
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A040185</u> <u>001</u>	Oct 28, 1996
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>A040101</u> <u>001</u>	Jul 19, 1996
<u>AB</u>	+	<u>EQ 10MG BASE</u>	<u>A040101</u> <u>002</u>	Jul 19, 1996
<u>AB</u>	TEVA PHARMS	<u>EQ 5MG BASE</u>	<u>A040120</u> <u>001</u>	Jul 11, 1996
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A040120</u> <u>002</u>	Jul 11, 1996
	<u>PROCOMP</u>			
<u>AB</u>	JUBILANT CADISTA	<u>EQ 5MG BASE</u>	<u>A040268</u> <u>001</u>	Feb 27, 1998
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A040268</u> <u>002</u>	Feb 27, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 349 (of 424)

PROGESTERONE

CAPSULE; ORAL PROMETRIUM ABBOTT LABS	100MG 200MG	N019781 001 N019781 002	May 14, 1998 Oct 15, 1999
GEL; VAGINAL CRINONE WATSON LABS	4% 8%	N020701 001 N020701 002	Jul 31, 1997 Jul 31, 1997
INJECTABLE; INJECTION <u>PROGESTERONE</u>			
AO APP PHARMS	<u>50MG/ML</u>	A075906 001	Apr 25, 2001
AO HIKMA FARMACEUTICA	<u>50MG/ML</u>	A091033 001	Oct 28, 2010
AO LUITPOLD	<u>50MG/ML</u>	A090845 001	Jun 22, 2009
AO + WATSON LABS (UTAH)	<u>50MG/ML</u>	<u>N017362 002</u>	
INSERT; VAGINAL ENDOMETRIN + FERRING	100MG	N022057 001	Jun 21, 2007

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION <u>PROMETHAZINE HYDROCHLORIDE</u>			
AP BAXTER HLTHCARE	<u>25MG/ML</u>	A083312 001	
AP	<u>50MG/ML</u>	A083312 002	
AP HOSPIRA	<u>25MG/ML</u>	A040372 001	Jun 08, 2000
AP LUITPOLD	<u>25MG/ML</u>	A040515 001	Mar 19, 2003
AP + TEVA PARENTERAL	<u>25MG/ML</u>	A040454 001	Aug 22, 2002
AP +	<u>50MG/ML</u>	A040454 002	Aug 22, 2002
AP WOCKHARDT	<u>25MG/ML</u>	A040785 001	Sep 26, 2008
AP	<u>50MG/ML</u>	A040785 002	Sep 26, 2008
AP X-GEN PHARMS	<u>25MG/ML</u>	A040737 001	Apr 24, 2008
AP	<u>50MG/ML</u>	A040737 002	Apr 24, 2008
SUPPOSITORY; RECTAL <u>PROMETHAZINE HYDROCHLORIDE</u>			
AB G AND W LABS	<u>12.5MG</u>	A040428 002	Mar 31, 2003
AB +	<u>25MG</u>	A040428 001	Feb 05, 2002
AB PADDOCK LLC	<u>12.5MG</u>	A040479 001	Jun 24, 2003
AB	<u>25MG</u>	A040479 002	Jun 24, 2003
AB PERRIGO NEW YORK	<u>12.5MG</u>	A040500 001	Jun 30, 2003
AB	<u>25MG</u>	A040500 002	Jun 30, 2003
AB TARO	<u>12.5MG</u>	A040603 001	Oct 26, 2006
AB	<u>25MG</u>	A040603 002	Oct 26, 2006
PROMETHEGAN + G AND W LABS	50MG	A087165 001	Aug 14, 1987
SYRUP; ORAL <u>PROMETHAZINE HYDROCHLORIDE</u>			
AA AMNEAL PHARMS	<u>6.25MG/5ML</u>	A040882 001	Dec 30, 2009
AA HI TECH PHARMA	<u>6.25MG/5ML</u>	A040026 001	Sep 25, 1998
AA SUN PHARM INDs INC	<u>6.25MG/5ML</u>	A040891 001	Mar 13, 2009
AA TARO	<u>6.25MG/5ML</u>	A040718 001	Apr 04, 2007
AA VINTAGE	<u>6.25MG/5ML</u>	A040643 001	Apr 26, 2006
PROMETHAZINE PLAIN			
AA + WOCKHARDT	<u>6.25MG/5ML</u>	A087953 001	Nov 15, 1982
TABLET; ORAL <u>PROMETHAZINE HYDROCHLORIDE</u>			
AB AMNEAL PHARMS NY	<u>12.5MG</u>	A091179 001	Dec 13, 2010
AB	<u>25MG</u>	A091179 002	Dec 13, 2010
AB	<u>50MG</u>	A091179 003	Dec 13, 2010
AB EMCURE PHARMS USA	<u>12.5MG</u>	A040673 001	Mar 05, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 350 (of 424)

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	EMCURE PHARMS USA	<u>25MG</u>	<u>A040673</u> <u>002</u>	Mar 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A040673</u> <u>003</u>	Mar 05, 2008
<u>AB</u>	IMPAK LABS	<u>12.5MG</u>	<u>A040724</u> <u>001</u>	Feb 12, 2008
<u>AB</u>		<u>25MG</u>	<u>A040724</u> <u>002</u>	Feb 12, 2008
<u>AB</u>		<u>50MG</u>	<u>A040791</u> <u>001</u>	May 20, 2008
<u>AB</u>	KVK TECH	<u>12.5MG</u>	<u>A040712</u> <u>002</u>	May 04, 2007
<u>AB</u>		<u>25MG</u>	<u>A040712</u> <u>001</u>	Jul 31, 2006
<u>AB</u>		<u>50MG</u>	<u>A040713</u> <u>001</u>	Jul 31, 2006
<u>AB</u>	MYLAN	<u>12.5MG</u>	<u>A091054</u> <u>001</u>	Aug 30, 2011
<u>AB</u>		<u>25MG</u>	<u>A091054</u> <u>002</u>	Aug 30, 2011
<u>AB</u>		<u>50MG</u>	<u>A091054</u> <u>003</u>	Aug 30, 2011
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A084234</u> <u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A084176</u> <u>001</u>	
<u>AB</u>	+ SUN PHARM IND'S INC	<u>12.5MG</u>	<u>A040863</u> <u>001</u>	Dec 30, 2008
<u>AB</u>		<u>25MG</u>	<u>A040863</u> <u>002</u>	Dec 30, 2008
<u>AB</u>		<u>50MG</u>	<u>A040863</u> <u>003</u>	Dec 30, 2008
<u>AB</u>	VINTAGE PHARMS	<u>12.5MG</u>	<u>A040622</u> <u>001</u>	Jul 18, 2006
<u>AB</u>		<u>25MG</u>	<u>A040622</u> <u>002</u>	Jul 18, 2006
<u>AB</u>		<u>50MG</u>	<u>A040622</u> <u>003</u>	Jul 18, 2006
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A083426</u> <u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>A083711</u> <u>001</u>	
<u>AB</u>	ZYDUS PHARMS USA	<u>12.5MG</u>	<u>A040596</u> <u>001</u>	Nov 18, 2005
<u>AB</u>		<u>25MG</u>	<u>A040596</u> <u>002</u>	Nov 18, 2005
<u>AB</u>		<u>50MG</u>	<u>A040596</u> <u>003</u>	Nov 18, 2005

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	PAR PHARM	<u>225MG</u>	<u>A078540</u> <u>001</u>	Oct 18, 2010
<u>AB</u>		<u>325MG</u>	<u>A078540</u> <u>002</u>	Oct 18, 2010
<u>AB</u>		<u>425MG</u>	<u>A078540</u> <u>003</u>	Oct 18, 2010
<u>AB</u>	<u>RYTHMOL SR</u>			
<u>AB</u>	GLAXOSMITHKLINE LLC	<u>225MG</u>	<u>N021416</u> <u>001</u>	Sep 04, 2003
<u>AB</u>		<u>325MG</u>	<u>N021416</u> <u>002</u>	Sep 04, 2003
<u>AB</u>	+ <u>425MG</u>		<u>N021416</u> <u>003</u>	Sep 04, 2003

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	MUTUAL PHARM	<u>150MG</u>	<u>A075998</u> <u>001</u>	Nov 29, 2001
<u>AB</u>		<u>225MG</u>	<u>A075998</u> <u>002</u>	Nov 29, 2001
<u>AB</u>		<u>300MG</u>	<u>A075998</u> <u>003</u>	Nov 29, 2001
<u>AB</u>	NESHER PHARMS	<u>150MG</u>	<u>A076193</u> <u>001</u>	Feb 07, 2002
<u>AB</u>		<u>225MG</u>	<u>A076193</u> <u>002</u>	Feb 07, 2002
<u>AB</u>		<u>300MG</u>	<u>A076193</u> <u>003</u>	Feb 07, 2002
<u>AB</u>	PLIVA	<u>150MG</u>	<u>A076550</u> <u>001</u>	Apr 23, 2004
<u>AB</u>		<u>225MG</u>	<u>A076550</u> <u>002</u>	Apr 23, 2004
<u>AB</u>		<u>300MG</u>	<u>A076550</u> <u>003</u>	Apr 23, 2004
<u>AB</u>	VINTAGE PHARMS	<u>150MG</u>	<u>A075938</u> <u>001</u>	Oct 17, 2002
<u>AB</u>		<u>225MG</u>	<u>A075938</u> <u>002</u>	Oct 17, 2002
<u>AB</u>		<u>300MG</u>	<u>A075938</u> <u>003</u>	Oct 17, 2002
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075203</u> <u>001</u>	Oct 24, 2000
<u>AB</u>		<u>225MG</u>	<u>A075203</u> <u>002</u>	Oct 24, 2000
<u>AB</u>	<u>RYTHMOL</u>			
<u>AB</u>	GLAXOSMITHKLINE LLC	<u>150MG</u>	<u>N019151</u> <u>001</u>	Nov 27, 1989
<u>AB</u>		<u>225MG</u>	<u>N019151</u> <u>003</u>	Nov 20, 1992
<u>AB</u>	+ <u>300MG</u>		<u>N019151</u> <u>002</u>	Nov 27, 1989

PRESCRIPTION DRUG PRODUCT LIST

3 - 351 (of 424)

PROPANTHELINE BROMIDE

TABLET; ORAL
 PROPANTHELINE BROMIDE
 + ROXANE 15MG

A080927 002

PROPARACAINe HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

<u>ATC</u>	<u>ALCAINE</u>	<u>0.5%</u>	<u>A080027</u> <u>001</u>
<u>ATC</u>	<u>OPHTHAINE</u>	<u>0.5%</u>	<u>N008883</u> <u>001</u>
<u>ATC</u>	+ APOTHECON	<u>0.5%</u>	<u>N012583</u> <u>001</u>
<u>ATC</u>	<u>OPHTHETIC</u>		
<u>ATC</u>	+ ALLERGAN	<u>0.5%</u>	<u>A040074</u> <u>001</u> Sep 29, 1995
<u>ATC</u>	<u>PROPARACAINe HYDROCHLORIDE</u>		<u>A040277</u> <u>001</u> Mar 16, 2000
<u>ATC</u>	BAUSCH AND LOMB	<u>0.5%</u>	
<u>ATC</u>	TAYLOR PHARMA	<u>0.5%</u>	

PROPOFOL

INJECTABLE; INJECTION

<u>AB</u>	<u>DIPRIVAN</u>	<u>10MG/ML</u>	<u>N019627</u> <u>002</u>	Jun 11, 1996
<u>AB</u>	<u>PROPOFOL</u>		<u>A077908</u> <u>001</u>	Mar 17, 2006
<u>AB</u>	HOSPIRA	<u>10MG/ML</u>	<u>A075102</u> <u>001</u>	Jan 04, 1999
<u>AB</u>	TEVA PARENTERAL	<u>10MG/ML</u>		

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

<u>AB</u>	<u>INDERAL LA</u>			
<u>AB</u>	AKRIMAX PHARMS	<u>60MG</u>	<u>N018553</u> <u>004</u>	Mar 18, 1987
<u>AB</u>		<u>80MG</u>	<u>N018553</u> <u>002</u>	Apr 19, 1983
<u>AB</u>		<u>120MG</u>	<u>N018553</u> <u>003</u>	Apr 19, 1983
<u>AB</u>	+	<u>160MG</u>	<u>N018553</u> <u>001</u>	Apr 19, 1983
<u>AB</u>	<u>PROPRANOLOL HYDROCHLORIDE</u>			
<u>AB</u>	ACTAVIS ELIZABETH	<u>60MG</u>	<u>A078494</u> <u>001</u>	Aug 10, 2007
<u>AB</u>		<u>80MG</u>	<u>A078494</u> <u>002</u>	Aug 10, 2007
<u>AB</u>		<u>120MG</u>	<u>A078494</u> <u>003</u>	Aug 10, 2007
<u>AB</u>		<u>160MG</u>	<u>A078494</u> <u>004</u>	Aug 10, 2007
<u>AB</u>	GLAXOSMITHKLINE	<u>60MG</u>	<u>A078703</u> <u>001</u>	Jul 15, 2011
<u>AB</u>		<u>80MG</u>	<u>A078703</u> <u>002</u>	Jul 15, 2011
<u>AB</u>		<u>120MG</u>	<u>A078703</u> <u>003</u>	Jul 15, 2011
<u>AB</u>		<u>160MG</u>	<u>A078703</u> <u>004</u>	Jul 15, 2011
<u>AB</u>	MYLAN	<u>60MG</u>	<u>A078022</u> <u>001</u>	Feb 15, 2007
<u>AB</u>		<u>80MG</u>	<u>A078022</u> <u>002</u>	Feb 15, 2007
<u>AB</u>		<u>120MG</u>	<u>A078022</u> <u>003</u>	Feb 15, 2007
<u>AB</u>		<u>160MG</u>	<u>A078022</u> <u>004</u>	Feb 15, 2007
<u>AB</u>	PAR PHARM	<u>60MG</u>	<u>A078065</u> <u>001</u>	Jan 26, 2007
<u>AB</u>		<u>80MG</u>	<u>A078065</u> <u>002</u>	Jan 26, 2007
<u>AB</u>		<u>120MG</u>	<u>A078065</u> <u>003</u>	Jan 26, 2007
<u>AB</u>		<u>160MG</u>	<u>A078065</u> <u>004</u>	Jan 26, 2007
<u>AB</u>	UPSHER SMITH	<u>60MG</u>	<u>A078311</u> <u>001</u>	Mar 06, 2009
<u>AB</u>		<u>80MG</u>	<u>A078311</u> <u>002</u>	Mar 06, 2009
<u>AB</u>		<u>120MG</u>	<u>A078311</u> <u>003</u>	Mar 06, 2009
<u>AB</u>		<u>160MG</u>	<u>A078311</u> <u>004</u>	Mar 06, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>60MG</u>	<u>A090321</u> <u>001</u>	Mar 25, 2011
<u>AB</u>		<u>80MG</u>	<u>A090321</u> <u>002</u>	Mar 25, 2011
<u>AB</u>		<u>120MG</u>	<u>A090321</u> <u>003</u>	Mar 25, 2011
<u>AB</u>		<u>160MG</u>	<u>A090321</u> <u>004</u>	Mar 25, 2011
<u>BX</u>	INNOPRAN XL			
<u>BX</u>	GLAXOSMITHKLINE LLC	<u>80MG</u>	<u>N021438</u> <u>001</u>	Mar 12, 2003
<u>BX</u>		<u>120MG</u>	<u>N021438</u> <u>002</u>	Mar 12, 2003

PREScription DRUG PRODUCT LIST

3 - 352 (of 424)

PROPRANOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

<u>AP</u>	APP PHARMS	<u>1MG/ML</u>	<u>A075826</u>	<u>001</u>	Aug 31, 2001
<u>AP</u> +	BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>N016419</u>	<u>001</u>	
<u>AP</u>	BEDFORD	<u>1MG/ML</u>	<u>A075792</u>	<u>001</u>	Aug 29, 2000
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077760</u>	<u>001</u>	Jan 31, 2008
<u>AP</u>	SANDOZ	<u>1MG/ML</u>	<u>A076400</u>	<u>001</u>	Feb 26, 2003

SOLUTION; ORAL

PROPRANOLOL HYDROCHLORIDE

+ ROXANE 20MG/5ML A070979 001 May 15, 1987
+ 40MG/5ML A070690 001 May 15, 1987

TABLET; ORAL

INDERAL

AB AKRIMAX PHARMS **40MG** **N016418 002**
AB **60MG** **N016418 009** Oct 18, 1982
AB + **80MG** **N016418 004**

PROPRANOLOL HYDROCHLORIDE

<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>	PLIVA	<u>10MG</u>	<u>A071972</u>	<u>001</u>	Apr 06, 1988
<u>AB</u>		<u>20MG</u>	<u>A071973</u>	<u>001</u>	Apr 06, 1988
<u>AB</u>		<u>40MG</u>	<u>A071974</u>	<u>001</u>	Apr 06, 1988
<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>	VINTAGE PHARMS	<u>10MG</u>	<u>A070217</u>	<u>001</u>	Aug 01, 1986
<u>AB</u>		<u>20MG</u>	<u>A070218</u>	<u>001</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>A070220</u>	<u>001</u>	Sep 24, 1986
<u>AB</u>		<u>80MG</u>	<u>A070221</u>	<u>001</u>	Apr 14, 1986
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A070175</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>20MG</u>	<u>A070176</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>40MG</u>	<u>A070177</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>80MG</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
BD WEST WARD 50MG A080154 001

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

+ APP PHARMS 10MG/ML A089454 001 Apr 07, 1987

PRESCRIPTION DRUG PRODUCT LIST

3 - 353 (of 424)

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	ROXANE	<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>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>VIVACTIL</u>				
<u>AB</u>	ODYSSEY PHARMS	<u>5MG</u>	<u>A073644</u>	<u>001</u>	Aug 24, 1995
<u>AB</u>	<u>+</u>	<u>10MG</u>	<u>A073645</u>	<u>001</u>	Aug 24, 1995

PYRAZINAMIDE

TABLET; ORAL

PYRAZINAMIDE

<u>AB</u>	<u>+</u>	DAVA PHARMS INC	<u>500MG</u>	<u>A080157</u>	<u>001</u>	
<u>AB</u>		MIKART	<u>500MG</u>	<u>A081319</u>	<u>001</u>	Jun 30, 1992

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION

MESTINON

<u>AP</u>	<u>+</u>	VALEANT PHARM INTL	<u>5MG/ML</u>	<u>N009830</u>	<u>001</u>	
<u>AP</u>		REGONOL				

<u>AP</u>	SANDOZ	<u>5MG/ML</u>	<u>N017398</u>	<u>001</u>	
-----------	--------	---------------	----------------	------------	--

SYRUP; ORAL

MESTINON

+ VALEANT PHARM INTL	60MG/ 5ML
----------------------	-----------

N015193 001

TABLET; ORAL

MESTINON

<u>AB</u>	<u>+</u>	VALEANT PHARM INTL	<u>60MG</u>	<u>N009829</u>	<u>002</u>	
-----------	----------	--------------------	-------------	----------------	------------	--

PYRIDOSTIGMINE BROMIDE

<u>AB</u>	COREPHARMA	<u>60MG</u>	<u>A040457</u>	<u>001</u>	Dec 26, 2002
<u>AB</u>	IMPAX LABS	<u>60MG</u>	<u>A040502</u>	<u>001</u>	Apr 24, 2003

TABLET, EXTENDED RELEASE; ORAL

MESTINON

+ VALEANT PHARM INTL	180MG
----------------------	-------

N011665 001

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

PYRIDOXINE HYDROCHLORIDE

+ APP PHARMS	100MG/ML
--------------	----------

A080618 001

PYRIMETHAMINE

TABLET; ORAL

DARAPRIM

+ COREPHARMA	25MG
--------------	------

N008578 001

QUAZEPAM

TABLET; ORAL

DORAL

+ QUESTCOR PHARMS	15MG
-------------------	------

N018708 001 Dec 27, 1985

QUETIAPINE FUMARATE

TABLET; ORAL

SEROQUEL

+ ASTRazeneca	EQ 25MG BASE
	EQ 50MG BASE
	EQ 100MG BASE
	EQ 200MG BASE
+	EQ 300MG BASE

N020639 001	Sep 26, 1997
N020639 007	Oct 04, 2005
N020639 002	Sep 26, 1997
N020639 003	Sep 26, 1997
N020639 005	Jul 26, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 354 (of 424)

QUETIAPINE FUMARATE

TABLET; ORAL SEROQUEL ASTRAZENECA	EQ 400MG BASE	N020639 006	Oct 04, 2005
TABLET, EXTENDED RELEASE; ORAL SEROQUEL XR ASTRAZENECA	EQ 50MG BASE EQ 150MG BASE EQ 200MG BASE EQ 300MG BASE EQ 400MG BASE	N022047 001 N022047 005 N022047 002 N022047 003 N022047 004	May 17, 2007 Aug 11, 2008 May 17, 2007 May 17, 2007 May 17, 2007
+ +			

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL <u>ACCUPRIL</u> PFIZER PHARMS	<u>EQ 5MG BASE</u>	<u>N019885 001</u>	Nov 19, 1991
	<u>EQ 10MG BASE</u>	<u>N019885 002</u>	Nov 19, 1991
	<u>EQ 20MG BASE</u>	<u>N019885 003</u>	Nov 19, 1991
+ +	<u>EQ 40MG BASE</u>	<u>N019885 004</u>	Nov 19, 1991
<u>QUINAPRIL HYDROCHLORIDE</u>			
APOTEX	<u>EQ 5MG BASE</u>	<u>A076240 001</u>	Jan 26, 2006
	<u>EQ 10MG BASE</u>	<u>A076240 002</u>	Jan 26, 2006
	<u>EQ 20MG BASE</u>	<u>A076240 003</u>	Jan 26, 2006
	<u>EQ 40MG BASE</u>	<u>A076240 004</u>	Jan 26, 2006
INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078457 001</u>	Aug 24, 2007
	<u>EQ 10MG BASE</u>	<u>A078457 002</u>	Aug 24, 2007
	<u>EQ 20MG BASE</u>	<u>A078457 003</u>	Aug 24, 2007
	<u>EQ 40MG BASE</u>	<u>A078457 004</u>	Aug 24, 2007
LUPIN	<u>EQ 5MG BASE</u>	<u>A077690 001</u>	Jun 20, 2006
	<u>EQ 10MG BASE</u>	<u>A077690 002</u>	Jun 20, 2006
	<u>EQ 20MG BASE</u>	<u>A077690 003</u>	Jun 20, 2006
	<u>EQ 40MG BASE</u>	<u>A077690 004</u>	Jun 20, 2006
MYLAN	<u>EQ 5MG BASE</u>	<u>A076036 001</u>	Jan 28, 2005
	<u>EQ 5MG BASE</u>	<u>A076694 001</u>	Dec 23, 2004
	<u>EQ 10MG BASE</u>	<u>A076036 002</u>	Jan 28, 2005
	<u>EQ 10MG BASE</u>	<u>A076694 002</u>	Dec 23, 2004
	<u>EQ 20MG BASE</u>	<u>A076036 003</u>	Jan 28, 2005
	<u>EQ 20MG BASE</u>	<u>A076694 003</u>	Dec 23, 2004
	<u>EQ 40MG BASE</u>	<u>A076036 004</u>	Jan 28, 2005
	<u>EQ 40MG BASE</u>	<u>A076694 004</u>	Dec 23, 2004
RANBAXY	<u>EQ 5MG BASE</u>	<u>A076607 001</u>	Dec 15, 2004
	<u>EQ 10MG BASE</u>	<u>A076607 002</u>	Dec 15, 2004
	<u>EQ 20MG BASE</u>	<u>A076607 003</u>	Dec 15, 2004
	<u>EQ 40MG BASE</u>	<u>A076607 004</u>	Dec 15, 2004
SANDOZ	<u>EQ 5MG BASE</u>	<u>A076803 001</u>	Mar 02, 2005
	<u>EQ 10MG BASE</u>	<u>A076803 002</u>	Mar 02, 2005
	<u>EQ 20MG BASE</u>	<u>A076803 003</u>	Mar 02, 2005
	<u>EQ 40MG BASE</u>	<u>A076803 004</u>	Mar 02, 2005
SUN PHARM IND LTD	<u>EQ 5MG BASE</u>	<u>A090800 001</u>	Jun 18, 2009
	<u>EQ 10MG BASE</u>	<u>A090800 002</u>	Jun 18, 2009
	<u>EQ 20MG BASE</u>	<u>A090800 003</u>	Jun 18, 2009
	<u>EQ 40MG BASE</u>	<u>A090800 004</u>	Jun 18, 2009
TEVA	<u>EQ 5MG BASE</u>	<u>A075504 001</u>	Aug 24, 2007
	<u>EQ 10MG BASE</u>	<u>A075504 002</u>	Aug 24, 2007
	<u>EQ 20MG BASE</u>	<u>A075504 003</u>	Aug 24, 2007
	<u>EQ 40MG BASE</u>	<u>A075504 004</u>	Aug 24, 2007
WATSON LABS FLORIDA	<u>EQ 5MG BASE</u>	<u>A076049 001</u>	Jan 14, 2005
	<u>EQ 10MG BASE</u>	<u>A076049 002</u>	Jan 14, 2005
	<u>EQ 20MG BASE</u>	<u>A076049 003</u>	Jan 14, 2005
	<u>EQ 40MG BASE</u>	<u>A076049 004</u>	Jan 14, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 355 (of 424)

QUINIDINE GLUCONATE

INJECTABLE; INJECTION QUINIDINE GLUCONATE			
+ LILLY	80MG/ML	N007529 002	Feb 10, 1989
TABLET, EXTENDED RELEASE; ORAL QUINIDINE GLUCONATE			
BX + MUTUAL PHARM	324MG	A089338 001	Feb 11, 1987
BX WATSON LABS	324MG	A087810 001	Sep 29, 1982

QUINIDINE SULFATE

TABLET; ORAL <u>QUINIDINE SULFATE</u>			
<u>AB</u> MUTUAL PHARM	<u>200MG</u>	<u>A081030</u> <u>001</u>	Apr 14, 1989
<u>AB</u>	<u>300MG</u>	<u>A081031</u> <u>001</u>	Apr 14, 1989
<u>AB</u> SANDOZ	<u>200MG</u>	<u>A088072</u> <u>002</u>	
<u>AB</u>	<u>300MG</u>	<u>A088072</u> <u>001</u>	Sep 26, 1983
<u>AB</u> + WATSON LABS	<u>200MG</u>	<u>A083288</u> <u>001</u>	
<u>AB</u> +	<u>300MG</u>	<u>A085583</u> <u>001</u>	
TABLET, EXTENDED RELEASE; ORAL QUINIDINE SULFATE			
+ TEVA PHARMS	300MG	A040045 001	Jun 30, 1994

QUININE SULFATE

CAPSULE; ORAL QUALAQUIN			
+ AR HOLDING CO INC	324MG	N021799 001	Aug 12, 2005

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL ACIPHEX			
+ EISAI INC	20MG	N020973 002	Aug 19, 1999

RALOXIFENE HYDROCHLORIDE

TABLET; ORAL EVISTA			
+ LILLY	60MG	N020815 001	Dec 09, 1997

RALTEGRAVIR POTASSIUM

TABLET; ORAL ISENTRESS			
+ MERCK SHARP DOHME	EQ 400MG BASE	N022145 001	Oct 12, 2007
TABLET, CHEWABLE; ORAL ISENTRESS			
MERCK SHARP DOHME	EQ 25MG BASE	N203045 001	Dec 21, 2011
+	EQ 100MG BASE	N203045 002	Dec 21, 2011

RAMELTEON

TABLET; ORAL ROZEREM			
+ TAKEDA GLOBAL	8MG	N021782 001	Jul 22, 2005

RAMIPRIL

CAPSULE; ORAL <u>ALTACE</u>			
<u>AB</u> KING PHARMS	<u>1.25MG</u>	<u>N019901</u> <u>001</u>	Jan 28, 1991
<u>AB</u>	<u>2.5MG</u>	<u>N019901</u> <u>002</u>	Jan 28, 1991
<u>AB</u>	<u>5MG</u>	<u>N019901</u> <u>003</u>	Jan 28, 1991
<u>AB</u> +	<u>10MG</u>	<u>N019901</u> <u>004</u>	Jan 28, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 356 (of 424)

RAMIPRIL

CAPSULE; ORAL

RAMIPRIL

AB	APOTEX	<u>1.25MG</u>	<u>A079116</u> <u>001</u>	Jun 20, 2008
AB		<u>2.5MG</u>	<u>A079116</u> <u>002</u>	Jun 20, 2008
AB		<u>5MG</u>	<u>A079116</u> <u>003</u>	Jun 20, 2008
AB		<u>10MG</u>	<u>A079116</u> <u>004</u>	Jun 20, 2008
AB	AUROBINDO PHARMA LTD	<u>1.25MG</u>	<u>A091604</u> <u>001</u>	Jun 08, 2011
AB		<u>2.5MG</u>	<u>A091604</u> <u>002</u>	Jun 08, 2011
AB		<u>5MG</u>	<u>A091604</u> <u>003</u>	Jun 08, 2011
AB		<u>10MG</u>	<u>A091604</u> <u>004</u>	Jun 08, 2011
AB	CIPPLA	<u>1.25MG</u>	<u>A077004</u> <u>001</u>	Aug 07, 2008
AB		<u>2.5MG</u>	<u>A077004</u> <u>002</u>	Aug 07, 2008
AB		<u>5MG</u>	<u>A077004</u> <u>003</u>	Aug 07, 2008
AB		<u>10MG</u>	<u>A077004</u> <u>004</u>	Aug 07, 2008
AB	DR REDDYS LABS LTD	<u>1.25MG</u>	<u>A078191</u> <u>001</u>	Jun 18, 2008
AB		<u>2.5MG</u>	<u>A078191</u> <u>002</u>	Jun 18, 2008
AB		<u>5MG</u>	<u>A078191</u> <u>003</u>	Jun 18, 2008
AB		<u>10MG</u>	<u>A078191</u> <u>004</u>	Jun 18, 2008
AB	INVAGEN PHARMS	<u>1.25MG</u>	<u>A078745</u> <u>001</u>	Jun 18, 2008
AB		<u>2.5MG</u>	<u>A078745</u> <u>002</u>	Jun 18, 2008
AB		<u>5MG</u>	<u>A078745</u> <u>003</u>	Jun 18, 2008
AB		<u>10MG</u>	<u>A078745</u> <u>004</u>	Jun 18, 2008
AB	LUPIN	<u>1.25MG</u>	<u>A077626</u> <u>001</u>	Jun 09, 2008
AB		<u>2.5MG</u>	<u>A077626</u> <u>002</u>	Jun 09, 2008
AB		<u>5MG</u>	<u>A077626</u> <u>003</u>	Jun 09, 2008
AB		<u>10MG</u>	<u>A077626</u> <u>004</u>	Jun 09, 2008
AB	RANBAXY	<u>5MG</u>	<u>A078849</u> <u>001</u>	Mar 06, 2009
AB		<u>10MG</u>	<u>A078849</u> <u>002</u>	Mar 06, 2009
AB	ROXANE	<u>1.25MG</u>	<u>A077900</u> <u>001</u>	Jun 18, 2008
AB		<u>2.5MG</u>	<u>A077900</u> <u>002</u>	Jun 18, 2008
AB		<u>5MG</u>	<u>A077900</u> <u>003</u>	Jun 18, 2008
AB		<u>10MG</u>	<u>A077900</u> <u>004</u>	Jun 18, 2008
AB	SANDOZ	<u>1.25MG</u>	<u>A077514</u> <u>001</u>	Jun 18, 2008
AB		<u>2.5MG</u>	<u>A077514</u> <u>002</u>	Jun 18, 2008
AB		<u>5MG</u>	<u>A077514</u> <u>003</u>	Jun 18, 2008
AB		<u>10MG</u>	<u>A077514</u> <u>004</u>	Jun 18, 2008
AB	TEVA PHARMS	<u>1.25MG</u>	<u>A077470</u> <u>001</u>	Jun 18, 2008
AB		<u>2.5MG</u>	<u>A077470</u> <u>002</u>	Jun 18, 2008
AB		<u>5MG</u>	<u>A077470</u> <u>003</u>	Jun 18, 2008
AB		<u>10MG</u>	<u>A077470</u> <u>004</u>	Jun 18, 2008
AB	WATSON LABS	<u>1.25MG</u>	<u>A076549</u> <u>001</u>	Oct 24, 2005
AB		<u>2.5MG</u>	<u>A076549</u> <u>002</u>	Oct 24, 2005
AB		<u>5MG</u>	<u>A076549</u> <u>003</u>	Oct 24, 2005
AB		<u>10MG</u>	<u>A076549</u> <u>004</u>	Oct 24, 2005
AB	ZYDUS PHARMS USA	<u>1.25MG</u>	<u>A078832</u> <u>001</u>	Sep 02, 2008
AB		<u>2.5MG</u>	<u>A078832</u> <u>002</u>	Sep 02, 2008
AB		<u>5MG</u>	<u>A078832</u> <u>003</u>	Sep 02, 2008
AB		<u>10MG</u>	<u>A078832</u> <u>004</u>	Sep 02, 2008

TABLET; ORAL

ALTACE

AB	KING PHARMS	<u>1.25MG</u>	<u>N022021</u> <u>001</u>	Feb 27, 2007
AB		<u>2.5MG</u>	<u>N022021</u> <u>002</u>	Feb 27, 2007
AB		<u>5MG</u>	<u>N022021</u> <u>003</u>	Feb 27, 2007
AB	+	<u>10MG</u>	<u>N022021</u> <u>004</u>	Feb 27, 2007

RAMIPRIL

AB	MATRIX LABS LTD	<u>1.25MG</u>	<u>A090650</u> <u>001</u>	Jun 30, 2011
AB		<u>2.5MG</u>	<u>A090650</u> <u>002</u>	Jun 30, 2011
AB		<u>5MG</u>	<u>A090650</u> <u>003</u>	Jun 30, 2011
AB		<u>10MG</u>	<u>A090650</u> <u>004</u>	Jun 30, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 357 (of 424)

RAMIPRIL

TABLET; ORAL

RAMIPRIL

<u>AB</u>	ZYDUS PHARMS USA INC	<u>1.25MG</u>	<u>A090697</u>	<u>001</u>	Sep 24, 2009
<u>AB</u>		<u>2.5MG</u>	<u>A090697</u>	<u>002</u>	Sep 24, 2009
<u>AB</u>		<u>5MG</u>	<u>A090697</u>	<u>003</u>	Sep 24, 2009
<u>AB</u>		<u>10MG</u>	<u>A090697</u>	<u>004</u>	Sep 24, 2009

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A075742</u>	<u>001</u>	Nov 29, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075742</u>	<u>002</u>	Nov 29, 2000
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074655</u>	<u>001</u>	Oct 22, 1997
<u>AB</u>	+	<u>EQ 300MG BASE</u>	<u>A074655</u>	<u>002</u>	Oct 22, 1997

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

<u>AP</u>	BEDFORD	<u>EQ 25MG BASE/ML</u>	<u>A077458</u>	<u>001</u>	Feb 16, 2006
<u>AP</u>	BEN VENUE	<u>EQ 25MG BASE/ML</u>	<u>A074777</u>	<u>001</u>	Mar 02, 2005
<u> </u>	<u>ZANTAC</u>				
<u>AP</u>	+ GLAXOSMITHKLINE	<u>EQ 25MG BASE/ML</u>	<u>N019090</u>	<u>001</u>	Oct 19, 1984
	ZANTAC IN PLASTIC CONTAINER				
	+ GLAXOSMITHKLINE	<u>EQ 1MG BASE/ML</u>	<u>N019593</u>	<u>002</u>	Sep 27, 1991

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

<u>AA</u>	ACTAVIS MID ATLANTIC	<u>EQ 15MG BASE/ML</u>	<u>A076124</u>	<u>001</u>	Feb 21, 2007
<u>AA</u>	AMNEAL PHARMS	<u>EQ 15MG BASE/ML</u>	<u>A078312</u>	<u>001</u>	Sep 02, 2008
<u>AA</u>	APOTEX	<u>EQ 15MG BASE/ML</u>	<u>A077602</u>	<u>001</u>	Sep 17, 2007
<u>AA</u>	AUROBINDO PHARM	<u>EQ 15MG BASE/ML</u>	<u>A090623</u>	<u>001</u>	Jul 28, 2010
<u>AA</u>	CYPRESS PHARM	<u>EQ 15MG BASE/ML</u>	<u>A078684</u>	<u>001</u>	Aug 27, 2009
<u>AA</u>	DR REDDYS LABS LTD	<u>EQ 15MG BASE/ML</u>	<u>A090102</u>	<u>001</u>	May 26, 2009
<u>AA</u>	HI TECH PHARMA	<u>EQ 15MG BASE/ML</u>	<u>A091078</u>	<u>001</u>	Mar 22, 2011
<u>AA</u>	PHARM ASSOC	<u>EQ 15MG BASE/ML</u>	<u>A077405</u>	<u>001</u>	Sep 21, 2007
<u>AA</u>	RANBAXY	<u>EQ 15MG BASE/ML</u>	<u>A078448</u>	<u>001</u>	Dec 13, 2007
<u>AA</u>	SILARK	<u>EQ 15MG BASE/ML</u>	<u>A091288</u>	<u>001</u>	Dec 09, 2010
<u>AA</u>	SUN PHARM INDs INC	<u>EQ 15MG BASE/ML</u>	<u>A091091</u>	<u>001</u>	Sep 20, 2011
<u>AA</u>	TARO	<u>EQ 15MG BASE/ML</u>	<u>A077476</u>	<u>001</u>	Jun 13, 2011
<u>AA</u>	TOLMAR	<u>EQ 15MG BASE/ML</u>	<u>A090054</u>	<u>001</u>	Nov 15, 2010
<u>AA</u>	VINTAGE PHARMS	<u>EQ 15MG BASE/ML</u>	<u>A078890</u>	<u>001</u>	Jul 01, 2010
<u>AA</u>	WOCKHARDT	<u>EQ 15MG BASE/ML</u>	<u>A079211</u>	<u>001</u>	May 26, 2009
<u>AA</u>		<u>EQ 15MG BASE/ML</u>	<u>A079212</u>	<u>001</u>	Feb 23, 2009
<u> </u>	<u>ZANTAC</u>				
<u>AA</u>	+ GLAXOSMITHKLINE	<u>EQ 15MG BASE/ML</u>	<u>N019675</u>	<u>001</u>	Dec 30, 1988

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 150MG BASE</u>	<u>A077824</u>	<u>001</u>	Oct 13, 2006
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077824</u>	<u>002</u>	Oct 13, 2006
<u>AB</u>	APOTEX	<u>EQ 150MG BASE</u>	<u>A074680</u>	<u>001</u>	Sep 12, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074680</u>	<u>002</u>	Sep 12, 1997
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 150MG BASE</u>	<u>A076705</u>	<u>001</u>	Jul 27, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076705</u>	<u>002</u>	Jul 27, 2005
<u>AB</u>	GLENMARK GENERICS	<u>EQ 150MG BASE</u>	<u>A078542</u>	<u>001</u>	Nov 19, 2008
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078542</u>	<u>002</u>	Nov 19, 2008
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 150MG BASE</u>	<u>A075165</u>	<u>001</u>	Sep 30, 1998
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075165</u>	<u>002</u>	Sep 30, 1998
<u>AB</u>	MYLAN	<u>EQ 150MG BASE</u>	<u>A074023</u>	<u>001</u>	Aug 22, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074023</u>	<u>002</u>	Aug 22, 1997
<u>AB</u>	PAR PHARM	<u>EQ 150MG BASE</u>	<u>A075180</u>	<u>001</u>	Jan 28, 1999
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075180</u>	<u>002</u>	Jan 28, 1999
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074467</u>	<u>001</u>	Aug 29, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 358 (of 424)

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>EQ 300MG BASE</u>	<u>A074467 002</u>	Aug 29, 1997
<u>AB</u>	TEVA	<u>EQ 150MG BASE</u>	<u>A074488 001</u>	Jul 31, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074488 002</u>	Jul 31, 1997
<u>AB</u>	WATSON LABS	<u>EQ 150MG BASE</u>	<u>A074864 001</u>	Oct 20, 1997
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A077426 001</u>	Dec 19, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074864 002</u>	Oct 20, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077426 002</u>	Dec 19, 2005
<u>AB</u>	WOCKHARDT	<u>EQ 150MG BASE</u>	<u>A075208 001</u>	Dec 17, 1998
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078701 001</u>	Nov 12, 2009
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075208 002</u>	Dec 17, 1998
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078701 002</u>	Dec 11, 2009
	<u>ZANTAC 150</u>			
<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 150MG BASE</u>	<u>N018703 001</u>	Jun 09, 1983
	<u>ZANTAC 300</u>			
<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 300MG BASE</u>	<u>N018703 002</u>	Dec 09, 1985
	TABLET, EFFERVESCENT; ORAL			
	ZANTAC 25			
	+ GLAXOSMITHKLINE	EQ 25MG BASE	N020251 003	Apr 01, 2004

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

GILEAD 500MG

+ 1GM

N021526 002 Jan 27, 2006
N021526 001 Feb 12, 2007RASAGILINE MESYLATE

TABLET; ORAL

AZILECT

TEVA EQ 0.5MG BASE

+ EQ 1MG BASE

N021641 001 May 16, 2006
N021641 002 May 16, 2006REGADENOSON

SOLUTION; INTRAVENOUS

LEXISCAN

+ ASTELLAS 0.4MG/5ML (0.08MG/ML)

N022161 001 Apr 10, 2008

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ULTIVA

MYLAN INSTITUTIONAL EQ 1MG BASE/VIAL

EQ 2MG BASE/VIAL

+ EQ 5MG BASE/VIAL

N020630 001 Jul 12, 1996
N020630 002 Jul 12, 1996
N020630 003 Jul 12, 1996REPAGLINIDE

TABLET; ORAL

PRANDIN

NOVO NORDISK INC 0.5MG

1MG

+ 2MG

N020741 001 Dec 22, 1997
N020741 002 Dec 22, 1997
N020741 003 Dec 22, 1997RESERPINE

TABLET; ORAL

RESERPINE

BP SANDOZ 0.1MG

BP + 0.25MG

N009838 001
N009838 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 359 (of 424)

RESERPINE

TABLET; ORAL SERPALAN			
BP LANNETT	0.1MG	N010124 001	
BP	0.25MG	N010124 002	

RETAPAMULIN

OINTMENT; TOPICAL ALTABAX			
+ GLAXO GRP LTD	1%	N022055 001	Apr 12, 2007

RIBAVIRIN

CAPSULE; ORAL <u>REBETOL</u>			
<u>AB</u> + SCHERING PLOUGH RES	<u>200MG</u>	<u>N020903 002</u>	Jul 25, 2001
<u>RIBASPHERE</u>			
<u>AB</u> THREE RIVERS PHARMS	<u>200MG</u>	<u>A076203 001</u>	Apr 06, 2004
<u>RIBAVARIN</u>			
<u>AB</u> AUROBINDO PHARMA	<u>200MG</u>	<u>A079117 001</u>	Sep 17, 2009
<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
REBETOL			
+ SCHERING PLOUGH RES	200MG	N020903 001	Jun 03, 1998
FOR SOLUTION; INHALATION VIRAZOLE			
+ VALEANT PHARM INTL	6GM/VIAL	N018859 001	Dec 31, 1985
SOLUTION; ORAL REBETOL			
+ SCHERING	40MG/ML	N021546 001	Jul 29, 2003
CAPSULE; ORAL <u>COPEGUS</u>			
<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> SANDOZ	<u>200MG</u>	<u>A077743 001</u>	Oct 03, 2006
<u>AB</u> TEVA	<u>200MG</u>	<u>A077053 001</u>	Dec 05, 2005
<u>AB</u> THREE RIVERS PHARMS	<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> 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>600MG</u>	<u>A077094 003</u>	Mar 16, 2007
RIBAVIRIN			
ZYDUS PHARMS USA	500MG	A077094 004	Apr 18, 2008

RIFABUTIN

CAPSULE; ORAL MYCOBUTIN			
+ PHARMACIA AND UPJOHN	150MG	N050689 001	Dec 23, 1992

RIFAMPIN

CAPSULE; ORAL <u>RIFADIN</u>			
<u>AB</u> SANOFI AVENTIS US	<u>150MG</u>	<u>A062303 001</u>	
<u>AB</u> +	<u>300MG</u>	<u>N050420 001</u>	
<u>RIFAMPIN</u>			
<u>AB</u> LANNETT	<u>150MG</u>	<u>A065390 001</u>	Mar 28, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 360 (of 424)

RIFAMPIN

CAPSULE; ORAL

RIFAMPIN

<u>AB</u>	LANNETT	<u>300MG</u>	<u>A065390</u>	<u>002</u>	Mar 28, 2008
<u>AB</u>	SANDOZ	<u>150MG</u>	<u>A064150</u>	<u>002</u>	Jan 02, 1998
<u>AB</u>		<u>300MG</u>	<u>A064150</u>	<u>001</u>	May 28, 1997
<u>AB</u>	VERSAPHARM	<u>150MG</u>	<u>A065028</u>	<u>001</u>	Mar 14, 2001
<u>AB</u>		<u>300MG</u>	<u>A065028</u>	<u>002</u>	Mar 14, 2001

INJECTABLE; INJECTION

RIFADIN

<u>AP</u> + SANOFI AVENTIS US		<u>600MG/VIAL</u>	<u>N050627</u>	<u>001</u>	May 25, 1989
<u>AP</u>	BEDFORD	<u>600MG/VIAL</u>	<u>A064217</u>	<u>001</u>	Oct 29, 1999
<u>AP</u>	PFIZER	<u>600MG/VIAL</u>	<u>A065421</u>	<u>001</u>	May 22, 2008
<u>AP</u>	VERSAPHARM INC	<u>600MG/VIAL</u>	<u>A065502</u>	<u>001</u>	Sep 21, 2010

RIFAPENTINE

TABLET; ORAL

PRIFTIN

+ SANOFI AVENTIS US 150MG

N021024 001 Jun 22, 1998

RIFAXIMIN

TABLET; ORAL

XIFAXAN

+ SALIX PHARMS 200MG
+ 550MGN021361 001 May 25, 2004
N022554 001 Mar 24, 2010RILPIVIRINE HYDROCHLORIDE

TABLET; ORAL

EDURANT

+ TIBOTEC EQ 25MG BASE

N202022 001 May 20, 2011

RILUZOLE

TABLET; ORAL

RILUTEK

<u>AB</u> + SANOFI AVENTIS US		<u>50MG</u>	<u>N020599</u>	<u>001</u>	Dec 12, 1995
<u>AB</u>	IMPAK LABS	<u>50MG</u>	<u>A076173</u>	<u>001</u>	Jan 29, 2003

RIMANTADINE HYDROCHLORIDE

TABLET; ORAL

FLUMADINE

<u>AB</u> + CARACO		<u>100MG</u>	<u>N019649</u>	<u>001</u>	Sep 17, 1993
<u>AB</u>	COREPHARMA	<u>100MG</u>	<u>A075916</u>	<u>001</u>	Nov 02, 2001

AB IMPAX LABS 100MGA076132 001 Aug 30, 2002RIMEXOLONE

SUSPENSION/DROPS; OPHTHALMIC

+ ALCON 1% N020474 001 Dec 30, 1994

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

<u>AB</u>	WARNER CHILCOTT	<u>5MG</u>	<u>N020835</u>	<u>002</u>	Apr 14, 2000
<u>AB</u>		<u>30MG</u>	<u>N020835</u>	<u>001</u>	Mar 27, 1998
<u>AB</u> +		<u>35MG</u>	<u>N020835</u>	<u>003</u>	May 25, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 361 (of 424)

RISEDRONATE SODIUM

TABLET; ORAL

RISEDRONATE SODIUM

<u>AB</u>	TEVA PHARMS	<u>5MG</u>	<u>A077132</u>	<u>001</u>	Oct 05, 2007
<u>AB</u>		<u>30MG</u>	<u>A077132</u>	<u>002</u>	Oct 05, 2007
<u>AB</u>		<u>35MG</u>	<u>A077132</u>	<u>003</u>	Oct 05, 2007
	ACTONEL				
+ Warner Chilcott		150MG	N020835	005	Apr 22, 2008
	TABLET, DELAYED RELEASE; ORAL				
	ATELVIA				
+ Warner Chilcott		35MG	N022560	001	Oct 08, 2010

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</u>	<u>001</u>	Jun 10, 1996
	<u>RISPERIDONE</u>			
<u>AA</u> AMNEAL PHARMS	<u>1MG/ML</u>	<u>A091384</u>	<u>001</u>	May 25, 2011
<u>AA</u> APOTEX INC	<u>1MG/ML</u>	<u>A077719</u>	<u>001</u>	Jul 29, 2009
<u>AA</u> AUROBINDO PHARMA	<u>1MG/ML</u>	<u>A078452</u>	<u>001</u>	Sep 04, 2009
<u>AA</u> DR REDDYS LABS LTD	<u>1MG/ML</u>	<u>A078909</u>	<u>001</u>	Jul 29, 2009
<u>AA</u> PRECISION DOSE	<u>1MG/ML</u>	<u>A076797</u>	<u>001</u>	Jun 28, 2010
<u>AA</u> ROXANE	<u>1MG/ML</u>	<u>A076904</u>	<u>001</u>	Jul 29, 2009
<u>AA</u> TARO	<u>1MG/ML</u>	<u>A090347</u>	<u>001</u>	Feb 07, 2011
<u>AA</u> TEVA	<u>1MG/ML</u>	<u>A076440</u>	<u>001</u>	Jan 30, 2009
<u>AA</u> VINTAGE	<u>1MG/ML</u>	<u>A079158</u>	<u>001</u>	Dec 03, 2010
<u>AA</u> WOCKHARDT	<u>1MG/ML</u>	<u>A078744</u>	<u>001</u>	Oct 08, 2009

TABLET; ORAL

RISPERDAL

<u>AB</u> JANSSEN PHARMS	<u>0.25MG</u>	<u>N020272</u>	<u>008</u>	May 10, 1999
<u>AB</u>	<u>0.5MG</u>	<u>N020272</u>	<u>007</u>	Jan 27, 1999
<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

RISPERIDONE

<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

PRESCRIPTION DRUG PRODUCT LIST

3 - 362 (of 424)

RISPERIDONE

TABLET; ORAL

RISPERIDONE

AB	CIPLA	<u>0.25MG</u>	A077543	001	May 18, 2011
AB		<u>0.5MG</u>	A077543	002	May 18, 2011
AB		<u>1MG</u>	A077543	003	May 18, 2011
AB		<u>2MG</u>	A077543	004	May 18, 2011
AB		<u>3MG</u>	A077543	005	May 18, 2011
AB		<u>4MG</u>	A077543	006	May 18, 2011
AB	DR REDDYS LABS LTD	<u>0.25MG</u>	A076879	001	Oct 24, 2008
AB		<u>0.5MG</u>	A076879	002	Oct 24, 2008
AB		<u>1MG</u>	A076879	003	Oct 24, 2008
AB		<u>2MG</u>	A076879	004	Oct 24, 2008
AB		<u>3MG</u>	A076879	005	Oct 24, 2008
AB		<u>4MG</u>	A076879	006	Oct 24, 2008
AB	MYLAN	<u>0.25MG</u>	A076288	001	Sep 15, 2008
AB		<u>0.5MG</u>	A076288	002	Sep 15, 2008
AB		<u>1MG</u>	A076288	003	Sep 15, 2008
AB		<u>2MG</u>	A076288	004	Sep 15, 2008
AB		<u>3MG</u>	A076288	005	Sep 15, 2008
AB		<u>4MG</u>	A076288	006	Sep 15, 2008
AB	PLIVA HRVATSKA DOO	<u>0.25MG</u>	A077769	001	Oct 16, 2008
AB		<u>0.5MG</u>	A077769	002	Oct 16, 2008
AB		<u>1MG</u>	A077769	003	Oct 16, 2008
AB		<u>2MG</u>	A077769	004	Oct 16, 2008
AB		<u>3MG</u>	A077769	005	Oct 16, 2008
AB		<u>4MG</u>	A077769	006	Oct 16, 2008
AB	PRINSTON INC	<u>0.25MG</u>	A077493	001	Nov 29, 2011
AB		<u>0.5MG</u>	A077493	002	Nov 29, 2011
AB		<u>1MG</u>	A077493	003	Nov 29, 2011
AB		<u>2MG</u>	A077493	004	Nov 29, 2011
AB		<u>3MG</u>	A077493	005	Nov 29, 2011
AB		<u>4MG</u>	A077493	006	Nov 29, 2011
AB	SANDOZ	<u>0.25MG</u>	A078528	001	Oct 16, 2009
AB		<u>0.5MG</u>	A078528	002	Oct 16, 2009
AB		<u>1MG</u>	A078528	003	Oct 16, 2009
AB		<u>2MG</u>	A078528	004	Oct 16, 2009
AB		<u>3MG</u>	A078528	005	Oct 16, 2009
AB		<u>4MG</u>	A078528	006	Oct 16, 2009
AB	TEVA	<u>0.25MG</u>	A076228	001	Jun 30, 2008
AB		<u>0.5MG</u>	A076228	002	Jun 30, 2008
AB		<u>1MG</u>	A076228	003	Jun 30, 2008
AB		<u>2MG</u>	A076228	004	Jun 30, 2008
AB		<u>3MG</u>	A076228	005	Jun 30, 2008
AB		<u>4MG</u>	A076228	006	Jun 30, 2008
AB	TORRENT PHARMS	<u>0.25MG</u>	A079088	001	Oct 30, 2008
AB		<u>0.5MG</u>	A079088	002	Oct 30, 2008
AB		<u>1MG</u>	A079088	003	Oct 30, 2008
AB		<u>2MG</u>	A079088	004	Oct 30, 2008
AB		<u>3MG</u>	A079088	005	Oct 30, 2008
AB		<u>4MG</u>	A079088	006	Oct 30, 2008
AB	VINTAGE	<u>0.25MG</u>	A078707	001	Dec 29, 2008
AB		<u>0.5MG</u>	A078707	002	Dec 29, 2008
AB		<u>1MG</u>	A078707	003	Dec 29, 2008
AB		<u>2MG</u>	A078707	004	Dec 29, 2008
AB		<u>3MG</u>	A078707	005	Dec 29, 2008
AB		<u>4MG</u>	A078707	006	Dec 29, 2008
AB	WATSON LABS	<u>0.25MG</u>	A077860	001	Dec 05, 2008
AB		<u>0.5MG</u>	A077860	002	Dec 05, 2008
AB		<u>1MG</u>	A077860	003	Dec 05, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 363 (of 424)

RISPERIDONE

TABLET; ORAL

RISPERIDONE

AB	WATSON LABS	<u>2MG</u>	A077860	004	Dec 05, 2008
AB		<u>3MG</u>	A077860	005	Dec 05, 2008
AB		<u>4MG</u>	A077860	006	Dec 05, 2008
AB	WEST WARD PHARMS	<u>0.25MG</u>	A078740	001	May 29, 2009
AB		<u>0.5MG</u>	A078740	002	May 29, 2009
AB		<u>1MG</u>	A078740	003	May 29, 2009
AB		<u>2MG</u>	A078740	004	May 29, 2009
AB		<u>3MG</u>	A078740	005	May 29, 2009
AB		<u>4MG</u>	A078740	006	May 29, 2009
AB	WOCHARDT	<u>0.25MG</u>	A078871	001	Oct 09, 2008
AB		<u>0.5MG</u>	A078871	002	Oct 09, 2008
AB		<u>1MG</u>	A078871	003	Oct 09, 2008
AB		<u>2MG</u>	A078871	004	Oct 09, 2008
AB		<u>3MG</u>	A078871	005	Oct 09, 2008
AB		<u>4MG</u>	A078871	006	Oct 09, 2008
AB	ZYDUS PHARMS USA INC	<u>0.25MG</u>	A078040	001	Oct 16, 2008
AB		<u>0.5MG</u>	A078040	002	Oct 16, 2008
AB		<u>1MG</u>	A078040	003	Oct 16, 2008
AB		<u>2MG</u>	A078040	004	Oct 16, 2008
AB		<u>3MG</u>	A078040	005	Oct 16, 2008
AB		<u>4MG</u>	A078040	006	Oct 16, 2008

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

AB	JANSSEN PHARMS	<u>0.5MG</u>	N021444	001	Apr 02, 2003
AB	+	<u>1MG</u>	N021444	002	Apr 02, 2003
AB		<u>2MG</u>	N021444	003	Apr 02, 2003
AB		<u>3MG</u>	N021444	004	Dec 23, 2004
AB		<u>4MG</u>	N021444	005	Dec 23, 2004

RISPERIDONE

AB	DR REDDYS LABS LTD	<u>0.5MG</u>	A077328	001	Feb 24, 2009
AB		<u>1MG</u>	A077328	002	Oct 05, 2009
AB		<u>2MG</u>	A077328	003	Feb 24, 2009
AB	JUBILANT ORGANOSYS	<u>0.5MG</u>	A090839	001	Nov 04, 2011
AB		<u>1MG</u>	A090839	002	Nov 04, 2011
AB		<u>2MG</u>	A090839	003	Nov 04, 2011
AB		<u>3MG</u>	A090839	004	Nov 04, 2011
AB		<u>4MG</u>	A090839	005	Nov 04, 2011
AB	MYLAN	<u>0.5MG</u>	A091537	001	Mar 30, 2011
AB		<u>1MG</u>	A091537	002	Mar 30, 2011
AB		<u>2MG</u>	A091537	003	Mar 30, 2011
AB		<u>3MG</u>	A091537	004	Mar 30, 2011
AB		<u>4MG</u>	A091537	005	Mar 30, 2011
AB	PAR PHARM	<u>0.5MG</u>	A077494	002	Apr 30, 2009
AB		<u>1MG</u>	A077494	003	Oct 26, 2009
AB		<u>2MG</u>	A077494	004	Apr 30, 2009
AB		<u>3MG</u>	A077494	005	Apr 30, 2009
AB		<u>4MG</u>	A077494	006	Apr 30, 2009
AB	RANBAXY	<u>0.5MG</u>	A077542	001	Aug 06, 2010
AB		<u>1MG</u>	A077542	002	Aug 06, 2010
AB		<u>2MG</u>	A077542	003	Aug 06, 2010
AB		<u>3MG</u>	A078474	001	Aug 06, 2010
AB		<u>4MG</u>	A078474	002	Aug 06, 2010
AB	SANDOZ	<u>0.5MG</u>	A078116	001	Dec 22, 2009
AB		<u>1MG</u>	A078116	002	Dec 22, 2009
AB		<u>2MG</u>	A078116	003	Dec 22, 2009
AB		<u>3MG</u>	A078116	004	Dec 22, 2009
AB		<u>4MG</u>	A078116	005	Dec 22, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 364 (of 424)

RISPERIDONE

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERIDONE

<u>AB</u>	WATSON LABS FLORIDA	<u>0.5MG</u>	<u>A076996</u>	<u>001</u>	Apr 19, 2011
<u>AB</u>		<u>1MG</u>	<u>A076996</u>	<u>002</u>	Apr 19, 2011
<u>AB</u>		<u>2MG</u>	<u>A076996</u>	<u>003</u>	Apr 19, 2011
<u>AB</u>		<u>3MG</u>	<u>A076996</u>	<u>004</u>	Apr 19, 2011
<u>AB</u>		<u>4MG</u>	<u>A076996</u>	<u>005</u>	Apr 19, 2011
<u>AB</u>	ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A078516</u>	<u>001</u>	May 01, 2009
<u>AB</u>		<u>2MG</u>	<u>A078516</u>	<u>003</u>	May 01, 2009
	RISPERIDONE				
	PAR PHARM	0.25MG	A077494	001	Apr 30, 2009

RITONAVIR

CAPSULE; ORAL

NORVIR

+ ABBOTT	100MG	N020945	001	Jun 29, 1999
----------	-------	---------	-----	--------------

SOLUTION; ORAL

NORVIR

+ ABBOTT	80MG/ML	N020659	001	Mar 01, 1996
----------	---------	---------	-----	--------------

TABLET; ORAL

NORVIR

+ ABBOTT LABS	100MG	N022417	001	Feb 10, 2010
---------------	-------	---------	-----	--------------

RIVAROXABAN

TABLET; ORAL

XARELTO

+ JANSSEN PHARMS	10MG	N022406	001	Jul 01, 2011
	15MG	N202439	001	Nov 04, 2011
+	20MG	N202439	002	Nov 04, 2011

RIVASTIGMINE

FILM, EXTENDED RELEASE; TRANSDERMAL

EXELON

NOVARTIS	4.6MG/24HR	N022083	001	Jul 06, 2007
+	9.5MG/24HR	N022083	002	Jul 06, 2007

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

EXELON

<u>AB</u> + NOVARTIS	<u>EQ 1.5MG BASE</u>	<u>N020823</u>	<u>003</u>	Apr 21, 2000
<u>AB</u>	<u>EQ 3MG BASE</u>	<u>N020823</u>	<u>004</u>	Apr 21, 2000
<u>AB</u>	<u>EQ 4.5MG BASE</u>	<u>N020823</u>	<u>005</u>	Apr 21, 2000
<u>AB</u>	<u>EQ 6MG BASE</u>	<u>N020823</u>	<u>006</u>	Apr 21, 2000

RIVASTIGMINE TARTRATE

<u>AB</u> DR REDDYS LABS INC	<u>EQ 1.5MG BASE</u>	<u>A077130</u>	<u>001</u>	Oct 31, 2007
<u>AB</u>	<u>EQ 3MG BASE</u>	<u>A077130</u>	<u>002</u>	Oct 31, 2007
<u>AB</u>	<u>EQ 4.5MG BASE</u>	<u>A077130</u>	<u>003</u>	Oct 31, 2007
<u>AB</u>	<u>EQ 6MG BASE</u>	<u>A077130</u>	<u>004</u>	Oct 31, 2007
<u>AB</u> SUN PHARM INDs	<u>EQ 1.5MG BASE</u>	<u>A077131</u>	<u>001</u>	Oct 22, 2007
<u>AB</u>	<u>EQ 3MG BASE</u>	<u>A077131</u>	<u>002</u>	Oct 22, 2007
<u>AB</u>	<u>EQ 4.5MG BASE</u>	<u>A077131</u>	<u>003</u>	Oct 22, 2007
<u>AB</u>	<u>EQ 6MG BASE</u>	<u>A077131</u>	<u>004</u>	Oct 22, 2007
<u>AB</u> WATSON LABS	<u>EQ 1.5MG BASE</u>	<u>A077129</u>	<u>001</u>	Jan 08, 2008
<u>AB</u>	<u>EQ 3MG BASE</u>	<u>A077129</u>	<u>002</u>	Jan 08, 2008
<u>AB</u>	<u>EQ 4.5MG BASE</u>	<u>A077129</u>	<u>003</u>	Jan 08, 2008
<u>AB</u>	<u>EQ 6MG BASE</u>	<u>A077129</u>	<u>004</u>	Jan 08, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 365 (of 424)

RIVASTIGMINE TARTRATE

SOLUTION; ORAL
 EXELON
 + NOVARTIS EQ 2MG BASE/ML N021025 001 Apr 21, 2000

RIZATRIPTAN BENZOATE

TABLET; ORAL
 MAXALT
 MERCK EQ 5MG BASE N020864 001 Jun 29, 1998
 + EQ 10MG BASE N020864 002 Jun 29, 1998

TABLET, ORALLY DISINTEGRATING; ORAL
 MAXALT-MLT
 MERCK EQ 5MG BASE N020865 001 Jun 29, 1998
 + EQ 10MG BASE N020865 002 Jun 29, 1998

ROCURONIUM BROMIDE

INJECTABLE; INJECTION
ROCURONIUM BROMIDE
AP APP PHARMS 50MG/5ML (10MG/ML) A078651 001 Dec 29, 2008
AP 100MG/10ML (10MG/ML) A078651 002 Dec 29, 2008
AP BIONICHE PHARMA USA 50MG/5ML (10MG/ML) A079199 001 Nov 26, 2008
AP 100MG/10ML (10MG/ML) A079199 002 Nov 26, 2008
AP HOSPIRA 50MG/5ML (10MG/ML) A078519 001 Nov 26, 2008
AP 100MG/10ML (10MG/ML) A078519 002 Nov 26, 2008
AP SAGENT STRIDES 50MG/5ML (10MG/ML) A091458 001 Jul 28, 2010
AP 100MG/10ML (10MG/ML) A091458 002 Jul 28, 2010
AP SANDOZ 50MG/5ML (10MG/ML) A079195 001 Dec 05, 2008
AP 100MG/10ML (10MG/ML) A079195 002 Dec 05, 2008
AP TEVA PARENTERAL 50MG/5ML (10MG/ML) A078717 001 Nov 26, 2008
AP 100MG/10ML (10MG/ML) A078717 002 Nov 26, 2008
ZEMURON
AP + SCHERING 50MG/5ML (10MG/ML) N020214 001 Mar 17, 1994
AP + 100MG/10ML (10MG/ML) N020214 003 Mar 17, 1994

ROFLUMILAST

TABLET; ORAL
 DALIRESP
 + FOREST RES INST INC 500MCG N022522 001 Feb 28, 2011

ROMIDEPSIN

POWDER; IV (INFUSION)
 + CELGENE 10MG/VIAL N022393 001 Nov 05, 2009

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL
REQUIP
AB + GLAXOSMITHKLINE EQ 0.25MG BASE N020658 001 Sep 19, 1997
AB EQ 0.5MG BASE N020658 002 Sep 19, 1997
AB EQ 1MG BASE N020658 003 Sep 19, 1997
AB EQ 2MG BASE N020658 004 Sep 19, 1997
AB EQ 3MG BASE N020658 006 Jan 27, 1999
AB EQ 4MG BASE N020658 007 Jan 27, 1999
AB EQ 5MG BASE N020658 005 Sep 19, 1997
ROPINIROLE HYDROCHLORIDE
AB ALEMBIC LTD EQ 0.25MG BASE A090429 001 Mar 24, 2010
AB EQ 0.5MG BASE A090429 002 Mar 24, 2010
AB EQ 1MG BASE A090429 003 Mar 24, 2010
AB EQ 2MG BASE A090429 004 Mar 24, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 366 (of 424)

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

AB	ALEMBIC LTD	<u>EQ 3MG BASE</u>	<u>A090429 005</u>	Mar 24, 2010
AB		<u>EQ 4MG BASE</u>	<u>A090429 006</u>	Mar 24, 2010
AB		<u>EQ 5MG BASE</u>	<u>A090429 007</u>	Mar 24, 2010
AB	COREPHARMA	<u>EQ 0.25MG BASE</u>	<u>A078230 001</u>	May 20, 2008
AB		<u>EQ 0.5MG BASE</u>	<u>A078230 002</u>	May 20, 2008
AB		<u>EQ 1MG BASE</u>	<u>A078230 003</u>	May 20, 2008
AB		<u>EQ 2MG BASE</u>	<u>A078230 004</u>	May 20, 2008
AB		<u>EQ 3MG BASE</u>	<u>A078230 005</u>	May 20, 2008
AB		<u>EQ 4MG BASE</u>	<u>A078230 006</u>	May 20, 2008
AB		<u>EQ 5MG BASE</u>	<u>A078230 007</u>	May 20, 2008
AB	GLENMARK GENERICS	<u>EQ 0.25MG BASE</u>	<u>A090135 001</u>	Feb 25, 2010
AB		<u>EQ 0.5MG BASE</u>	<u>A090135 002</u>	Feb 25, 2010
AB		<u>EQ 1MG BASE</u>	<u>A090135 003</u>	Feb 25, 2010
AB		<u>EQ 2MG BASE</u>	<u>A090135 004</u>	Feb 25, 2010
AB		<u>EQ 3MG BASE</u>	<u>A090135 005</u>	Feb 25, 2010
AB		<u>EQ 4MG BASE</u>	<u>A090135 006</u>	Feb 25, 2010
AB		<u>EQ 5MG BASE</u>	<u>A090135 007</u>	Feb 25, 2010
AB	MYLAN	<u>EQ 0.25MG BASE</u>	<u>A078881 001</u>	May 05, 2008
AB		<u>EQ 0.5MG BASE</u>	<u>A078881 002</u>	May 05, 2008
AB		<u>EQ 1MG BASE</u>	<u>A078881 003</u>	May 05, 2008
AB		<u>EQ 2MG BASE</u>	<u>A078881 004</u>	May 05, 2008
AB		<u>EQ 3MG BASE</u>	<u>A078881 005</u>	May 05, 2008
AB		<u>EQ 4MG BASE</u>	<u>A078881 006</u>	May 05, 2008
AB		<u>EQ 5MG BASE</u>	<u>A078881 007</u>	May 19, 2008
AB	PRINSTON INC	<u>EQ 0.25MG BASE</u>	<u>A078110 001</u>	May 05, 2008
AB		<u>EQ 0.5MG BASE</u>	<u>A078110 002</u>	May 05, 2008
AB		<u>EQ 1MG BASE</u>	<u>A078110 003</u>	May 05, 2008
AB		<u>EQ 2MG BASE</u>	<u>A078110 004</u>	May 05, 2008
AB		<u>EQ 3MG BASE</u>	<u>A078110 005</u>	May 05, 2008
AB		<u>EQ 4MG BASE</u>	<u>A078110 006</u>	May 05, 2008
AB		<u>EQ 5MG BASE</u>	<u>A078110 007</u>	Jul 11, 2008
AB	ROXANE	<u>EQ 0.25MG BASE</u>	<u>A077852 001</u>	May 05, 2008
AB		<u>EQ 0.5MG BASE</u>	<u>A077852 002</u>	May 05, 2008
AB		<u>EQ 1MG BASE</u>	<u>A077852 003</u>	May 05, 2008
AB		<u>EQ 2MG BASE</u>	<u>A077852 004</u>	May 05, 2008
AB		<u>EQ 3MG BASE</u>	<u>A077852 005</u>	May 05, 2008
AB		<u>EQ 4MG BASE</u>	<u>A077852 006</u>	May 05, 2008
AB		<u>EQ 5MG BASE</u>	<u>A077852 007</u>	May 19, 2008
AB	TEVA	<u>EQ 0.25MG BASE</u>	<u>A077460 001</u>	May 05, 2008
AB		<u>EQ 0.5MG BASE</u>	<u>A077460 002</u>	May 05, 2008
AB		<u>EQ 1MG BASE</u>	<u>A077460 003</u>	May 05, 2008
AB		<u>EQ 2MG BASE</u>	<u>A077460 004</u>	May 05, 2008
AB		<u>EQ 3MG BASE</u>	<u>A077460 005</u>	May 05, 2008
AB		<u>EQ 4MG BASE</u>	<u>A077460 006</u>	May 05, 2008
AB		<u>EQ 5MG BASE</u>	<u>A077460 007</u>	May 19, 2008
AB	WOCKHARDT	<u>EQ 0.25MG BASE</u>	<u>A079050 001</u>	May 29, 2008
AB		<u>EQ 0.5MG BASE</u>	<u>A079050 002</u>	May 29, 2008
AB		<u>EQ 1MG BASE</u>	<u>A079050 003</u>	May 29, 2008
AB		<u>EQ 2MG BASE</u>	<u>A079050 004</u>	May 29, 2008
AB		<u>EQ 3MG BASE</u>	<u>A079050 005</u>	May 29, 2008
AB		<u>EQ 4MG BASE</u>	<u>A079050 006</u>	May 29, 2008
AB		<u>EQ 5MG BASE</u>	<u>A079050 007</u>	May 29, 2008
AB	ZYDUS PHARMS USA INC	<u>EQ 0.25MG BASE</u>	<u>A090411 001</u>	Jun 01, 2009
AB		<u>EQ 0.5MG BASE</u>	<u>A090411 002</u>	Jun 01, 2009
AB		<u>EQ 1MG BASE</u>	<u>A090411 003</u>	Jun 01, 2009
AB		<u>EQ 2MG BASE</u>	<u>A090411 004</u>	Jun 01, 2009
AB		<u>EQ 3MG BASE</u>	<u>A090411 005</u>	Jun 01, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 367 (of 424)

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>	ZYDUS PHARMS USA INC	<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

+ SMITHKLINE BEECHAM	EQ 2MG BASE	N022008 001	Jun 13, 2008
	EQ 4MG BASE	N022008 003	Jun 13, 2008
	EQ 6MG BASE	N022008 006	Apr 10, 2009
	EQ 8MG BASE	N022008 004	Jun 13, 2008
	EQ 12MG BASE	N022008 005	Oct 31, 2008

ROPIVACAINE HYDROCHLORIDE MONOHYDRATE

INJECTABLE; INJECTION

NAROPIN

APP PHARMS	2MG/ML	N020533 001	Sep 24, 1996
	5MG/ML	N020533 003	Sep 24, 1996
	7.5MG/ML	N020533 004	Sep 24, 1996
+	10MG/ML	N020533 005	Sep 24, 1996

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

SB PHARMCO	EQ 2MG BASE	N021071 002	May 25, 1999
	EQ 4MG BASE	N021071 003	May 25, 1999
+	EQ 8MG BASE	N021071 004	May 25, 1999

ROSUVASTATIN CALCIUM

TABLET; ORAL

CRESTOR

IPR	5MG	N021366 002	Aug 12, 2003
	10MG	N021366 003	Aug 12, 2003
	20MG	N021366 004	Aug 12, 2003
+	40MG	N021366 005	Aug 12, 2003

RUBIDIUM CHLORIDE RB-82

INJECTABLE; INJECTION

CARDIOGEN-82

BRACCO	N/A	N019414 001	Dec 29, 1989
--------	-----	-------------	--------------

RUFINAMIDE

SUSPENSION; ORAL

BANZEL

+ EISAI INC	40MG/ML	N201367 001	Mar 03, 2011
-------------	---------	-------------	--------------

TABLET; ORAL

BANZEL

EISAI INC	200MG	N021911 002	Nov 14, 2008
+	400MG	N021911 003	Nov 14, 2008

RUXOLITINIB PHOSPHATE

TABLET; ORAL

JAKAFI

INCYTE CORP	EQ 5MG BASE	N202192 001	Nov 16, 2011
	EQ 10MG BASE	N202192 002	Nov 16, 2011
	EQ 15MG BASE	N202192 003	Nov 16, 2011
	EQ 20MG BASE	N202192 004	Nov 16, 2011
+	EQ 25MG BASE	N202192 005	Nov 16, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 368 (of 424)

SACROSIDASE

SOLUTION; ORAL
 SUCRAID
 + QOL MEDCL 8,500 IU/ML N020772 001 Apr 09, 1998

SAFFLOWER OIL; SOYBEAN OIL

INJECTABLE; INJECTION
 LIPOSYN II 10%
 + HOSPIRA 5%;5% (5GM/100ML) N018997 001 Aug 27, 1984
 LIPOSYN II 20%
 + HOSPIRA 10%;10% (10GM/100ML) N018991 001 Aug 27, 1984

SALMETEROL XINAFOATE

POWDER; INHALATION
 SEREVENT
 + GLAXO GRP LTD EQ 0.05MG BASE/INH N020692 001 Sep 19, 1997

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION
 QUADRAMET
 + EUSA PHARMA USA 50mCi/ML N020570 001 Mar 28, 1997

SAPROPTERIN DIHYDROCHLORIDE

TABLET; ORAL
 KUVAN
 + BIOMARIN PHARM 100MG N022181 001 Dec 13, 2007

SAQUINAVIR MESYLATE

CAPSULE; ORAL
 INVIRASE
 + HOFFMANN LA ROCHE EQ 200MG BASE N020628 001 Dec 06, 1995

TABLET; ORAL
 INVIRASE
 + ROCHE EQ 500MG BASE N021785 001 Dec 17, 2004

SAXAGLIPTIN HYDROCHLORIDE

TABLET; ORAL
 ONGLYZA
 BRISTOL MYERS SQUIBB EQ 2.5MG BASE N022350 001 Jul 31, 2009
 + EQ 5MG BASE N022350 002 Jul 31, 2009

SCOPOLAMINE

FILM, EXTENDED RELEASE; TRANSDERMAL
 TRANSDERM SCOP
 + NOVARTIS 1MG/72HR N017874 001

SECOBARBITAL SODIUM

CAPSULE; ORAL
 SECONAL SODIUM
 + MARATHON PHARMS 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

PRESCRIPTION DRUG PRODUCT LIST

3 - 369 (of 424)

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

ELDEPRYL

<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

SELEGILINE HYDROCHLORIDE

<u>AB</u> + APOTEX INC	<u>5MG</u>	<u>A074871 001</u>	Jun 06, 1997
<u>AB</u> DAVA PHARMS INC	<u>5MG</u>	<u>A074641 001</u>	Aug 02, 1996
<u>AB</u> MYLAN	<u>5MG</u>	<u>A074866 001</u>	Nov 26, 1997
<u>AB</u> STASON	<u>5MG</u>	<u>A074912 001</u>	Apr 30, 1998

TABLET, ORALLY DISINTEGRATING; ORAL
ZELAPAR

+ VALEANT PHARM INTL	1.25MG	N021479 001	Jun 14, 2006
----------------------	--------	-------------	--------------

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

SELENIUM SULFIDE

<u>AT</u> PERRIGO NEW YORK	<u>2.5%</u>	<u>A089996 001</u>	Jan 10, 1991
<u>AT</u> WOCKHARDT	<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>	

SERTACONAZOLE NITRATE

CREAM; TOPICAL

ERTACZO

+ ORTHO JANSSEN	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
<u>AA</u> RANBAXY	<u>EQ 20MG BASE/ML</u>	<u>A078053 001</u>	Feb 05, 2007
<u>ZOLOFT</u>			
<u>AA</u> + PFIZER	<u>EQ 20MG BASE/ML</u>	<u>N020990 001</u>	Dec 07, 1999

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

<u>AB</u> APOTEX INC	<u>EQ 25MG BASE</u>	<u>A076882 001</u>	Feb 06, 2007
	<u>EQ 50MG BASE</u>	<u>A076882 002</u>	Feb 06, 2007
	<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>EQ 50MG BASE</u>	<u>A077206 002</u>	Feb 06, 2007
	<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>EQ 50MG BASE</u>	<u>A078677 002</u>	Mar 04, 2009
	<u>EQ 100MG BASE</u>	<u>A078677 003</u>	Mar 04, 2009
<u>AB</u> DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A076442 001</u>	Apr 30, 2007
	<u>EQ 50MG BASE</u>	<u>A076442 002</u>	Apr 30, 2007
	<u>EQ 100MG BASE</u>	<u>A076442 003</u>	Apr 30, 2007
<u>AB</u> HIKMA PHARMS	<u>EQ 25MG BASE</u>	<u>A077864 001</u>	Aug 10, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 370 (of 424)

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

<u>AB</u>	HIKMA PHARMS	<u>EQ 50MG BASE</u>	<u>A077864</u> <u>002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077864</u> <u>003</u>	Aug 10, 2009
<u>AB</u>	INVAGEN PHARMS	<u>EQ 25MG BASE</u>	<u>A077397</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077397</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077397</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	LUPIN	<u>EQ 25MG BASE</u>	<u>A077670</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077670</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077670</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	MATRIX LABS LTD	<u>EQ 25MG BASE</u>	<u>A078626</u> <u>001</u>	Jan 31, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078626</u> <u>002</u>	Jan 31, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078626</u> <u>003</u>	Jan 31, 2008
<u>AB</u>	MYLAN	<u>EQ 25MG BASE</u>	<u>A076540</u> <u>001</u>	Mar 20, 2007
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A076671</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076540</u> <u>002</u>	Mar 20, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076671</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076540</u> <u>003</u>	Mar 20, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076671</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	RANBAXY	<u>EQ 25MG BASE</u>	<u>A077977</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077977</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077977</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	SUN PHARM INDs (IN)	<u>EQ 25MG BASE</u>	<u>A078108</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078108</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078108</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076465</u> <u>001</u>	Aug 11, 2006
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076465</u> <u>002</u>	Aug 11, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076465</u> <u>003</u>	Aug 11, 2006
<u>AB</u>	TORRENT PHARMS	<u>EQ 25MG BASE</u>	<u>A077765</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077765</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077765</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	WATSON LABS	<u>EQ 25MG BASE</u>	<u>A077663</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077663</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077663</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	WOCKHARDT	<u>EQ 25MG BASE</u>	<u>A078403</u> <u>001</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078403</u> <u>002</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078403</u> <u>003</u>	Jan 08, 2008
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A077106</u> <u>001</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077106</u> <u>002</u>	Feb 06, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077106</u> <u>003</u>	Feb 06, 2007
<u>AB</u>	<u>ZOLOFT</u>			
<u>AB</u>	PFIZER	<u>EQ 25MG BASE</u>	<u>N019839</u> <u>005</u>	Mar 06, 1996
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N019839</u> <u>001</u>	Dec 30, 1991
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N019839</u> <u>002</u>	Dec 30, 1991
<u>SERTRALINE HYDROCHLORIDE</u>				
	RANBAXY	EQ 150MG BASE	A077977	004
		EQ 200MG BASE	A077977	005

SEVELAMER CARBONATE

FOR SUSPENSION; ORAL

RENVELA

GENZYME

800MG/PACKET

N022318 001 Aug 12, 2009

+

2.4GM/PACKET

N022318 002 Feb 18, 2009

TABLET; ORAL

RENVELA

+ GENZYME

800MG

N022127 001 Oct 19, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 371 (of 424)

SEVELAMER HYDROCHLORIDE

TABLET; ORAL				
RENAGEL				
GENZYME	400MG		N021179 001	Jul 12, 2000
+	800MG		N021179 002	Jul 12, 2000

SEVOFLURANE

LIQUID; INHALATION				
<u>SEVOFLURANE</u>				
<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> <u>SOJOURN</u>				
<u>AN</u> PIRAMAL CRITICAL	<u>100%</u>		<u>A077867 001</u>	May 02, 2007
<u>AN</u> <u>ULTANE</u>				
<u>AN</u> + ABBOTT	<u>100%</u>		<u>N020478 001</u>	Jun 07, 1995

SILDENAFIL CITRATE

SOLUTION; INTRAVENOUS				
REVATIO				
+ PFIZER	EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)		N022473 001	Nov 18, 2009
TABLET; ORAL				
REVATIO				
+ PFIZER	EQ 20MG BASE		N021845 001	Jun 03, 2005
VIAGRA				
PFIZER IRELAND	EQ 25MG BASE		N020895 001	Mar 27, 1998
	EQ 50MG BASE		N020895 002	Mar 27, 1998
+	EQ 100MG BASE		N020895 003	Mar 27, 1998

SILODOSIN

CAPSULE; ORAL				
RAPAFLO				
WATSON LABS	4MG		N022206 001	Oct 08, 2008
+	8MG		N022206 002	Oct 08, 2008

SILVER SULFADIAZINE

CREAM; TOPICAL				
<u>SILVADENE</u>				
<u>AB</u> + KING PHARMS	<u>1%</u>		<u>N017381 001</u>	
<u>SSD</u>				
<u>AB</u> DR REDDYS LA	<u>1%</u>		<u>N018578 001</u>	Feb 25, 1982
<u>THERMAZENE</u>				
<u>AB</u> COVIDIEN	<u>1%</u>		<u>N018810 001</u>	Dec 23, 1985
SSD AF				
BX DR REDDYS LA	1%		N018578 003	Jul 11, 1990

SIMVASTATIN

TABLET; ORAL				
<u>SIMVASTATIN</u>				
<u>AB</u> ACCORD HLTHCARE	<u>10MG</u>		<u>A078155 002</u>	Feb 26, 2008
	<u>20MG</u>		<u>A078155 003</u>	Feb 26, 2008
	<u>40MG</u>		<u>A078155 004</u>	Feb 26, 2008
	<u>80MG</u>		<u>A078155 001</u>	Feb 26, 2008
<u>AB</u> AUROBINDO PHARMA	<u>5MG</u>		<u>A077691 001</u>	Dec 20, 2006
	<u>10MG</u>		<u>A077691 002</u>	Dec 20, 2006
	<u>20MG</u>		<u>A077691 003</u>	Dec 20, 2006
	<u>40MG</u>		<u>A077691 004</u>	Dec 20, 2006
	<u>80MG</u>		<u>A077691 005</u>	Dec 20, 2006
<u>AB</u> BLU CARIBE	<u>5MG</u>		<u>A078034 001</u>	Dec 20, 2006
	<u>10MG</u>		<u>A078034 002</u>	Dec 20, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 372 (of 424)

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

AB	BLU CARIBE	<u>20MG</u>	<u>A078034</u> <u>003</u>	Dec 20, 2006
AB		<u>40MG</u>	<u>A078034</u> <u>004</u>	Dec 20, 2006
AB		<u>80MG</u>	<u>A078034</u> <u>005</u>	Dec 20, 2006
AB	DR REDDYS LABS INC	<u>5MG</u>	<u>A077752</u> <u>005</u>	Jan 23, 2008
AB		<u>10MG</u>	<u>A077752</u> <u>001</u>	Dec 20, 2006
AB		<u>20MG</u>	<u>A077752</u> <u>002</u>	Dec 20, 2006
AB		<u>40MG</u>	<u>A077752</u> <u>003</u>	Dec 20, 2006
AB		<u>80MG</u>	<u>A077752</u> <u>004</u>	Dec 20, 2006
AB	IVAX SUB TEVA PHARMS	<u>5MG</u>	<u>A076052</u> <u>001</u>	Jun 23, 2006
AB		<u>10MG</u>	<u>A076052</u> <u>002</u>	Jun 23, 2006
AB		<u>20MG</u>	<u>A076052</u> <u>003</u>	Jun 23, 2006
AB		<u>40MG</u>	<u>A076052</u> <u>004</u>	Jun 23, 2006
AB		<u>80MG</u>	<u>A076052</u> <u>005</u>	Dec 20, 2006
AB	LUPIN	<u>5MG</u>	<u>A078103</u> <u>005</u>	Apr 14, 2009
AB		<u>10MG</u>	<u>A078103</u> <u>001</u>	May 11, 2007
AB		<u>20MG</u>	<u>A078103</u> <u>002</u>	May 11, 2007
AB		<u>40MG</u>	<u>A078103</u> <u>003</u>	May 11, 2007
AB		<u>80MG</u>	<u>A078103</u> <u>004</u>	May 11, 2007
AB	MATRIX LABS LTD	<u>5MG</u>	<u>A090868</u> <u>001</u>	Jun 08, 2010
AB		<u>10MG</u>	<u>A090868</u> <u>002</u>	Jun 08, 2010
AB		<u>20MG</u>	<u>A090868</u> <u>003</u>	Jun 08, 2010
AB		<u>40MG</u>	<u>A090868</u> <u>004</u>	Jun 08, 2010
AB		<u>80MG</u>	<u>A090868</u> <u>005</u>	Jun 08, 2010
AB	MICRO LABS LTD	<u>5MG</u>	<u>A090383</u> <u>001</u>	Sep 16, 2011
AB		<u>10MG</u>	<u>A090383</u> <u>002</u>	Sep 16, 2011
AB		<u>20MG</u>	<u>A090383</u> <u>003</u>	Sep 16, 2011
AB		<u>40MG</u>	<u>A090383</u> <u>004</u>	Sep 16, 2011
AB		<u>80MG</u>	<u>A090383</u> <u>005</u>	Sep 16, 2011
AB	RANBAXY	<u>5MG</u>	<u>A076285</u> <u>001</u>	Dec 20, 2006
AB		<u>10MG</u>	<u>A076285</u> <u>002</u>	Dec 20, 2006
AB		<u>20MG</u>	<u>A076285</u> <u>003</u>	Dec 20, 2006
AB		<u>40MG</u>	<u>A076285</u> <u>004</u>	Dec 20, 2006
AB		<u>80MG</u>	<u>A076285</u> <u>005</u>	Jun 23, 2006
AB	WATSON LABS	<u>5MG</u>	<u>A076685</u> <u>001</u>	Dec 20, 2006
AB		<u>10MG</u>	<u>A076685</u> <u>002</u>	Dec 20, 2006
AB		<u>20MG</u>	<u>A076685</u> <u>003</u>	Dec 20, 2006
AB		<u>40MG</u>	<u>A076685</u> <u>004</u>	Dec 20, 2006
AB		<u>80MG</u>	<u>A076685</u> <u>005</u>	Dec 20, 2006
AB	ZYDUS PHARMS USA	<u>5MG</u>	<u>A077837</u> <u>001</u>	Dec 20, 2006
AB		<u>10MG</u>	<u>A077837</u> <u>002</u>	Dec 20, 2006
AB		<u>20MG</u>	<u>A077837</u> <u>003</u>	Dec 20, 2006
AB		<u>40MG</u>	<u>A077837</u> <u>004</u>	Dec 20, 2006
AB		<u>80MG</u>	<u>A077837</u> <u>005</u>	Dec 20, 2006
AB	<u>ZOCOR</u>			
AB	MERCK	<u>5MG</u>	<u>N019766</u> <u>001</u>	Dec 23, 1991
AB		<u>10MG</u>	<u>N019766</u> <u>002</u>	Dec 23, 1991
AB		<u>20MG</u>	<u>N019766</u> <u>003</u>	Dec 23, 1991
AB		<u>40MG</u>	<u>N019766</u> <u>004</u>	Dec 23, 1991
AB	+	<u>80MG</u>	<u>N019766</u> <u>005</u>	Jul 10, 1998

SIMVASTATIN; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JUVISYNC

MERCK SHARP DOHME

10MG;EQ 100MG BASE
20MG;EQ 100MG BASE
40MG;EQ 100MG BASE

+

N202343 001 Oct 07, 2011
N202343 002 Oct 07, 2011
N202343 003 Oct 07, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 373 (of 424)

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 + WYETH PHARMS INC	1MG/ML	N021083 001	Sep 15, 1999
TABLET; ORAL RAPAMUNE WYETH PHARMS INC	0.5MG 1MG + 2MG	N021110 004 N021110 001 N021110 002	Jan 25, 2010 Aug 25, 2000 Aug 22, 2002

SITAGLIPTIN PHOSPHATE

TABLET; ORAL JANUVIA MERCK CO INC	EQ 25MG BASE EQ 50MG BASE + EQ 100MG BASE	N021995 001 N021995 002 N021995 003	Oct 16, 2006 Oct 16, 2006 Oct 16, 2006
---	---	---	--

SODIUM ACETATE ANHYDROUS

INJECTABLE; INJECTION
 SODIUM ACETATE IN PLASTIC CONTAINER
 + HOSPIRA 2MEQ/ML N018893 001 May 04, 1983

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; IV (INFUSION)
 + UCYCLYD 10%;10% (5GM/50ML;5GM/50ML) N020645 001 Feb 17, 2005

SODIUM BICARBONATE

INJECTABLE; INJECTION
 SODIUM BICARBONATE
 + HOSPIRA 0.9MEQ/ML
 + 1MEQ/ML A077394 001 Nov 09, 2005
 A077394 002 Nov 09, 2005

SODIUM CHLORIDE

INJECTABLE; INJECTION <u>BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> <u>AP APP PHARMS 9MG/ML</u> <u>AP + HOSPIRA 9MG/ML</u>	<u>A088911 001</u> Feb 07, 1985 <u>N018800 001</u> Oct 29, 1982
<u>SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u> <u>AP B BRAUN 450MG/100ML</u> <u>AP BAXTER HLTHCARE 450MG/100ML</u> <u>AP HOSPIRA 450MG/100ML</u> <u>AP 450MG/100ML</u>	<u>N019635 001</u> Mar 09, 1988 <u>N018016 001</u> <u>N018090 001</u> <u>N019759 001</u> Jun 08, 1988
<u>SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u> <u>AP + APP PHARMS 9MG/ML</u> <u>AP + B BRAUN 900MG/100ML</u> <u>AP + 900MG/100ML</u> <u>AP + BAXTER HLTHCARE 9MG/ML</u> <u>AP 9MG/ML</u>	<u>A088912 001</u> Jan 10, 1985 <u>N017464 001</u> <u>N019635 002</u> Mar 09, 1988 <u>N016677 004</u> Oct 30, 1985 <u>N020178 002</u> Dec 07, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 374 (of 424)

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u> + BAXTER HLTHCARE	<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 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>N016366</u> <u>001</u>	
<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> + TARO PHARMS IRELAND	<u>9MG/ML</u>	<u>A077407</u> <u>001</u>	Aug 11, 2006
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ MALLINCKRODT	45MG/50ML (9MG/ML)	N021569 001	Jul 27, 2006
	112.5MG/125ML (9MG/ML)	N021569 002	Jul 27, 2006
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	3GM/100ML	N019022 001	Nov 01, 1983
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML	N019022 002	Nov 01, 1983
SODIUM CHLORIDE IN PLASTIC CONTAINER			
HOSPIRA	2.5MEQ/ML	N018897 001	Jul 20, 1984

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER	
BAXTER HLTHCARE	900MG/100ML

N019319 002 May 17, 1985

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AT</u> BAXTER HLTHCARE	<u>450MG/100ML</u>	<u>N017864</u> <u>001</u>
<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> HOSPIRA	<u>900MG/100ML</u>	<u>N017514</u> <u>001</u>
<u>AT</u>	<u>900MG/100ML</u>	<u>N018314</u> <u>001</u>

SODIUM CHROMATE CR-51

INJECTABLE; INJECTION

CHROMITOPE SODIUM

BRACCO

200uCi/ML

N013993 001

SODIUM FERRIC GLUCONATE COMPLEX

INJECTABLE; INJECTION

FERRLECIT

<u>AB</u> + SANOFI AVENTIS US	<u>62.5MG/5ML</u>	<u>N020955</u> <u>001</u>	Feb 18, 1999
<u>SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE</u>			
<u>AB</u> GENERAMEDIX	<u>62.5MG/5ML</u>	<u>A078215</u> <u>001</u>	Mar 31, 2011

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

<u>AA</u> + GE HEALTHCARE	<u>100uCi</u>	<u>N017630</u> <u>001</u>	
<u>AA</u> MALLINCKRODT	<u>100uCi</u>	<u>A071909</u> <u>001</u>	Feb 28, 1989
<u>AA</u>	<u>200uCi</u>	<u>A071910</u> <u>001</u>	Feb 28, 1989
<u>AA</u> + SYNCOR PHARMS	<u>100uCi</u>	<u>N018671</u> <u>001</u>	May 27, 1982
<u>AA</u> +	<u>200uCi</u>	<u>N018671</u> <u>002</u>	May 27, 1982

SOLUTION; ORAL

SODIUM IODIDE I 123

+ GE HEALTHCARE

2mCi/ML

N017630 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 375 (of 424)

SODIUM IODIDE I-131

CAPSULE; ORAL			
SODIUM IODIDE I 131			
+ MALLINCKRODT	0.8-100mCi	N016517	001
SODIUM IODIDE I-131			
DRAXIMAGE	9-100mCi	N021305	006 May 19, 2005
	2-200mCi	N021305	004 Nov 18, 2004
SOLUTION; ORAL			
HICON			
+ DRAXIMAGE	1-1000mCi/ML	N021305	005 Apr 04, 2006
+	1-500mCi/0.5ML	N021305	003 Jan 24, 2003
+	1-250mCi/0.25ML	N021305	002 Jan 24, 2003
SODIUM IODIDE I 131			
+ MALLINCKRODT	3.5-150mCi/VIAL	N016515	001

SODIUM LACTATE

INJECTABLE; INJECTION			
SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER			
AP BAXTER HLTHCARE	1.87GM/100ML	N016692	001
SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER			
AP B BRAUN	1.87GM/100ML	N020004	001 Apr 21, 1992
SODIUM LACTATE IN PLASTIC CONTAINER			
+ HOSPIRA	5MEQ/ML	N018947	001 Sep 05, 1984

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

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION			
NITROPRESS			
+ HOSPIRA	25MG/ML	A071961	001 Aug 01, 1988

SODIUM OXYBATE

SOLUTION; ORAL			
XYREM			
+ JAZZ	500MG/ML	N021196	001 Jul 17, 2002

SODIUM PHENYLBUTYRATE

POWDER; ORAL			
BUPHENYL			
+ MEDICIS	3GM/TEASPOONFUL	N020573	001 Apr 30, 1996
TABLET; ORAL			
BUPHENYL			
AB + MEDICIS	500MG	N020572	001 May 13, 1996
SODIUM PHENYLBUTYRATE			
AB AMPOLGEN	500MG	A090910	001 Nov 18, 2011

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			
AB NOVEL LABS INC	0.398GM;1.102GM	A079247	001 Dec 30, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 376 (of 424)

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

OSMOPREP

AB + 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

AA KVK TECH 454GM/BOT A040905 001 Mar 30, 2009

KAYEXALATE

AA + SANOFI AVENTIS US 453.6GM/BOT N011287 001

KIONEX

AA PADDOCK LLC 454GM/BOT A040029 001 Feb 06, 1998

SODIUM POLYSTYRENE SULFONATE

AA CAROLINA MEDCL 454GM/BOT A089910 001 Jan 19, 1989

AA CEDAR PHARMS 453.6GM/BOT A090313 001 Dec 21, 2011

AA CITRUSPHRMA 454GM/BOT A040909 001 Dec 03, 2008

SUSPENSION; ORAL, RECTAL

KIONEX

AA PADDOCK LLC 15GM/60ML A040028 001 Sep 17, 2007

SODIUM POLYSTYRENE SULFONATE

AA PADDOCK LLC 15GM/60ML A090590 001 May 13, 2011

AA ROXANE 15GM/60ML A089049 001 Nov 17, 1986

SPS

AA + CAROLINA MEDCL 15GM/60ML A087859 001 Dec 08, 1982

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL
BIONICHE PHARMA 20MG/2ML (10MG/ML) A040541 001 Nov 12, 2004

+ 60MG/2ML (30MG/ML) A040541 002 Nov 12, 2004

SOLIFENACIN SUCCINATE

TABLET; ORAL

VESICARE

ASTELLAS 5MG N021518 001 Nov 19, 2004

+ 10MG N021518 002 Nov 19, 2004

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

GENOTROPIN
BX + PHARMACIA AND UPJOHN 5.8MG/VIAL N020280 006 Aug 24, 1995

GENOTROPIN PRESERVATIVE FREE
BX PHARMACIA AND UPJOHN 1.5MG/VIAL N020280 004 Aug 24, 1995

HUMATROPE
BX + LILLY 5MG/VIAL N019640 004 Mar 08, 1987

BX 6MG/VIAL N019640 005 Feb 04, 1999

NORDITROPIN
BX NOVO NORDISK INC 5MG/1.5ML N021148 001 Jun 20, 2000

BX 10MG/1.5ML N021148 002 Jun 20, 2000

NORDITROPIN FLEXPRO
BX NOVO NORDISK INC 5MG/1.5ML N021148 008 Mar 01, 2010

BX 10MG/1.5ML N021148 009 Mar 01, 2010

NORDITROPIN NORDIFLEX
BX NOVO NORDISK INC 5MG/1.5ML N021148 004 Oct 01, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 377 (of 424)

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION NORDITROPIN NORDIFLEX				
BX	NOVO NORDISK INC	10MG/1.5ML	N021148 005	Oct 01, 2004
	NUTROPIN			
BX	GENENTECH	5MG/VIAL	N020168 001	Nov 17, 1993
	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	4MG/VIAL	N020604 003	Jul 25, 1997
BX		5MG/VIAL	N020604 002	Aug 23, 1996
BX		6MG/VIAL	N020604 001	Aug 23, 1996
	TEV-TROPIN			
BX	+ FERRING	5MG/VIAL	N019774 002	Jan 04, 2002
	VALTROPIN			
BX	LG LIFE	5MG/VIAL	N021905 001	Apr 19, 2007
	GENOTROPIN			
+ PHARMACIA AND UPJOHN		13.8MG/VIAL	N020280 007	Oct 23, 1996
GENOTROPIN PRESERVATIVE		FREE		
PHARMACIA AND UPJOHN		0.2MG/VIAL	N020280 001	Jan 27, 1998
		0.4MG/VIAL	N020280 002	Jan 27, 1998
		0.6MG/VIAL	N020280 003	Jan 27, 1998
		0.8MG/VIAL	N020280 005	Jan 27, 1998
		1MG/VIAL	N020280 008	Jan 27, 1998
		1.2MG/VIAL	N020280 009	Jan 27, 1998
		1.4MG/VIAL	N020280 010	Jan 27, 1998
		1.6MG/VIAL	N020280 011	Jan 27, 1998
		1.8MG/VIAL	N020280 012	Jan 27, 1998
+		2MG/VIAL	N020280 013	Jan 27, 1998
HUMATROPE				
+ LILLY		12MG/VIAL	N019640 006	Feb 04, 1999
+		24MG/VIAL	N019640 007	Feb 04, 1999
NORDITROPIN				
+ NOVO NORDISK INC		15MG/1.5ML	N021148 003	Jun 20, 2000
NORDITROPIN FLEXPRO				
NOVO NORDISK INC		15MG/1.5ML	N021148 010	Mar 01, 2010
NORDITROPIN NORDIFLEX				
NOVO NORDISK INC		15MG/1.5ML	N021148 006	Oct 01, 2004
		30MG/3ML	N021148 007	Mar 10, 2009
NUTROPIN				
+ GENENTECH		10MG/VIAL	N020168 002	Nov 17, 1993
NUTROPIN AQ				
+ GENENTECH		5MG/2ML (2.5MG/ML)	N020522 003	Jan 03, 2008
+		10MG/2ML (5MG/ML)	N020522 001	Dec 29, 1995
+		20MG/2ML (10MG/ML)	N020522 004	Jan 03, 2008
NUTROPIN AQ PEN				
+ GENENTECH		10MG/2ML (5MG/ML)	N020522 002	Apr 22, 2002
SAIZEN				
+ EMD SERONO		8.8MG/VIAL	N019764 003	Aug 29, 2000
ZORBTIVE				
+ EMD SERONO		8.8MG/VIAL	N021597 004	Dec 01, 2003

SORAFENIB TOSYLATE

TABLET; ORAL NEXAVAR			
+ BAYER HLTHCARE	EQ 200MG BASE	N021923 001	Dec 20, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 378 (of 424)

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

+ ACADEMIC PHARMS 150MG/10ML (15MG/ML) N022306 001 Jul 02, 2009

TABLET; ORAL

BETAPACE

<u>AB1</u>	BAYER HLTHCARE	<u>80MG</u>	<u>N019865</u> <u>001</u>	Oct 30, 1992
<u>AB1</u>		<u>120MG</u>	<u>N019865</u> <u>005</u>	Apr 20, 1994
<u>AB1</u> +		<u>160MG</u>	<u>N019865</u> <u>002</u>	Oct 30, 1992
<u>AB1</u>		<u>240MG</u>	<u>N019865</u> <u>003</u>	Oct 30, 1992

SORINE

<u>AB1</u>	UPSHER SMITH	<u>80MG</u>	<u>A075500</u> <u>001</u>	Apr 27, 2001
<u>AB1</u>		<u>120MG</u>	<u>A075500</u> <u>004</u>	Apr 27, 2001
<u>AB1</u>		<u>160MG</u>	<u>A075500</u> <u>002</u>	Apr 27, 2001
<u>AB1</u>		<u>240MG</u>	<u>A075500</u> <u>003</u>	Apr 27, 2001

SOTALOL HYDROCHLORIDE

<u>AB1</u>	APOTEX INC	<u>80MG</u>	<u>A076140</u> <u>001</u>	Sep 26, 2002
<u>AB1</u>		<u>120MG</u>	<u>A076140</u> <u>002</u>	Sep 26, 2002
<u>AB1</u>		<u>160MG</u>	<u>A076140</u> <u>003</u>	Sep 26, 2002
<u>AB1</u>		<u>240MG</u>	<u>A076140</u> <u>004</u>	Sep 26, 2002
<u>AB1</u>	IMPAK PHARMS	<u>80MG</u>	<u>A075663</u> <u>001</u>	Nov 07, 2000
<u>AB1</u>		<u>120MG</u>	<u>A075663</u> <u>002</u>	Nov 07, 2000
<u>AB1</u>		<u>160MG</u>	<u>A075663</u> <u>003</u>	Nov 07, 2000
<u>AB1</u>		<u>240MG</u>	<u>A075663</u> <u>004</u>	Nov 07, 2000
<u>AB1</u>	MYLAN	<u>80MG</u>	<u>A075237</u> <u>001</u>	May 01, 2000
<u>AB1</u>		<u>120MG</u>	<u>A075237</u> <u>002</u>	May 01, 2000
<u>AB1</u>		<u>160MG</u>	<u>A075237</u> <u>003</u>	May 01, 2000
<u>AB1</u>		<u>240MG</u>	<u>A075237</u> <u>004</u>	May 01, 2000
<u>AB1</u>	SANDOZ	<u>80MG</u>	<u>A075366</u> <u>001</u>	May 01, 2000
<u>AB1</u>		<u>120MG</u>	<u>A075366</u> <u>002</u>	May 01, 2000
<u>AB1</u>		<u>160MG</u>	<u>A075366</u> <u>003</u>	May 01, 2000
<u>AB1</u>		<u>240MG</u>	<u>A075366</u> <u>004</u>	May 01, 2000
<u>AB1</u>	TEVA	<u>80MG</u>	<u>A075429</u> <u>001</u>	May 01, 2000
<u>AB1</u>		<u>120MG</u>	<u>A075429</u> <u>002</u>	May 01, 2000
<u>AB1</u>		<u>160MG</u>	<u>A075429</u> <u>003</u>	May 01, 2000
<u>AB1</u>		<u>240MG</u>	<u>A075429</u> <u>004</u>	May 01, 2000
<u>AB1</u>	VINTAGE PHARMS	<u>80MG</u>	<u>A075563</u> <u>001</u>	Nov 07, 2003
<u>AB1</u>		<u>120MG</u>	<u>A075563</u> <u>002</u>	Nov 07, 2003
<u>AB1</u>		<u>160MG</u>	<u>A075563</u> <u>003</u>	Nov 07, 2003
<u>AB1</u>		<u>240MG</u>	<u>A075563</u> <u>004</u>	Nov 07, 2003

BETAPACE AF

<u>AB2</u>	BAYER HLTHCARE	<u>80MG</u>	<u>N021151</u> <u>001</u>	Feb 22, 2000
<u>AB2</u>		<u>120MG</u>	<u>N021151</u> <u>002</u>	Feb 22, 2000
<u>AB2</u> +		<u>160MG</u>	<u>N021151</u> <u>003</u>	Feb 22, 2000

SOTALOL HYDROCHLORIDE

<u>AB2</u>	AMNEAL PHARM	<u>80MG</u>	<u>A077070</u> <u>001</u>	Nov 04, 2005
<u>AB2</u>		<u>120MG</u>	<u>A077070</u> <u>002</u>	Nov 04, 2005
<u>AB2</u>		<u>160MG</u>	<u>A077070</u> <u>003</u>	Nov 04, 2005
<u>AB2</u>	APOTEX	<u>80MG</u>	<u>A076214</u> <u>001</u>	Aug 27, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 379 (of 424)

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

<u>AB2</u>	APOTEX	<u>120MG</u>	<u>A076214</u> <u>002</u>	Aug 27, 2003
<u>AB2</u>		<u>160MG</u>	<u>A076214</u> <u>003</u>	Aug 27, 2003
<u>AB2</u>	MYLAN	<u>80MG</u>	<u>A077616</u> <u>001</u>	Feb 07, 2007
<u>AB2</u>		<u>120MG</u>	<u>A077616</u> <u>002</u>	Feb 07, 2007
<u>AB2</u>		<u>160MG</u>	<u>A077616</u> <u>003</u>	Feb 07, 2007
<u>AB2</u>	TEVA	<u>80MG</u>	<u>A076883</u> <u>001</u>	Jul 26, 2004
<u>AB2</u>		<u>120MG</u>	<u>A076883</u> <u>002</u>	Jul 26, 2004
<u>AB2</u>		<u>160MG</u>	<u>A076883</u> <u>003</u>	Jul 26, 2004

SOYBEAN OIL

INJECTABLE; INJECTION

INTRALIPID 10%

<u>AP</u>	+ FRESENIUS	<u>10%</u>	<u>N017643</u> <u>001</u>	
<u>AP</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>	Aug 07, 1996
<u>AP</u>	<u>INTRALIPID 30%</u>			
<u>AP</u>	+ FRESENIUS	<u>30%</u>	<u>N019942</u> <u>001</u>	Dec 30, 1993
<u>AP</u>	<u>LIPOSYN III 10%</u>			
<u>AP</u>	+ HOSPIRA	<u>10%</u>	<u>N018969</u> <u>001</u>	Sep 24, 1984
<u>AP</u>	<u>LIPOSYN III 20%</u>			
<u>AP</u>	+ HOSPIRA	<u>20%</u>	<u>N018970</u> <u>001</u>	Sep 25, 1984
<u>AP</u>	<u>LIPOSYN III 30%</u>			
<u>AP</u>	+ HOSPIRA	<u>30%</u>	<u>N020181</u> <u>001</u>	Jan 13, 1998
<u>AP</u>	<u>NUTRILIPID 10%</u>			
<u>AP</u>	+ B BRAUN	<u>10%</u>	<u>N019531</u> <u>001</u>	May 28, 1993
<u>AP</u>	<u>NUTRILIPID 20%</u>			
<u>AP</u>	+ B BRAUN	<u>20%</u>	<u>N019531</u> <u>002</u>	May 28, 1993

SPINOSAD

SUSPENSION; TOPICAL

NATROBA

+ PARAPRO PHARMS 0.9%

N022408 001 Jan 18, 2011

SPIRONOLACTONE

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
<u>AB</u>	<u>SPIRONOLACTONE</u>			
<u>AB</u>	ACTAVIS ELIZABETH	<u>25MG</u>	<u>A040353</u> <u>003</u>	Mar 15, 2006
<u>AB</u>		<u>50MG</u>	<u>A040353</u> <u>001</u>	Jul 29, 1999
<u>AB</u>		<u>100MG</u>	<u>A040353</u> <u>002</u>	Jul 29, 1999
<u>AB</u>	AMNEAL PHARMS	<u>25MG</u>	<u>A091426</u> <u>001</u>	Jun 08, 2010
<u>AB</u>		<u>50MG</u>	<u>A091426</u> <u>002</u>	Jun 08, 2010
<u>AB</u>		<u>100MG</u>	<u>A091426</u> <u>003</u>	Jun 08, 2010
<u>AB</u>	MUTUAL PHARM	<u>25MG</u>	<u>A089424</u> <u>001</u>	Jul 23, 1986
<u>AB</u>		<u>50MG</u>	<u>A089424</u> <u>002</u>	Aug 11, 1999
<u>AB</u>		<u>100MG</u>	<u>A089424</u> <u>003</u>	Aug 11, 1999
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A040424</u> <u>001</u>	Aug 20, 2001
<u>AB</u>		<u>50MG</u>	<u>A040424</u> <u>002</u>	Aug 20, 2001
<u>AB</u>		<u>100MG</u>	<u>A040424</u> <u>003</u>	Aug 20, 2001
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A086809</u> <u>001</u>	
<u>AB</u>	VINTAGE	<u>25MG</u>	<u>A040750</u> <u>001</u>	Aug 29, 2006
<u>AB</u>		<u>50MG</u>	<u>A040750</u> <u>002</u>	Aug 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A040750</u> <u>003</u>	Aug 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 380 (of 424)

STAVUDINE

CAPSULE; ORAL

STAVUDINE

<u>AB</u>	AUROBINDO PHARMA	<u>15MG</u>	<u>A077672</u> <u>003</u>	Dec 29, 2008
<u>AB</u>		<u>20MG</u>	<u>A077672</u> <u>004</u>	Dec 29, 2008
<u>AB</u>		<u>30MG</u>	<u>A077672</u> <u>001</u>	Dec 29, 2008
<u>AB</u>		<u>40MG</u>	<u>A077672</u> <u>002</u>	Dec 29, 2008
<u>AB</u>	HETERO DRUGS	<u>15MG</u>	<u>A078957</u> <u>001</u>	Dec 29, 2008
<u>AB</u>		<u>20MG</u>	<u>A078957</u> <u>002</u>	Dec 29, 2008
<u>AB</u>		<u>30MG</u>	<u>A078957</u> <u>003</u>	Dec 29, 2008
<u>AB</u>		<u>40MG</u>	<u>A078957</u> <u>004</u>	Dec 29, 2008
<u>AB</u>	MATRIX LABS LTD	<u>30MG</u>	<u>A078775</u> <u>001</u>	Jan 05, 2009
<u>AB</u>		<u>40MG</u>	<u>A078775</u> <u>002</u>	Jan 05, 2009
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A079069</u> <u>001</u>	Dec 29, 2008
<u>AB</u>		<u>20MG</u>	<u>A079069</u> <u>002</u>	Dec 29, 2008
<u>AB</u>		<u>30MG</u>	<u>A079069</u> <u>003</u>	Dec 29, 2008
<u>AB</u>		<u>40MG</u>	<u>A079069</u> <u>004</u>	Dec 29, 2008
	<u>ZERIT</u>			
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>15MG</u>	<u>N020412</u> <u>002</u>	Jun 24, 1994
<u>AB</u>		<u>20MG</u>	<u>N020412</u> <u>003</u>	Jun 24, 1994
<u>AB</u>		<u>30MG</u>	<u>N020412</u> <u>004</u>	Jun 24, 1994
<u>AB</u> +		<u>40MG</u>	<u>N020412</u> <u>005</u>	Jun 24, 1994

FOR SOLUTION; ORAL

STAVUDINE

<u>AA</u>	AUROBINDO PHARMA	<u>1MG/ML</u>	<u>A077774</u> <u>001</u>	Dec 29, 2008
<u>AA</u>	CIPLA LTD	<u>1MG/ML</u>	<u>A078030</u> <u>001</u>	Mar 20, 2009
	<u>ZERIT</u>			
<u>AA</u> +	BRISTOL MYERS SQUIBB	<u>1MG/ML</u>	<u>N020413</u> <u>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</u> <u>001</u>	Oct 27, 1982
	<u>STERILE WATER FOR INJECTION IN PLASTIC CONTAINER</u>			
<u>AP</u>	APP PHARMS	<u>100%</u>	<u>A088400</u> <u>001</u>	Jan 16, 1984
<u>AP</u> +	B BRAUN	<u>100%</u>	<u>N019633</u> <u>001</u>	Feb 29, 1988
<u>AP</u> +	BAXTER HLTHCARE	<u>100%</u>	<u>N018632</u> <u>002</u>	Apr 19, 1988
<u>AP</u> +		<u>100%</u>	<u>N018632</u> <u>001</u>	Jun 30, 1982
<u>AP</u> +	HOSPIRA	<u>100%</u>	<u>N018233</u> <u>001</u>	
<u>AP</u> +		<u>100%</u>	<u>N018801</u> <u>001</u>	Oct 27, 1982
<u>AP</u> +		<u>100%</u>	<u>N019869</u> <u>001</u>	Dec 26, 1989
<u>AP</u>	TARO PHARMS IRELAND	<u>100%</u>	<u>A077393</u> <u>001</u>	Aug 11, 2006

STERILE WATER FOR IRRIGATION

LIQUID; IRRIGATION

STERILE WATER

<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N017428</u> <u>001</u>	
	<u>STERILE WATER IN PLASTIC CONTAINER</u>			
<u>AT</u>	B BRAUN	<u>100%</u>	<u>N016734</u> <u>001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N017866</u> <u>001</u>	
<u>AT</u>	HOSPIRA	<u>100%</u>	<u>N017513</u> <u>001</u>	
<u>AT</u>		<u>100%</u>	<u>N018313</u> <u>001</u>	

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

+ X GEN PHARMS EQ 1GM BASE/VIAL

A064210 001 Jun 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 381 (of 424)

STREPTOZOCIN

INJECTABLE; INJECTION
 ZANOSAR
 + TEVA PARENTERAL 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
 + LUNDBECK INC 100MG N019998 002 Jan 30, 1991

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION
ANECTINE
AP + SANDOZ 20MG/ML N008453 002
QUELICIN
AP + HOSPIRA 20MG/ML N008845 006
QUELICIN PRESERVATIVE FREE
AP + HOSPIRA 20MG/ML N008845 001
 QUELICIN PRESERVATIVE FREE
 + HOSPIRA 100MG/ML N008845 004

SUCRALFATE

SUSPENSION; ORAL
 CARAFATE
 + APTALIS PHARMA US 1GM/10ML N019183 001 Dec 16, 1993

TABLET; ORAL
CARAFATE
AB + APTALIS PHARMA US 1GM N018333 001
SUCRALFATE
AB NOSTRUM LABS 1GM A074415 001 Jun 08, 1998
AB TEVA 1GM A070848 001 Mar 29, 1996

SUFENTANIL CITRATE

INJECTABLE; INJECTION
SUFENTA PRESERVATIVE FREE
AP + AKORN EQ 0.05MG BASE/ML N019050 001 May 04, 1984
SUFENTANIL CITRATE
AP BAXTER HLTHCARE EQ 0.05MG BASE/ML A074413 001 Dec 15, 1995
AP HOSPIRA EQ 0.05MG BASE/ML A074534 001 Dec 11, 1996

SULCONAZOLE NITRATE

CREAM; TOPICAL
 EXELDERM
 + RANBAXY 1% N018737 001 Feb 28, 1989

SOLUTION; TOPICAL
 EXELDERM
 + RANBAXY 1% N018738 001 Aug 30, 1985

SULFACETAMIDE SODIUM

LOTION; TOPICAL
KLARON
AB + SANOFI AVVENTIS US 10% N019931 001 Dec 23, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 382 (of 424)

SULFACETAMIDE SODIUM

LOTION; TOPICAL

SULFACETAMIDE SODIUM

<u>AB</u>	ALTANA	<u>10%</u>	<u>A077015</u>	<u>001</u>	Nov 17, 2006
<u>AB</u>	PERRIGO CO TENNESSEE	<u>10%</u>	<u>A078649</u>	<u>001</u>	Mar 23, 2009
<u>AB</u>	TARO	<u>10%</u>	<u>A078668</u>	<u>001</u>	May 20, 2009

OINTMENT; OPHTHALMIC

CETAMIDE

<u>AT</u>	+ ALCON	<u>10%</u>	<u>A080021</u>	<u>001</u>
-----------	---------	------------	----------------	------------

SULFACETAMIDE SODIUM

<u>AT</u>	FERA PHARMS	<u>10%</u>	<u>A080029</u>	<u>001</u>
-----------	-------------	------------	----------------	------------

SOLUTION/DROPS; OPHTHALMIC

BLEPH-10

<u>AT</u>	+ ALLERGAN	<u>10%</u>	<u>A080028</u>	<u>001</u>
-----------	------------	------------	----------------	------------

SULFACETAMIDE SODIUM

<u>AT</u>	ALCON UNIVERSAL	<u>10%</u>	<u>A089560</u>	<u>001</u>	Oct 18, 1988
<u>AT</u>	BAUSCH AND LOMB	<u>10%</u>	<u>A040066</u>	<u>001</u>	Dec 28, 1994

SULFADIAZINE

TABLET; ORAL

SULFADIAZINE

+ SANDOZ	500MG		A040091	001	Jul 29, 1994
----------	-------	--	---------	-----	--------------

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

+ TEVA PARENTERAL	80MG/ML;16MG/ML		A073303	001	Oct 31, 1991
-------------------	-----------------	--	---------	-----	--------------

SUSPENSION; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

<u>AB</u>	AUROBINDO PHARMA	<u>200MG/5ML;40MG/5ML</u>	<u>A091348</u>	<u>001</u>	Jun 08, 2010
<u>AB</u>	+ HI TECH PHARMA	<u>200MG/5ML;40MG/5ML</u>	<u>A074650</u>	<u>001</u>	Dec 29, 1997
<u>AB</u>	VINTAGE	<u>200MG/5ML;40MG/5ML</u>	<u>A077785</u>	<u>001</u>	Jan 24, 2007

SULFATRIM PEDIATRIC

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>200MG/5ML;40MG/5ML</u>	<u>N018615</u>	<u>001</u>	Jan 07, 1983
-----------	----------------------	---------------------------	----------------	------------	--------------

TABLET; ORAL

BACTRIM

<u>AB</u>	MUTUAL PHARM	<u>400MG;80MG</u>	<u>N017377</u>	<u>001</u>
-----------	--------------	-------------------	----------------	------------

BACTRIM DS

<u>AB</u>	+ MUTUAL PHARM	<u>800MG;160MG</u>	<u>N017377</u>	<u>002</u>
-----------	----------------	--------------------	----------------	------------

SEPTRA

<u>AB</u>	MONARCH PHARMS	<u>400MG;80MG</u>	<u>N017376</u>	<u>001</u>
-----------	----------------	-------------------	----------------	------------

SEPTRA DS

<u>AB</u>	MONARCH PHARMS	<u>800MG;160MG</u>	<u>N017376</u>	<u>002</u>
-----------	----------------	--------------------	----------------	------------

SULFAMETHOPRIM

<u>AB</u>	NOVEL LABS INC	<u>400MG;80MG</u>	<u>A070022</u>	<u>001</u>	Feb 15, 1985
-----------	----------------	-------------------	----------------	------------	--------------

SULFAMETHOPRIM-DS

<u>AB</u>	NOVEL LABS INC	<u>800MG;160MG</u>	<u>A070032</u>	<u>001</u>	Feb 15, 1985
-----------	----------------	--------------------	----------------	------------	--------------

SULFAMETHOXAZOLE AND TRIMETHOPRIM

<u>AB</u>	AMNEAL PHARMS NY	<u>400MG;80MG</u>	<u>A076899</u>	<u>001</u>	Jan 27, 2005
-----------	------------------	-------------------	----------------	------------	--------------

800MG;160MG

<u>AB</u>	AUROBINDO PHARMA	<u>400MG;80MG</u>	<u>A090624</u>	<u>001</u>	Feb 16, 2010
-----------	------------------	-------------------	----------------	------------	--------------

800MG;160MG

<u>AB</u>	GLENMARK GENERICS	<u>400MG;80MG</u>	<u>A090828</u>	<u>002</u>	Dec 22, 2010
-----------	-------------------	-------------------	----------------	------------	--------------

800MG;160MG

<u>AB</u>	MUTUAL PHARM	<u>800MG;160MG</u>	<u>A071017</u>	<u>001</u>	Aug 25, 1986
-----------	--------------	--------------------	----------------	------------	--------------

400MG;80MG

<u>AB</u>	VINTAGE	<u>400MG;80MG</u>	<u>A078060</u>	<u>002</u>	Jan 25, 2007
-----------	---------	-------------------	----------------	------------	--------------

800MG;160MG

<u>AB</u>	VISTA PHARMS	<u>400MG;80MG</u>	<u>A076817</u>	<u>001</u>	Jan 25, 2007
-----------	--------------	-------------------	----------------	------------	--------------

800MG;160MG

<u>AB</u>		<u>800MG;160MG</u>	<u>A076817</u>	<u>002</u>	Oct 07, 2005
-----------	--	--------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 383 (of 424)

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

<u>AB</u>	SANDOZ	<u>800MG;160MG</u>	<u>N018598 004</u>	May 19, 1982
<u>AB</u>	TEVA	<u>800MG;160MG</u>	<u>A070037 001</u>	Jun 02, 1987
<u>SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH</u>				
<u>AB</u>	PLANTEX	<u>400MG;80MG</u>	<u>A070030 001</u>	Jun 02, 1987

SULFANILAMIDE

CREAM; VAGINAL

AVC

+ AZUR PHARMA 15%

N006530 003 Jan 27, 1987

SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE

+ PHARMACIA AND UPJOHN 250MG/5ML

A086983 001

TABLET; ORAL

AZULFIDINE

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>500MG</u>	<u>N007073 001</u>	
<u>SULFASALAZINE</u>				
<u>AB</u>	VINTAGE PHARMS	<u>500MG</u>	<u>A040349 001</u>	Jan 11, 2002
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A085828 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A087197 001</u>	

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>500MG</u>	<u>N007073 002</u>	Apr 06, 1983
<u>SULFASALAZINE</u>				
<u>AB</u>	VINTAGE PHARMS	<u>500MG</u>	<u>A075339 001</u>	Jan 11, 2002

SULINDAC

TABLET; ORAL

CLINORIL

<u>AB</u>	+ MERCK	<u>200MG</u>	<u>N017911 002</u>	
<u>SULINDAC</u>				
<u>AB</u>	EPIC PHARMA	<u>150MG</u>	<u>A072710 001</u>	Mar 25, 1991
<u>AB</u>		<u>200MG</u>	<u>A072711 001</u>	Mar 25, 1991
<u>AB</u>	MUTUAL PHARM	<u>150MG</u>	<u>A072050 001</u>	Apr 17, 1991
<u>AB</u>		<u>200MG</u>	<u>A072051 001</u>	Apr 17, 1991
<u>AB</u>	MYLAN	<u>150MG</u>	<u>A073039 002</u>	Jun 22, 1993
<u>AB</u>		<u>200MG</u>	<u>A073039 001</u>	Jun 22, 1993
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A071891 001</u>	Apr 03, 1990
<u>AB</u>		<u>200MG</u>	<u>A071795 001</u>	Apr 03, 1990

SUMATRIPTAN

SPRAY; NASAL

IMITREX

+ GLAXOSMITHKLINE 5MG/SPRAY

N020626 001 Aug 26, 1997
N020626 003 Aug 26, 1997SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX STATDOSE

<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>N020080 003</u>	Dec 23, 1996
<u>SUMATRIPTAN SUCCINATE</u>				
<u>AB</u>	SUN PHARM INDs INC	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A090358 001</u>	Jun 21, 2011
<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>N020080 001</u>	Dec 28, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 384 (of 424)

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

SUMATRIPTAN SUCCINATE

<u>AP</u>	APP PHARMS	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079242 001</u>	Mar 02, 2009
<u>AP</u>		<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079240 002</u>	Sep 18, 2009
<u>AP</u>		<u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u>	<u>A079240 001</u>	Sep 18, 2009
<u>AP</u>	BEDFORD	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A079123 001</u>	Feb 06, 2009
<u>AP</u>	BEDFORD LABS	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A090310 001</u>	Aug 11, 2010
<u>AP</u>	JHP PHARMS	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A077871 001</u>	Jul 09, 2009
<u>AP</u>	PAR PHARM	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A077332 001</u>	Oct 09, 2009
<u>AP</u>	SAGENT STRIDES	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A090314 001</u>	Jun 10, 2010
<u>AP</u>		<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A090641 001</u>	Jul 28, 2010
<u>AP</u>	SANDOZ	<u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u>	<u>A078067 002</u>	Feb 06, 2009
<u>AP</u>		<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A078067 001</u>	Feb 06, 2009
<u>AP</u>	TEVA PARENTERAL	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A077907 001</u>	Feb 06, 2009
<u>AP</u>	WOCKHARDT	<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>A078593 001</u>	Feb 06, 2009
	ALSUMA			
+ MERIDIAN MEDCL		<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>N022377 001</u>	Jun 29, 2010
IMITREX STATDOSE				
+ GLAXOSMITHKLINE		<u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u>	<u>N020080 002</u>	Feb 01, 2006
SUMAVEL DOSEPRO				
+ ZOGENIX INC		<u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u>	<u>N022239 001</u>	Jul 15, 2009

TABLET; ORAL

IMITREX

<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 25MG BASE</u>	<u>N020132 002</u>	Jun 01, 1995
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N020132 003</u>	Jun 01, 1995
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N020132 001</u>	Jun 01, 1995
	<u>SUMATRIPTAN SUCCINATE</u>			
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A078327 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078327 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078327 003</u>	Aug 10, 2009
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 25MG BASE</u>	<u>A076847 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076847 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076847 003</u>	Aug 10, 2009
<u>AB</u>	MYLAN	<u>EQ 25MG BASE</u>	<u>A077163 001</u>	Nov 02, 2009
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A077744 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077163 002</u>	Nov 02, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077744 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077163 003</u>	Nov 02, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077744 003</u>	Aug 10, 2009
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 25MG BASE</u>	<u>A078284 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078284 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078284 003</u>	Aug 10, 2009
<u>AB</u>	RANBAXY	<u>EQ 25MG BASE</u>	<u>A076554 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076554 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076572 001</u>	Feb 09, 2009
<u>AB</u>	SANDOZ	<u>EQ 25MG BASE</u>	<u>A076976 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076976 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076976 003</u>	Aug 10, 2009
<u>AB</u>	SUN PHARM INDs	<u>EQ 25MG BASE</u>	<u>A078295 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078295 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078295 003</u>	Aug 10, 2009
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076840 001</u>	Feb 09, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076840 002</u>	Feb 09, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076840 003</u>	Feb 09, 2009
<u>AB</u>	WATSON LABS	<u>EQ 25MG BASE</u>	<u>A076933 001</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076933 002</u>	Aug 10, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076933 003</u>	Aug 10, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 385 (of 424)

SUNITINIB MALATE

CAPSULE; ORAL SUTENT CPPI CV	EQ 12.5MG BASE EQ 25MG BASE EQ 37.5MG BASE EQ 50MG BASE	N021938 001 N021938 002 N021938 004 N021938 003	Jan 26, 2006 Jan 26, 2006 Mar 31, 2009 Jan 26, 2006
+			

TACROLIMUS

CAPSULE; ORAL <u>PROGRAF</u> AB ASTELLAS	<u>EQ 0.5MG BASE</u>	<u>N050708 003</u>	Aug 24, 1998
AB +	<u>EQ 1MG BASE</u>	<u>N050708 001</u>	Apr 08, 1994
	<u>EQ 5MG BASE</u>	<u>N050708 002</u>	Apr 08, 1994
<u>TACROLIMUS</u>			
AB ACCORD HLTHCARE	<u>0.5MG</u>	<u>A091195 001</u>	Aug 31, 2011
AB	<u>1MG</u>	<u>A091195 002</u>	Aug 31, 2011
AB	<u>5MG</u>	<u>A091195 003</u>	Aug 31, 2011
AB DR REDDYS LABS LTD	<u>EQ 0.5MG BASE</u>	<u>A090509 001</u>	May 12, 2010
AB	<u>EQ 1MG BASE</u>	<u>A090509 002</u>	May 12, 2010
AB	<u>EQ 5MG BASE</u>	<u>A090509 003</u>	May 12, 2010
AB MYLAN	<u>EQ 0.5MG BASE</u>	<u>A090596 001</u>	Sep 17, 2010
AB	<u>EQ 1MG BASE</u>	<u>A090596 002</u>	Sep 17, 2010
AB	<u>EQ 5MG BASE</u>	<u>A090596 003</u>	Sep 17, 2010
AB SANDOZ	<u>EQ 0.5MG BASE</u>	<u>A065461 001</u>	Aug 10, 2009
AB	<u>EQ 1MG BASE</u>	<u>A065461 002</u>	Aug 10, 2009
AB	<u>EQ 5MG BASE</u>	<u>A065461 003</u>	Aug 10, 2009
AB WATSON LABS	<u>EQ 5MG BASE</u>	<u>A090402 001</u>	Jul 01, 2010

INJECTABLE; INJECTION PROGRAF + ASTELLAS	EQ 5MG BASE/ML	N050709 001	Apr 08, 1994
--	----------------	-------------	--------------

OINTMENT; TOPICAL PROTOPIC ASTELLAS	0.03%	N050777 001	Dec 08, 2000
+ LILLY	0.1%	N050777 002	Dec 08, 2000

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

TALC

AEROSOL, METERED; INTRAPLEURAL SCLEROSOL + BRYAN	400MG/SPRAY	N020587 001	Dec 24, 1997
POWDER; INTRAPLEURAL TALC + BRYAN	5GM/BOT	N021388 001	Dec 15, 2003

TAMOXIFEN CITRATE

TABLET; ORAL <u>TAMOXIFEN CITRATE</u> AB AEGIS PHARMS	<u>EQ 10MG BASE</u>	<u>A076398 001</u>	Mar 31, 2003
	<u>EQ 20MG BASE</u>	<u>A076398 002</u>	Mar 31, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 386 (of 424)

TAMOXIFEN CITRATE

TABLET; ORAL

TAMOXIFEN CITRATE

<u>AB</u>	APOTEX	<u>EQ 10MG BASE</u>	<u>A090878 001</u>	Sep 23, 2011
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090878 002</u>	Sep 23, 2011
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A074732 002</u>	Feb 20, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A074732 001</u>	Feb 20, 2003
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075797 001</u>	Feb 20, 2003
<u>AB</u>	+ TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A074858 001</u>	Feb 20, 2003
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>A070929 001</u>	Feb 20, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A070929 002</u>	Feb 20, 2003
<u>AB</u>	WATSON LABS FLORIDA	<u>EQ 10MG BASE</u>	<u>A076179 001</u>	Feb 20, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076179 002</u>	Feb 20, 2003

TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

FLOMAX

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>0.4MG</u>	<u>N020579 001</u>	Apr 15, 1997
<u>TAMSULOSIN HYDROCHLORIDE</u>				
<u>AB</u>	IMPAX LABS	<u>0.4MG</u>	<u>A090377 001</u>	Mar 02, 2010
<u>AB</u>	MYLAN	<u>0.4MG</u>	<u>A090408 001</u>	Apr 27, 2010
<u>AB</u>	SANDOZ	<u>0.4MG</u>	<u>A078015 001</u>	Apr 27, 2010
<u>AB</u>	SUN PHARM INDs LTD	<u>0.4MG</u>	<u>A090931 001</u>	Jul 15, 2010
<u>AB</u>	SYNTHON PHARMS	<u>0.4MG</u>	<u>A078801 001</u>	Apr 27, 2010
<u>AB</u>	TEVA PHARMS	<u>0.4MG</u>	<u>A077630 001</u>	Apr 27, 2010
<u>AB</u>	WOCKHARDT	<u>0.4MG</u>	<u>A078938 001</u>	Apr 27, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.4MG</u>	<u>A078225 001</u>	Apr 27, 2010

TAPENTADOL HYDROCHLORIDE

TABLET; ORAL

NUCYNTA

JANSSEN PHARMS	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

JANSSEN PHARMS	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

TAZAROTENE

CREAM; TOPICAL

AVAGE

+ ALLERGAN	0.1%	N021184 003	Sep 30, 2002
TAZORAC			
+ ALLERGAN	0.05%	N021184 001	Sep 29, 2000
+	0.1%	N021184 002	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	PHARMALUCENCE	N/A	N017776 001
----	---------------	-----	-------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 387 (of 424)

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION

TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT

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

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETECAP + BRACCO N/A N018963 001 Jan 21, 1987
TECHNETIUM TC-99M MEBROFENINAP PHARMALUCENCE N/A A078242 001 Jan 29, 2008TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION

DRAXIMAGE MDP-25

+ DRAXIMAGE N/A N018035 002 Feb 27, 2004

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

CIS-MDPAP PHARMALUCENCE N/A N018124 001MDP-BRACCOAP BRACCO N/A N018107 001TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3

+ MALLINCKRODT N/A N019882 001 Jun 15, 1990

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+ MALLINCKRODT N/A N018321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

AN-DTPAAP PHARMALUCENCE N/A N017714 001DTPAAP DRAXIMAGE N/A N018511 001 Dec 29, 1989

PRESCRIPTION DRUG PRODUCT LIST

3 - 388 (of 424)

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO

<u>AP</u>	<u>PHARMALUCENCE</u>	<u>N/A</u>	<u>N019039 001</u>	Jun 30, 1987
<u>AP</u>	<u>TECHNECAN PYP KIT</u>			

MALLINCKRODTN/AN017538 001TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

MALLINCKRODT

N/A

N019981 001 Jun 10, 1991

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE

<u>AP</u>	+ LANTHEUS MEDCL	<u>N/A</u>	<u>N019785 001</u>	Dec 21, 1990
	<u>TECHNETIUM TC 99M SESTAMIBI</u>			
<u>AP</u>	CARDINAL HEALTH 414	<u>N/A</u>	<u>A078809 001</u>	Apr 28, 2009
<u>AP</u>	DRAKIMAGE	<u>N/A</u>	<u>A078806 001</u>	Apr 29, 2009
<u>AP</u>	PHARMALUCENCE	<u>10-30mCi</u>	<u>A079157 001</u>	Jul 10, 2009
	<u>TECHNETIUM TC-99 SESTAMIBI</u>			
<u>AP</u>	MALLINCKRODT	<u>N/A</u>	<u>A078098 001</u>	Sep 22, 2008

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL

TECHNELITE

+ LANTHEUS MEDCL	0.0083-2.7 CI/GENERATOR	N017771 001
ULTRA-TECHNEKOW FM		

+ MALLINCKRODT	0.25-3 CI/GENERATOR	N017243 002
----------------	---------------------	-------------

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 KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOID

+ PHARMALUCENCE	N/A	N017858 001
-----------------	-----	-------------

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW

+ GE HEALTHCARE	N/A	N020372 001	Feb 09, 1996
MYOVIEW 30ML			

+ GE HEALTHCARE	N/A	N020372 002	Jul 07, 2005
-----------------	-----	-------------	--------------

TELAPREVIR

TABLET; ORAL

INCIVEK

+ VERTEX PHARMS	375MG	N201917 001	May 23, 2011
-----------------	-------	-------------	--------------

TELAVANCIN HYDROCHLORIDE

POWDER; IV (INFUSION)

THERAVANCE INC	EQ 250MG BASE/VIAL	N022110 001	Sep 11, 2009
+	EQ 750MG BASE/VIAL	N022110 002	Sep 11, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 389 (of 424)

TELBIVUDINE

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

TELMISARTAN

TABLET; ORAL
 MICARDIS
 BOEHRINGER INGELHEIM 20MG N020850 003 Apr 04, 2000
 40MG N020850 001 Nov 10, 1998
 + 80MG N020850 002 Nov 10, 1998

TEMAZEPAM

CAPSULE; ORAL
RESTORIL
AB MALLINCKRODT INC 7.5MG N018163 003 Oct 25, 1991
AB 15MG N018163 001
AB 22.5MG N018163 004 Nov 02, 2004
AB + 30MG N018163 002
TEMAZEPAM
AB ACTAVIS ELIZABETH 15MG A071638 001 Aug 07, 1987
AB 30MG A071620 001 Aug 07, 1987
AB MUTUAL PHARM 7.5MG A078581 001 Sep 08, 2009
AB 22.5MG A071175 002 Sep 14, 2009
AB MYLAN 7.5MG A070920 002 May 21, 2010
AB 15MG A070920 004 Jul 07, 1986
AB 22.5MG A070920 003 Jun 12, 2009
AB 30MG A070920 001 Jul 10, 1986
AB NOVEL LABS INC 15MG A071456 001 Apr 21, 1987
AB 30MG A071457 001 Apr 21, 1987
AB SANDOZ 15MG A071427 001 Jan 12, 1988
AB 30MG A071428 001 Jan 12, 1988
AB WATSON LABS 15MG A071446 001 May 21, 1993
AB 30MG A071447 001 May 21, 1993

TEMOZOLOMIDE

CAPSULE; ORAL
TEMODAR
AB SCHERING 5MG N021029 001 Aug 11, 1999
AB 20MG N021029 002 Aug 11, 1999
AB 100MG N021029 003 Aug 11, 1999
AB 140MG N021029 005 Oct 19, 2006
AB 180MG N021029 006 Oct 19, 2006
AB + 250MG N021029 004 Aug 11, 1999
TEMOZOLOMIDE
AB BARR 5MG A078879 001 Mar 01, 2010
AB 20MG A078879 002 Mar 01, 2010
AB 100MG A078879 003 Mar 01, 2010
AB 140MG A078879 005 Mar 01, 2010
AB 180MG A078879 006 Mar 01, 2010
AB 250MG A078879 004 Mar 01, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 390 (of 424)

TEMOZOLOMIDE

POWDER; INTRAVENOUS
 TEMODAR
 + SCHERING 100MG/VIAL N022277 001 Feb 27, 2009

TEMSIROLIMUS

SOLUTION; INTRAVENOUS
 TORISEL
 + WYETH PHARMS INC 25MG/ML (25MG/ML) N022088 001 May 30, 2007

TENIPOSIDE

INJECTABLE; INJECTION
 VUMON
 + BRISTOL MYERS SQUIBB 10MG/ML N020119 001 Jul 14, 1992

TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL
 VIREAD
 + GILEAD 300MG N021356 001 Oct 26, 2001

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL			
<u>HYTRIN</u>			
AB ABBOTT	<u>EQ 1MG BASE</u>	N020347 001	Dec 14, 1994
AB +	<u>EQ 2MG BASE</u>	N020347 002	Dec 14, 1994
AB	<u>EQ 5MG BASE</u>	N020347 003	Dec 14, 1994
AB	<u>EQ 10MG BASE</u>	N020347 004	Dec 14, 1994
<u>TERAZOSIN HYDROCHLORIDE</u>			
AB APOTEX	<u>EQ 1MG BASE</u>	A075498 001	Apr 12, 2001
AB	<u>EQ 2MG BASE</u>	A075498 002	Apr 12, 2001
AB	<u>EQ 5MG BASE</u>	A075498 003	Apr 12, 2001
AB	<u>EQ 10MG BASE</u>	A075498 004	Apr 12, 2001
AB IVAX SUB TEVA PHARMS	<u>EQ 1MG BASE</u>	A075614 002	Jan 30, 2001
AB	<u>EQ 2MG BASE</u>	A075614 001	Jan 30, 2001
AB	<u>EQ 5MG BASE</u>	A075614 003	Jan 30, 2001
AB	<u>EQ 10MG BASE</u>	A075614 004	Jan 30, 2001
AB JUBILANT CADISTA	<u>EQ 1MG BASE</u>	A075317 001	Dec 20, 2004
AB	<u>EQ 2MG BASE</u>	A075317 002	Dec 20, 2004
AB	<u>EQ 5MG BASE</u>	A075317 003	Dec 20, 2004
AB	<u>EQ 10MG BASE</u>	A075317 004	Dec 20, 2004
AB MYLAN	<u>EQ 1MG BASE</u>	A075140 002	Feb 11, 2000
AB	<u>EQ 2MG BASE</u>	A075140 003	Feb 11, 2000
AB	<u>EQ 5MG BASE</u>	A075140 001	Feb 11, 2000
AB	<u>EQ 10MG BASE</u>	A075140 004	Feb 11, 2000
AB RANBAXY	<u>EQ 1MG BASE</u>	A076021 001	Aug 22, 2002
AB	<u>EQ 2MG BASE</u>	A076021 002	Aug 22, 2002
AB	<u>EQ 5MG BASE</u>	A076021 003	Aug 22, 2002
AB	<u>EQ 10MG BASE</u>	A076021 004	Aug 22, 2002
AB SANDOZ	<u>EQ 1MG BASE</u>	A074823 001	Mar 30, 1998
AB	<u>EQ 2MG BASE</u>	A074823 002	Mar 30, 1998
AB	<u>EQ 5MG BASE</u>	A074823 003	Mar 30, 1998
AB	<u>EQ 10MG BASE</u>	A074823 004	Mar 30, 1998

CAPSULE; ORAL			
<u>HYTRIN</u>			
AB Abbott	<u>EQ 1MG BASE</u>	N019057 001	Aug 07, 1987
+ +	<u>EQ 2MG BASE</u>	N019057 002	Aug 07, 1987
	<u>EQ 5MG BASE</u>	N019057 003	Aug 07, 1987
	<u>EQ 10MG BASE</u>	N019057 004	Aug 07, 1987

PRESCRIPTION DRUG PRODUCT LIST

3 - 391 (of 424)

TERBINAFINE HYDROCHLORIDE

GRANULE; ORAL LAMISIL			
NOVARTIS	EQ 125MG BASE/PACKET	N022071 001	Sep 28, 2007
+	EQ 187.5MG BASE/PACKET	N022071 002	Sep 28, 2007
TABLET; ORAL <u>LAMISIL</u>			
<u>AB</u> + NOVARTIS	<u>EQ 250MG BASE</u>	<u>N020539 001</u>	May 10, 1996
<u>TERBINAFINE HYDROCHLORIDE</u>			
APOTEX	<u>EQ 250MG BASE</u>	<u>A078199 001</u>	Jul 02, 2007
AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A078297 001</u>	Jul 02, 2007
BRECKENRIDGE PHARM	<u>EQ 250MG BASE</u>	<u>A077714 001</u>	Jun 04, 2010
DR REDDYS LABS INC	<u>EQ 250MG BASE</u>	<u>A076390 001</u>	Jul 02, 2007
GLENMARK GENERICS	<u>EQ 250MG BASE</u>	<u>A078157 001</u>	Jul 02, 2007
HARRIS PHARM	<u>EQ 250MG BASE</u>	<u>A077919 001</u>	Jul 02, 2007
INVAGEN PHARMS	<u>EQ 250MG BASE</u>	<u>A077533 001</u>	Jul 02, 2007
MYLAN	<u>EQ 250MG BASE</u>	<u>A077136 001</u>	Jul 02, 2007
	<u>EQ 250MG BASE</u>	<u>A077195 001</u>	Jul 02, 2007
ORCHID HLTHCARE	<u>EQ 250MG BASE</u>	<u>A078163 001</u>	Jul 02, 2007
TEVA	<u>EQ 250MG BASE</u>	<u>A076377 001</u>	Jul 02, 2007
WATSON LABS	<u>EQ 250MG BASE</u>	<u>A077137 001</u>	Jul 02, 2007
WOCKHARDT	<u>EQ 250MG BASE</u>	<u>A078229 001</u>	Jul 02, 2007

TERBUTALINE SULFATE

INJECTABLE; INJECTION <u>TERBUTALINE SULFATE</u>			
AKORN	<u>1Mg/ML</u>	<u>A078151 001</u>	Jan 07, 2008
APP PHARMS	<u>1Mg/ML</u>	<u>A076887 001</u>	May 26, 2004
+ BEDFORD	<u>1Mg/ML</u>	<u>A076770 001</u>	Apr 23, 2004
HIKMA FARMACEUTICA	<u>1Mg/ML</u>	<u>A078630 001</u>	May 20, 2009
TEVA PARENTERAL	<u>1Mg/ML</u>	<u>A076853 001</u>	Jul 20, 2004
TABLET; ORAL <u>TERBUTALINE SULFATE</u>			
IMPAX LABS	<u>2.5MG</u>	<u>A075877 001</u>	Jun 26, 2001
+	<u>5MG</u>	<u>A075877 002</u>	Jun 26, 2001
LANNETT	<u>2.5MG</u>	<u>A077152 001</u>	Mar 25, 2005
	<u>5MG</u>	<u>A077152 002</u>	Mar 25, 2005

TERCONAZOLE

CREAM; VAGINAL <u>TERAZOL 3</u>			
<u>AB</u> + JANSSEN PHARMS	<u>0.8%</u>	<u>N019964 001</u>	Feb 21, 1991
<u>TERAZOL 7</u>			
<u>AB</u> + JANSSEN PHARMS	<u>0.4%</u>	<u>N019579 001</u>	Dec 31, 1987
<u>TERCONAZOLE</u>			
ALTANA	<u>0.4%</u>	<u>A076712 001</u>	Feb 18, 2005
TARO	<u>0.4%</u>	<u>A076043 001</u>	Jan 19, 2005
	<u>0.8%</u>	<u>A075953 001</u>	Apr 06, 2004
TERCONAZOLE			
<u>BX</u> + NYCOMED US	0.8%	N021735 001	Oct 01, 2004
SUPPOSITORY; VAGINAL <u>TERAZOL 3</u>			
<u>AB</u> + JANSSEN PHARMS	<u>80MG</u>	<u>N019641 001</u>	May 24, 1988
<u>TERCONAZOLE</u>			
ALTANA	<u>80MG</u>	<u>A076850 001</u>	Jul 12, 2006
PERRIGO NEW YORK	<u>80MG</u>	<u>A077149 001</u>	Mar 17, 2006
TARO	<u>80MG</u>	<u>A077553 001</u>	Mar 09, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 392 (of 424)

TERIPARATIDE RECOMBINANT HUMAN

INJECTABLE; SUBCUTANEOUS
 FORTEO
 + LILLY 0.6MG/2.4ML (0.25MG/ML) N021318 002 Jun 25, 2008

TESAMORELIN ACETATE

POWDER; SUBCUTANEOUS
 EGRIFTA
 + THERATECHNOLOGIES EQ 1MG BASE/VIAL N022505 001 Nov 10, 2010

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL
 ANDRODERM
 + WATSON LABS 2MG/24HR N020489 003 Oct 20, 2011
 + 4MG/24HR N020489 004 Oct 20, 2011

GEL; TRANSDERMAL
 ANDROGEL
 BX + ABBOTT LABS 1% (5GM/PACKET) N021015 002 Feb 28, 2000
 TESTIM
 BX + AUXILIUM PHARMS 1% (5GM/PACKET) N021454 001 Oct 31, 2002
 ANDROGEL
 ABBOTT LABS 1% (2.5GM/PACKET) N021015 001 Feb 28, 2000

GEL, METERED; TRANSDERMAL
 ANDROGEL
 + ABBOTT LABS 1% (1.25GM/ACTIVATION) N021015 003 Sep 26, 2003
 + 1.62% (20.25MG/1.25GM ACTIVATION) N022309 001 Apr 29, 2011
 FORTESTA
 + ENDO PHARMS 10MG/0.5GM ACTIVATION N021463 001 Dec 29, 2010

PELLET; IMPLANTATION
 TESTOPEL
 + SLATE PHARMS 75MG A080911 001

SOLUTION, METERED; TRANSDERMAL
 AXIRON
 + ELI LILLY AND CO 30MG/1.5ML ACTIVATION N022504 001 Nov 23, 2010

TABLET, EXTENDED RELEASE; BUCCAL
 STRIANT
 + ACTIENT PHARMS 30MG N021543 001 Jun 19, 2003

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION
DEPO-TESTOSTERONE
AO + PHARMACIA AND UPJOHN 100MG/ML A085635 002
AO + 200MG/ML A085635 003
TESTOSTERONE CYPIONATE
AO BEDFORD 100MG/ML A090387 001 Jul 15, 2010
AO 200MG/ML A090387 002 Jul 15, 2010
AO PADDICK LLC 200MG/ML A040530 001 Jan 31, 2005
AO SANDOZ 100MG/ML A040615 001 Aug 10, 2006
AO 200MG/ML A040615 002 Aug 10, 2006
AO SYNERX PHARMA 200MG/ML A040652 001 Dec 11, 2006
AO WATSON LABS 200MG/ML A086030 001

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION
DELATESTREYL
AO + ENDO PHARM 200MG/ML N009165 003
TESTOSTERONE ENANTHATE
AO PADDICK LLC 200MG/ML A040575 001 Jun 14, 2006
AO SYNERX PHARMA 200MG/ML A040647 001 Oct 05, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 393 (of 424)

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

TESTOSTERONE ENANTHATE

<u>AO</u>	WATSON LABS	<u>200MG/ML</u>	<u>A085598 001</u>
-----------	-------------	-----------------	--------------------

TETRABENAZINE

TABLET; ORAL

XENAZINE

VALEANT INTL	12.5MG	N021894 001	Aug 15, 2008
+	25MG	N021894 002	Aug 15, 2008

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

<u>AB</u>	IMPAX LABS	<u>250MG</u>	<u>A060469 001</u>
<u>AB</u>		<u>500MG</u>	<u>A060469 003</u>
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>250MG</u>	<u>A060704 001</u>
<u>AB</u>	+	<u>500MG</u>	<u>A060704 002</u>
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A061837 001</u>
<u>AB</u>		<u>500MG</u>	<u>A061837 002</u>

TETRACYCLINE HYDROCHLORIDE

IMPAX LABS 100MG

A060469 002

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION; NASAL

TYZINE

+	NYCOMED US	0.05%	A086576 002
		0.1%	A086576 001

SPRAY; NASAL

TYZINE

+	NYCOMED US	0.1%	A086576 003
---	------------	------	-------------

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

<u>AP</u>	+	GE HEALTHCARE	<u>1mCi/ML</u>	<u>N018110 002</u>	Feb 27, 1996
<u>AP</u>	+	LANTHEUS MEDCL	<u>1mCi/ML</u>	<u>N017806 001</u>	
<u>AP</u>	+	MALLINCKRODT	<u>1mCi/ML</u>	<u>N018150 001</u>	

INJECTABLE; INTRAVENOUS

THALLOUS CHLORIDE TL 201

<u>AP</u>	+	LANTHEUS MEDCL	<u>2mCi/ML</u>	<u>N017806 002</u>	Oct 09, 1998
<u>AP</u>		MALLINCKRODT	<u>2mCi/ML</u>	<u>A077698 001</u>	Nov 09, 2006

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE; ORAL

THEO-24

BC	UCB INC	100MG	A087942 001	Aug 22, 1983
BC		200MG	A087943 001	Aug 22, 1983
BC		300MG	A087944 001	Aug 22, 1983
	THEOPHYLLINE			
BC	INWOOD LABS	100MG	A040052 001	Feb 14, 1994

PRESCRIPTION DRUG PRODUCT LIST

3 - 394 (of 424)

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE; ORAL THEOPHYLLINE			
BC	INWOOD LABS	200MG	A040052 003 Feb 14, 1994
BC	+	300MG	A040052 004 Feb 14, 1994
	THEO-24		
	UCB INC	400MG	A081034 001 Feb 28, 1992
	THEOPHYLLINE		
	INWOOD LABS	125MG	A040052 002 Feb 14, 1994
ELIXIR; ORAL ELIXOPHYLLIN			
	+	CARACO	80MG/15ML A085186 001
INJECTABLE; INJECTION <u>THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
AP	+	B BRAUN	<u>40MG/100ML</u> N019826 001 Aug 14, 1992
AP	+	B BRAUN	<u>80MG/100ML</u> N019826 002 Aug 14, 1992
AP	+	B BRAUN	<u>160MG/100ML</u> N019826 003 Aug 14, 1992
AP	+	B BRAUN	<u>320MG/100ML</u> N019826 006 Aug 14, 1992
<u>THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER</u>			
AP	+	BAXTER HLTHCARE	<u>4MG/ML</u> N018649 007 Jul 26, 1982
AP	+		<u>40MG/100ML</u> N018649 001 Jul 26, 1982
AP	+		<u>80MG/100ML</u> N018649 002 Jul 26, 1982
AP	+		<u>160MG/100ML</u> N018649 003 Jul 26, 1982
AP	+		<u>200MG/100ML</u> N018649 004 Jul 26, 1982
AP	+		<u>320MG/100ML</u> N018649 006 Nov 13, 1985
AP	+		<u>400MG/100ML</u> N018649 005 Jul 26, 1982
<u>THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
AP	+	HOSPIRA INC	<u>4MG/ML</u> N019211 007 Dec 14, 1984
AP	+		<u>40MG/100ML</u> N019211 001 Dec 14, 1984
AP	+		<u>160MG/100ML</u> N019211 003 Dec 14, 1984
AP	+		<u>320MG/100ML</u> N019211 006 Jan 20, 1988
SOLUTION; ORAL THEOPHYLLINE			
	+	SILARX	80MG/15ML A091156 001 Apr 13, 2011
TABLET; ORAL THEOLAIR			
	+	GRACEWAY	125MG A086399 001
	+		250MG A086399 002
TABLET, EXTENDED RELEASE; ORAL <u>THEOCHRON</u>			
AB		CARACO	<u>100MG</u> A088320 001 Feb 21, 1985
AB			<u>200MG</u> A088321 001 Feb 21, 1985
AB			<u>300MG</u> A087400 002 Jan 11, 1983
<u>THEOPHYLLINE</u>			
AB		ALEMBIC LTD	<u>300MG</u> A090430 001 Oct 27, 2010
AB			<u>450MG</u> A090430 002 Oct 27, 2010
AB		GLENMARK GENERICS	<u>400MG</u> A090355 001 Jul 13, 2010
AB			<u>600MG</u> A090355 002 Jul 13, 2010
AB		INWOOD LABS	<u>450MG</u> A040034 001 Apr 28, 1995
AB		NOSTRUM	<u>400MG</u> A040595 001 Apr 21, 2006
AB	+		<u>600MG</u> A040560 002 Apr 21, 2006
AB	+	PLIVA	<u>100MG</u> A089807 001 Apr 30, 1990
AB	+		<u>200MG</u> A089808 001 Apr 30, 1990
AB			<u>300MG</u> A089763 001 Apr 30, 1990
AB	+		<u>450MG</u> A081236 001 Nov 09, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 395 (of 424)

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

<u>AP</u> + APP PHARMS	<u>100MG/ML</u>	<u>A080556 001</u>
<u>AP</u> WATSON LABS	<u>100MG/ML</u>	<u>A080571 001</u>
THIAMINE HYDROCHLORIDE		
+ WATSON LABS	200MG/ML	A080571 002

THIOGUANINE

TABLET; ORAL

THIOGUANINE

+ GLAXOSMITHKLINE	40MG	N012429 001
-------------------	------	-------------

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

<u>AB</u> MUTUAL PHARM	<u>10MG</u>	<u>A089953 004</u>	Aug 01, 1986
<u>AB</u>	<u>25MG</u>	<u>A089953 003</u>	Aug 01, 1986
<u>AB</u>	<u>50MG</u>	<u>A089953 002</u>	Aug 01, 1986
<u>AB</u>	<u>100MG</u>	<u>A089953 001</u>	Oct 07, 1988
<u>AB</u> MYLAN	<u>10MG</u>	<u>A088004 002</u>	Mar 15, 1983
<u>AB</u>	<u>25MG</u>	<u>A088004 003</u>	Mar 15, 1983
<u>AB</u>	<u>50MG</u>	<u>A088004 004</u>	Mar 15, 1983
<u>AB</u> +	<u>100MG</u>	<u>A088004 001</u>	Nov 18, 1983

THIOTEP A

INJECTABLE; INJECTION

THIOTEP A

BEDFORD	15MG/VIAL	A075547 001	Apr 02, 2001
---------	-----------	-------------	--------------

THIOTHIXENE

CAPSULE; ORAL

NAVANE

<u>AB</u> PFIZER	<u>1MG</u>	<u>N016584 001</u>	
<u>AB</u>	<u>2MG</u>	<u>N016584 002</u>	
<u>AB</u> +	<u>5MG</u>	<u>N016584 003</u>	
<u>AB</u>	<u>10MG</u>	<u>N016584 004</u>	

THIOTHIXENE

<u>AB</u> MYLAN	<u>1MG</u>	<u>A071093 002</u>	Jun 23, 1987
<u>AB</u>	<u>2MG</u>	<u>A071093 003</u>	Jun 23, 1987
<u>AB</u>	<u>5MG</u>	<u>A071093 004</u>	Jun 23, 1987
<u>AB</u>	<u>10MG</u>	<u>A071093 001</u>	Jun 23, 1987
<u>AB</u> SANDOZ	<u>1MG</u>	<u>A071610 001</u>	Jun 24, 1987
<u>AB</u>	<u>2MG</u>	<u>A071570 001</u>	Jun 24, 1987
<u>AB</u>	<u>5MG</u>	<u>A071529 001</u>	Jun 24, 1987
<u>AB</u>	<u>10MG</u>	<u>A071530 001</u>	Jun 24, 1987
<u>AB</u> WATSON LABS	<u>2MG</u>	<u>A070601 001</u>	Jun 05, 1987
<u>AB</u>	<u>5MG</u>	<u>A070602 001</u>	Jun 05, 1987

THYROTROPIN ALFA

INJECTABLE; INJECTION

THYROID

+ GENZYME	1.1MG/VIAL	N020898 001	Nov 30, 1998
-----------	------------	-------------	--------------

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

<u>AB</u> CEPHALON	<u>2MG</u>	<u>N020646 005</u>	Apr 16, 1999
<u>AB</u> +	<u>4MG</u>	<u>N020646 001</u>	Sep 30, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 396 (of 424)

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

TIAGABINE HYDROCHLORIDE

<u>AB</u>	SUN PHARM IND'S	<u>2MG</u>	<u>A077555</u>	<u>001</u>	Nov 04, 2011
<u>AB</u>		<u>4MG</u>	<u>A077555</u>	<u>002</u>	Nov 04, 2011
	GABITRIL				
	CEPHALON	6MG	N020646	006	Nov 29, 2005
		8MG	N020646	007	Nov 29, 2005
		10MG	N020646	008	Nov 29, 2005
		12MG	N020646	002	Sep 30, 1997
		16MG	N020646	003	Sep 30, 1997

TICAGRELOR

TABLET; ORAL

BRILINTA

+ ASTRAZENECA LP 90MG

N022433 001 Jul 20, 2011

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>250MG</u>	<u>A075089</u>	<u>001</u>	Jul 01, 1999
<u>AB</u>	CARACO	<u>250MG</u>	<u>A075526</u>	<u>001</u>	Sep 26, 2002
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A075161</u>	<u>001</u>	Sep 13, 1999
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A075326</u>	<u>001</u>	Aug 20, 1999
<u>AB</u> + TEVA		<u>250MG</u>	<u>A075149</u>	<u>001</u>	Aug 20, 1999

TIGECYCLINE

INJECTABLE; IV (INFUSION)

TYGACIL

+ WYETH PHARMS INC 50MG/VIAL

N021821 001 Jun 15, 2005

TILOUDRONATE DISODIUM

TABLET; ORAL

SKELID

+ SANOFI AVENTIS US EQ 200MG BASE

N020707 001 Mar 07, 1997

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

+ SANTEN OY	EQ 0.25% BASE	N020439	001	Mar 31, 1995
+	EQ 0.5% BASE	N020439	002	Mar 31, 1995

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC

TIMOLOL MALEATE

<u>AB</u>	FALCON PHARMS	<u>EQ 0.25% BASE</u>	<u>N020963</u>	<u>001</u>	Oct 21, 1998
<u>AB</u>		<u>EQ 0.5% BASE</u>	<u>N020963</u>	<u>002</u>	Oct 21, 1998

TIMOPTIC-XE

<u>AB</u> + ATON		<u>EQ 0.25% BASE</u>	<u>N020330</u>	<u>001</u>	Nov 04, 1993
<u>AB</u> +		<u>EQ 0.5% BASE</u>	<u>N020330</u>	<u>002</u>	Nov 04, 1993

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

<u>AT</u>	AKORN	<u>EQ 0.25% BASE</u>	<u>A074515</u>	<u>001</u>	Mar 25, 1997
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A074466</u>	<u>001</u>	Mar 25, 1997
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A074516</u>	<u>001</u>	Mar 25, 1997
<u>AT</u>	APOTEX INC	<u>EQ 0.25% BASE</u>	<u>A075411</u>	<u>001</u>	Sep 08, 2000
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A075412</u>	<u>001</u>	Sep 08, 2000
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.25% BASE</u>	<u>A074778</u>	<u>001</u>	Mar 25, 1997
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A074776</u>	<u>001</u>	Mar 25, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 397 (of 424)

TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

<u>AT</u>	FALCON PHARMS	<u>EQ 0.25% BASE</u>	<u>A074261</u>	<u>001</u>	Apr 28, 1995
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A074262</u>	<u>001</u>	Apr 28, 1995
<u>AT</u>	FDC LTD	<u>EQ 0.25% BASE</u>	<u>A077259</u>	<u>001</u>	Apr 30, 2008
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A077259</u>	<u>002</u>	Apr 30, 2008
<u>AT</u>	HI TECH PHARMA	<u>EQ 0.5% BASE</u>	<u>A075163</u>	<u>001</u>	Sep 10, 2002
<u>AT</u>	PACIFIC PHARMA	<u>EQ 0.25% BASE</u>	<u>A074746</u>	<u>001</u>	Mar 25, 1997
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A074747</u>	<u>001</u>	Mar 25, 1997
<u>AT</u>	WOCKHARDT	<u>EQ 0.25% BASE</u>	<u>A078771</u>	<u>001</u>	Sep 28, 2009
<u>AT</u>		<u>EQ 0.5% BASE</u>	<u>A078771</u>	<u>002</u>	Sep 28, 2009
<u>TIMOPTIC</u>					
<u>AT</u>	+ ATON	<u>EQ 0.25% BASE</u>	<u>N018086</u>	<u>001</u>	
<u>AT</u>	+ ISTALOL	<u>EQ 0.5% BASE</u>	<u>N018086</u>	<u>002</u>	
BT	+ ISTA PHARMS	EQ 0.5% BASE	N021516	001	Jun 04, 2004
	TIMOPTIC IN OCUDOSE				
	+ ATON	EQ 0.25% BASE	N019463	001	Nov 05, 1986
	+	EQ 0.5% BASE	N019463	002	Nov 05, 1986
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

MISSION PHARMA	250MG	N021618	001	May 17, 2004
+	500MG	N021618	002	May 17, 2004

TINZAPARIN SODIUM

INJECTABLE; INJECTION

INNOHEP

+ LEO PHARMA AS	20,000 IU/ML	N020484	001	Jul 14, 2000
-----------------	--------------	---------	-----	--------------

TIOPRONIN

TABLET; ORAL

TIOPRONIN

+ MISSION PHARMA	100MG	N019569	001	Aug 11, 1988
------------------	-------	---------	-----	--------------

TIOTROPIUM BROMIDE MONOHYDRATE

POWDER; INHALATION

SPIRIVA

+ BOEHRINGER INGELHEIM	EQ 0.018MG BASE/INH	N021395	001	Jan 30, 2004
------------------------	---------------------	---------	-----	--------------

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
----------	---------------------------------------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 398 (of 424)

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION
 AGGRASTAT
 + MEDICURE EQ 12.5MG BASE/250ML (EQ 0.05MG
 BASE/ML) N020913 003 Apr 20, 2000

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL
 ZANAFLEX
 ACORDA EQ 2MG BASE N021447 001 Aug 29, 2002
 EQ 4MG BASE N021447 002 Aug 29, 2002
 + EQ 6MG BASE N021447 003 Aug 29, 2002

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>EQ 2MG BASE</u>	<u>A076282 001</u>	Dec 16, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076282 002</u>	Dec 16, 2003
<u>AB</u>	APOTEX	<u>EQ 2MG BASE</u>	<u>A076533 001</u>	Jan 16, 2004
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076533 002</u>	Jan 16, 2004
<u>AB</u>	CARACO	<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>	COREPHARMA	<u>EQ 2MG BASE</u>	<u>A076347 001</u>	Oct 11, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076347 002</u>	Oct 11, 2002
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 2MG BASE</u>	<u>A076286 001</u>	Jul 03, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076286 002</u>	Jul 03, 2002
<u>AB</u>	MYLAN	<u>EQ 2MG BASE</u>	<u>A076354 001</u>	Mar 28, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076354 002</u>	Mar 28, 2003
<u>AB</u>	SANDOZ	<u>EQ 2MG BASE</u>	<u>A076399 001</u>	Nov 26, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076280 002</u>	Jun 27, 2002
<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>ZANAFLEX</u>				
<u>AB</u>	+ ACORDA	<u>EQ 4MG BASE</u>	<u>N020397 001</u>	Nov 27, 1996

TOBRAMYCIN

OINTMENT; OPHTHALMIC
 TOBREX
 + ALCON 0 . 3 % N050555 001

SOLUTION; INHALATION
 TOBI
 + NOVARTIS PHARMS 300MG/5ML N050753 001 Dec 22, 1997

SOLUTION/DROPS; OPHTHALMIC

<u>AKTOB</u>				
<u>AT</u>	AKORN	<u>0 . 3 %</u>	<u>A064096 001</u>	Jan 31, 1996
<u>TOBRAMYCIN</u>				
<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>	NOVEX	<u>0 . 3 %</u>	<u>A065087 001</u>	Feb 25, 2002
<u>TOBREX</u>				
<u>AT</u>	ALCON	<u>0 . 3 %</u>	<u>A062535 001</u>	Dec 13, 1984
<u>AT</u>	+ FALCON PHARMS	<u>0 . 3 %</u>	<u>N050541 001</u>	

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION
TOBRAMYCIN SULFATE

<u>AP</u>	AKORN STRIDES	<u>EQ 40MG BASE/ML</u>	<u>A065407 001</u>	Mar 11, 2008
<u>AP</u>	APOTHECON	<u>EQ 10MG BASE/ML</u>	<u>A064021 001</u>	May 31, 1994
<u>AP</u>	APP PHARMS	<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

PRESCRIPTION DRUG PRODUCT LIST

3 - 400 (of 424)

TOLVAPTAN

TABLET; ORAL

SAMSCA

OTSKA AMERICA PHARM	15MG	N022275 001	May 19, 2009
+	30MG	N022275 002	May 19, 2009

TOPIRAMATE

CAPSULE; ORAL

TOPAMAX

AB JANSSEN PHARMS	<u>15MG</u>	<u>N020844 001</u>	Oct 26, 1998
AB +	<u>25MG</u>	<u>N020844 002</u>	Oct 26, 1998

TOPIRAMATE

AB MYLAN	<u>15MG</u>	<u>A078418 001</u>	Oct 14, 2009
AB	<u>25MG</u>	<u>A078418 002</u>	Oct 14, 2009
AB SANDOZ	<u>15MG</u>	<u>A079206 001</u>	Oct 14, 2009
AB	<u>25MG</u>	<u>A079206 002</u>	Oct 14, 2009
AB TEVA	<u>15MG</u>	<u>A076575 001</u>	Apr 17, 2009
AB	<u>25MG</u>	<u>A076575 002</u>	Apr 17, 2009
AB WATSON LABS	<u>15MG</u>	<u>A077868 001</u>	Apr 15, 2009
AB	<u>25MG</u>	<u>A077868 002</u>	Apr 15, 2009
AB ZYDUS PHARMS USA INC	<u>15MG</u>	<u>A078877 001</u>	Oct 14, 2009
AB	<u>25MG</u>	<u>A078877 002</u>	Oct 14, 2009

TABLET; ORAL

TOPAMAX

AB + JANSSEN PHARMS	<u>25MG</u>	<u>N020505 004</u>	Dec 24, 1996
AB	<u>50MG</u>	<u>N020505 005</u>	Dec 24, 1996
AB	<u>100MG</u>	<u>N020505 001</u>	Dec 24, 1996
AB	<u>200MG</u>	<u>N020505 002</u>	Dec 24, 1996

TOPIRAMATE

AB ACCORD HLTHCARE	<u>25MG</u>	<u>A076311 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A076311 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A076311 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A076311 004</u>	Mar 27, 2009
AB APOTEX INC	<u>25MG</u>	<u>A077733 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A077733 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A077733 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A077733 004</u>	Mar 27, 2009
AB AUROBINDO PHARMA	<u>25MG</u>	<u>A078462 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A078462 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A078462 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A078462 004</u>	Mar 27, 2009
AB CIPLA LTD	<u>25MG</u>	<u>A076343 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A076343 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A076343 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A076343 004</u>	Mar 27, 2009
AB GLENMARK GENERICS	<u>25MG</u>	<u>A077627 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A077627 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A077627 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A077627 004</u>	Mar 27, 2009
AB INVAGEN PHARMS	<u>25MG</u>	<u>A079162 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A079162 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A079162 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A079162 004</u>	Mar 27, 2009
AB MYLAN	<u>25MG</u>	<u>A076314 001</u>	Mar 27, 2009
AB	<u>50MG</u>	<u>A076314 002</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A076314 003</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A076314 004</u>	Mar 27, 2009
AB RANBAXY	<u>25MG</u>	<u>A076327 001</u>	Mar 27, 2009
AB	<u>100MG</u>	<u>A076327 002</u>	Mar 27, 2009
AB	<u>200MG</u>	<u>A076327 003</u>	Mar 27, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 401 (of 424)

TOPIRAMATE

TABLET; ORAL

TOPIRAMATE

<u>AB</u>	SUN PHARM INDS LTD	<u>25MG</u>	<u>A090278</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A090278</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A090278</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A090278</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	TEVA	<u>25MG</u>	<u>A076317</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A076317</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A076317</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A076317</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	TORRENT PHARMS	<u>25MG</u>	<u>A079153</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A079153</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079153</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A079153</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A090162</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A090162</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A090162</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>	UPSHER SMITH	<u>25MG</u>	<u>A078499</u>	<u>001</u>	Jan 07, 2010
<u>AB</u>		<u>50MG</u>	<u>A078499</u>	<u>002</u>	Jan 07, 2010
<u>AB</u>		<u>100MG</u>	<u>A078499</u>	<u>003</u>	Jan 07, 2010
<u>AB</u>		<u>200MG</u>	<u>A078499</u>	<u>004</u>	Jan 07, 2010
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A077643</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A077643</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A077643</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A077643</u>	<u>004</u>	Mar 27, 2009
<u>AB</u>	WOCKHARDT USA	<u>25MG</u>	<u>A090353</u>	<u>001</u>	Sep 01, 2010
<u>AB</u>		<u>50MG</u>	<u>A090353</u>	<u>002</u>	Sep 01, 2010
<u>AB</u>		<u>100MG</u>	<u>A090353</u>	<u>003</u>	Sep 01, 2010
<u>AB</u>		<u>200MG</u>	<u>A090353</u>	<u>004</u>	Sep 01, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078235</u>	<u>001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A078235</u>	<u>002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078235</u>	<u>003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078235</u>	<u>004</u>	Mar 27, 2009

TOPOTECAN HYDROCHLORIDE

CAPSULE; ORAL

HYCAMTIN

SMITHKLINE BEECHAM	EQ 0.25MG BASE	N020981	001	Oct 11, 2007
+	EQ 1MG BASE	N020981	002	Oct 11, 2007

INJECTABLE; INJECTION

HYCAMTIN

<u>AP</u>	+ GLAXOSMITHKLINE	<u>EQ 4MG BASE/VIAL</u>	<u>N020671</u>	<u>001</u>	May 28, 1996
<u>AP</u>	<u>TOPOTECAN HYDROCHLORIDE</u>				
<u>AP</u>	ACTAVIS TOTOWA	<u>EQ 4MG BASE/VIAL</u>	<u>A090620</u>	<u>001</u>	Dec 02, 2010
<u>AP</u>	APP PHARMS	<u>EQ 4MG BASE/VIAL</u>	<u>A091089</u>	<u>001</u>	Nov 29, 2010
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A201191</u>	<u>001</u>	Mar 09, 2011
<u>AP</u>	FRESENIUS KABI ONCOL	<u>EQ 4MG BASE/VIAL</u>	<u>A091376</u>	<u>001</u>	Nov 29, 2010
<u>AP</u>	SAGENT PHARMS	<u>EQ 4MG BASE/VIAL</u>	<u>A091284</u>	<u>001</u>	Jan 26, 2011
<u>AP</u>	THREE RIVERS PHARMS	<u>EQ 4MG BASE/VIAL</u>	<u>A091199</u>	<u>001</u>	Dec 01, 2010

SOLUTION; INTRAVENOUS

TOPOTECAN

+	HOSPIRA INC	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	N200582	001	Feb 02, 2011
---	-------------	----------------------------------	---------	-----	--------------

TOREMIFENE CITRATE

TABLET; ORAL

FARESTON

+	GTX INC	EQ 60MG BASE	N020497	001	May 29, 1997
---	---------	--------------	---------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 402 (of 424)

TORSEMIDE

INJECTABLE; INJECTION

TORSEMIDE

<u>AP</u>	+ BEDFORD LABS	<u>20MG/2ML (10MG/ML)</u>	<u>A078007 001</u>	Jun 11, 2008
<u>AP</u>	+ LUITPOLD	<u>50MG/5ML (10MG/ML)</u>	<u>A078007 002</u>	Jun 11, 2008
<u>AP</u>	LUITPOLD	<u>50MG/5ML (10MG/ML)</u>	<u>A090656 002</u>	Apr 21, 2010
<u>AP</u>		<u>20MG/2ML (10MG/ML)</u>	<u>A090656 001</u>	Apr 21, 2010

TABLET; ORAL

DEMADEX

<u>AB</u>	MEDA PHARMS	<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
<u>AB</u>	HETERO DRUGS	<u>5MG</u>	<u>A079234 001</u>	Jan 27, 2009
<u>AB</u>		<u>10MG</u>	<u>A079234 002</u>	Jan 27, 2009
<u>AB</u>		<u>20MG</u>	<u>A079234 003</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079234 004</u>	Jan 27, 2009
<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>A076226 001</u>	May 27, 2003
<u>AB</u>		<u>10MG</u>	<u>A076226 002</u>	May 27, 2003
<u>AB</u>		<u>20MG</u>	<u>A076226 003</u>	May 27, 2003
<u>AB</u>		<u>100MG</u>	<u>A076226 004</u>	May 27, 2003
<u>AB</u>	PLIVA PHARM IND	<u>5MG</u>	<u>A076346 001</u>	May 30, 2003
<u>AB</u>		<u>10MG</u>	<u>A076346 002</u>	May 30, 2003
<u>AB</u>		<u>20MG</u>	<u>A076346 003</u>	May 30, 2003
<u>AB</u>		<u>100MG</u>	<u>A076346 004</u>	Oct 19, 2004
<u>AB</u>	ROXANE	<u>5MG</u>	<u>A076943 001</u>	Mar 01, 2005
<u>AB</u>		<u>10MG</u>	<u>A076943 002</u>	Mar 01, 2005
<u>AB</u>		<u>20MG</u>	<u>A076943 003</u>	Mar 01, 2005
<u>AB</u>	SUN PHARM INDNS	<u>5MG</u>	<u>A078478 001</u>	Feb 26, 2008
<u>AB</u>		<u>10MG</u>	<u>A078478 002</u>	Feb 26, 2008
<u>AB</u>		<u>20MG</u>	<u>A078478 003</u>	Feb 26, 2008
<u>AB</u>		<u>100MG</u>	<u>A078478 004</u>	Feb 26, 2008
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076110 001</u>	May 14, 2002
<u>AB</u>		<u>10MG</u>	<u>A076110 002</u>	May 14, 2002
<u>AB</u>		<u>20MG</u>	<u>A076110 003</u>	May 14, 2002
<u>AB</u>		<u>100MG</u>	<u>A076110 004</u>	May 14, 2002
<u>AB</u>	VINTAGE PHARMS	<u>5MG</u>	<u>A090613 001</u>	Mar 22, 2011
<u>AB</u>		<u>10MG</u>	<u>A090613 002</u>	Mar 22, 2011
<u>AB</u>		<u>20MG</u>	<u>A090613 003</u>	Mar 22, 2011
<u>AB</u>		<u>100MG</u>	<u>A090613 004</u>	Mar 22, 2011

TRAMADOL HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
CONZIP

<u>+</u>	CIPHER PHARMS INC	<u>100MG</u>	<u>N022370 001</u>	May 07, 2010
		<u>150MG</u>	<u>N022370 004</u>	Aug 01, 2011
		<u>200MG</u>	<u>N022370 002</u>	May 07, 2010
		<u>300MG</u>	<u>N022370 003</u>	May 07, 2010

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>50MG</u>	<u>A075980 001</u>	Nov 21, 2002
-----------	------------	-------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 403 (of 424)

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB</u>	ALVOGEN	<u>50MG</u>	<u>A202075</u>	<u>001</u>	Nov 28, 2011
<u>AB</u>	AMNEAL PHARMS	<u>50MG</u>	<u>A076003</u>	<u>001</u>	Jun 20, 2002
<u>AB</u>	APOTEX	<u>50MG</u>	<u>A075981</u>	<u>001</u>	Jul 10, 2002
<u>AB</u>	CARACO	<u>50MG</u>	<u>A075964</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	MALLINCKRODT	<u>50MG</u>	<u>A075983</u>	<u>001</u>	Jun 25, 2002
<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>A076100</u>	<u>001</u>	Jun 20, 2002
<u>AB</u>	MYLAN	<u>50MG</u>	<u>A075986</u>	<u>001</u>	Jun 21, 2002
<u>AB</u>	NORTHSTAR HLTHCARE	<u>50MG</u>	<u>A078935</u>	<u>001</u>	May 26, 2010
<u>AB</u>	PLIVA	<u>50MG</u>	<u>A075982</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>A075968</u>	<u>001</u>	Jun 25, 2002
<u>AB</u>	TEVA	<u>50MG</u>	<u>A075977</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>A075962</u>	<u>001</u>	Jun 24, 2002
<u>AB</u>	ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A090404</u>	<u>001</u>	Jan 31, 2011

ULTRAM

<u>AB</u>	+ JANSSEN PHARMS	<u>50MG</u>	<u>N020281</u>	<u>002</u>	Mar 03, 1995
-----------	------------------	-------------	----------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB1</u>	LUPIN LTD	<u>100MG</u>	<u>A200503</u>	<u>001</u>	Aug 29, 2011
<u>AB1</u>		<u>200MG</u>	<u>A200503</u>	<u>002</u>	Aug 29, 2011
<u>AB1</u>		<u>300MG</u>	<u>A200503</u>	<u>003</u>	Aug 29, 2011
<u>AB1</u>	PAR PHARM	<u>100MG</u>	<u>A078783</u>	<u>001</u>	Nov 13, 2009
<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

ULTRAM ER

<u>AB1</u>	+ VALEANT INTL	<u>100MG</u>	<u>N021692</u>	<u>001</u>	Sep 08, 2005
<u>AB1</u>		<u>200MG</u>	<u>N021692</u>	<u>002</u>	Sep 08, 2005
<u>AB1</u>		<u>300MG</u>	<u>N021692</u>	<u>003</u>	Sep 08, 2005

RYZOLT

<u>AB2</u>	+ PURDUE PHARMA	<u>100MG</u>	<u>N021745</u>	<u>001</u>	Dec 30, 2008
<u>AB2</u>		<u>200MG</u>	<u>N021745</u>	<u>002</u>	Dec 30, 2008
<u>AB2</u>		<u>300MG</u>	<u>N021745</u>	<u>003</u>	Dec 30, 2008

TRAMADOL HYDROCHLORIDE

<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

TABLET, ORALLY DISINTEGRATING; ORAL

TRAMADOL HYDROCHLORIDE

+ SHIONOGI INC		50MG	N021693	001	May 05, 2005
----------------	--	------	---------	-----	--------------

TRANDOLAPRIL

TABLET; ORAL

MAVIK

<u>AB</u>	ABBOTT	<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>	CIPLA	<u>1MG</u>	<u>A077307</u>	<u>002</u>	Jun 12, 2007
<u>AB</u>		<u>2MG</u>	<u>A077307</u>	<u>001</u>	Jun 12, 2007
<u>AB</u>		<u>4MG</u>	<u>A077307</u>	<u>003</u>	Jun 12, 2007
<u>AB</u>	COREPHARMA	<u>1MG</u>	<u>A077256</u>	<u>001</u>	Jun 12, 2007
<u>AB</u>		<u>2MG</u>	<u>A077256</u>	<u>002</u>	Jun 12, 2007

PRESCRIPTION DRUG PRODUCT LIST

3 - 404 (of 424)

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

<u>AB</u>	COREPHARMA	<u>4MG</u>	<u>A077256</u>	<u>003</u>	Jun 12, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A078493</u>	<u>001</u>	Aug 25, 2008
<u>AB</u>		<u>2MG</u>	<u>A078493</u>	<u>002</u>	Aug 25, 2008
<u>AB</u>		<u>4MG</u>	<u>A078493</u>	<u>003</u>	Aug 25, 2008
<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>	INVAGEN PHARMS	<u>1MG</u>	<u>A078320</u>	<u>001</u>	Jun 12, 2007
<u>AB</u>		<u>2MG</u>	<u>A078320</u>	<u>002</u>	Jun 12, 2007
<u>AB</u>		<u>4MG</u>	<u>A078320</u>	<u>003</u>	Jun 12, 2007
<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>	MYLAN	<u>1MG</u>	<u>A078346</u>	<u>001</u>	Apr 28, 2008
<u>AB</u>		<u>2MG</u>	<u>A078346</u>	<u>002</u>	Apr 28, 2008
<u>AB</u>		<u>4MG</u>	<u>A078346</u>	<u>003</u>	Apr 28, 2008
<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>	ABBOTT	<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
		<u>TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE</u>			
<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
		<u>TRANEXAMIC ACID</u>			
<u>AP</u>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A091657</u>	<u>001</u>	Nov 03, 2011
<u>AP</u>	PHARMAFORCE	<u>100MG/ML</u>	<u>A201885</u>	<u>001</u>	Aug 10, 2011
<u>AP</u>	VERSAPHARM INC	<u>100MG/ML</u>	<u>A202373</u>	<u>001</u>	Nov 17, 2011

TABLET; ORAL

LYSTEDA

+ FERRING PHARMS AS 650MG

N022430 001 Nov 13, 2009

TRANYLCYPROMINE SULFATE

TABLET; ORAL

PARNATE

<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 10MG BASE</u>	<u>N012342</u>	<u>003</u>	Aug 16, 1985
		<u>TRANYLCYPROMINE SULFATE</u>			
<u>AB</u>	PAR PHARM	<u>EQ 10MG BASE</u>	<u>A040640</u>	<u>001</u>	Jun 29, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 405 (of 424)

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC
 TRAVATAN Z
 + ALCON PHARMS LTD 0.004% N021994 001 Sep 21, 2006

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

<u>AB</u>	ALVOGEN	<u>50MG</u>	<u>A071636</u> <u>001</u>	Apr 18, 1988
<u>AB</u>		<u>100MG</u>	<u>A071514</u> <u>001</u>	Apr 18, 1988
<u>AB</u>	APOTEX	<u>50MG</u>	<u>A071258</u> <u>001</u>	Mar 25, 1987
<u>AB</u>	+ APOTEX INC	<u>100MG</u>	<u>A071196</u> <u>001</u>	Mar 25, 1987
<u>AB</u>		<u>150MG</u>	<u>A071196</u> <u>002</u>	Apr 26, 1999
<u>AB</u>		<u>300MG</u>	<u>A071196</u> <u>003</u>	Apr 26, 1999
<u>AB</u>	MATRIX LABS LTD	<u>50MG</u>	<u>A090514</u> <u>001</u>	Jun 02, 2009
<u>AB</u>		<u>100MG</u>	<u>A090514</u> <u>002</u>	Jun 02, 2009
<u>AB</u>		<u>150MG</u>	<u>A090514</u> <u>003</u>	Jun 02, 2009
<u>AB</u>		<u>300MG</u>	<u>A090514</u> <u>004</u>	Jun 02, 2009
<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>A073137</u> <u>002</u>	Mar 24, 1993
<u>AB</u>		<u>100MG</u>	<u>A073137</u> <u>001</u>	Mar 24, 1993
<u>AB</u>		<u>150MG</u>	<u>A073137</u> <u>003</u>	Dec 22, 1995
<u>AB</u>	PLIVA	<u>50MG</u>	<u>A071523</u> <u>001</u>	Dec 11, 1987
<u>AB</u>		<u>100MG</u>	<u>A071524</u> <u>001</u>	Dec 11, 1987
<u>AB</u>		<u>150MG</u>	<u>A071525</u> <u>001</u>	Mar 09, 1988
<u>AB</u>	VINTAGE	<u>50MG</u>	<u>A072192</u> <u>001</u>	Feb 02, 1989
<u>AB</u>		<u>100MG</u>	<u>A072193</u> <u>001</u>	Feb 02, 1989
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>A070857</u> <u>001</u>	Oct 10, 1986
<u>AB</u>		<u>100MG</u>	<u>A070858</u> <u>001</u>	Oct 10, 1986

TABLET, EXTENDED RELEASE; ORAL
 OLEPTRO
 + ANGELINI LLC 150MG N022411 001 Feb 02, 2010
 300MG N022411 002 Feb 02, 2010

TREPROSTINIL SODIUM

INJECTABLE; IV (INFUSION)-SC

UNITED THERAP	1MG/ML	N021272 001	May 21, 2002
	2.5MG/ML	N021272 002	May 21, 2002
	5MG/ML	N021272 003	May 21, 2002
+	10MG/ML	N021272 004	May 21, 2002

SOLUTION; INHALATION
 TYVASO
 + UNITED THERAP EQ 0.6MG BASE/ML N022387 001 Jul 30, 2009

TRETINOIN

CAPSULE; ORAL
 TRETINOIN
 + BARR 10MG A077684 001 Jun 22, 2007

CREAM; TOPICAL

<u>AB</u>	<u>AVITA</u>	<u>0.025%</u>	<u>N020404</u> <u>003</u>	Jan 14, 1997
<u>AB</u>	MYLAN BERTEK	<u>0.025%</u>		
<u>AB</u>	<u>RETIN-A</u>	<u>0.025%</u>	<u>N019049</u> <u>001</u>	Sep 16, 1988
<u>AB</u>	+ ORTHO JANSSEN	<u>0.1%</u>	<u>N017340</u> <u>001</u>	
<u>AB</u>	<u>TRETINOIN</u>	<u>0.025%</u>	<u>A075264</u> <u>001</u>	Dec 24, 1998
<u>AB</u>	TRIAK PHARMS LLC	<u>0.1%</u>	<u>A075213</u> <u>001</u>	Dec 24, 1998
<u>AB1</u>	<u>RETIN-A</u>	<u>0.05%</u>	<u>N017522</u> <u>001</u>	
<u>AB1</u>	+ ORTHO JANSSEN			

PRESCRIPTION DRUG PRODUCT LIST

3 - 406 (of 424)

TRETINOIN

CREAM; TOPICAL

TRETINOIN

<u>AB1</u>	TRIAx PHARMS LLC	<u>0.05%</u>	<u>A075265</u> <u>001</u>	Dec 24, 1998
	<u>RENOVA</u>			
<u>AB2</u> +	ORTHO JANSSEN	<u>0.05%</u>	<u>N019963</u> <u>001</u>	Dec 29, 1995
	<u>TRETINOIN</u>			
<u>AB2</u>	SPEAR PHARMS	<u>0.05%</u>	<u>A076498</u> <u>001</u>	Sep 15, 2005
	RENOVA			
+ ORTHO JANSSEN		0.02%	N021108 001	Aug 31, 2000
TRETINOIN				
+ TRIAX PHARMS LLC		0.0375%	A090098 001	Mar 22, 2010

GEL; TOPICAL

RETIN-A

<u>AB</u>	+ ORTHO JANSSEN	<u>0.01%</u>	<u>N017955</u> <u>001</u>	
<u>AB</u>	+	<u>0.025%</u>	<u>N017579</u> <u>002</u>	
	<u>TRETINOIN</u>			
<u>AB</u>	TRIAx PHARMS LLC	<u>0.01%</u>	<u>A075589</u> <u>001</u>	Jun 11, 2002
<u>AB</u>		<u>0.025%</u>	<u>A075529</u> <u>001</u>	Feb 22, 2000
	AVITA			
BT	MYLAN	0.025%	N020400 001	Jan 29, 1998
	ATRALIN			
+ DOW PHARM SCIENCES		0.05%	N022070 001	Jul 26, 2007
RETIN-A MICRO				
+ VALEANT INTL		0.04%	N020475 002	May 10, 2002
+		0.1%	N020475 001	Feb 07, 1997

SOLUTION; TOPICAL

RETIN-A

<u>AT</u>	+ ORTHO JANSSEN	<u>0.05%</u>	<u>N016921</u> <u>001</u>	
<u>AT</u>	<u>TRETINOIN</u>		<u>A075260</u> <u>001</u>	Jan 25, 1999
	WOCKHARDT	<u>0.05%</u>		

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; INHALATION

AZMACORT

+ ABBOTT		0.1MG/INH	N018117 001	Apr 23, 1982
----------	--	-----------	-------------	--------------

CREAM; TOPICAL

KENALOG

<u>AT</u>	+ APOTHECON	<u>0.025%</u>	<u>N011601</u> <u>003</u>	
<u>AT</u>	+	<u>0.1%</u>	<u>N011601</u> <u>006</u>	

TRIACET

<u>AT</u>	TEVA	<u>0.025%</u>	<u>A084908</u> <u>001</u>	
<u>AT</u>		<u>0.5%</u>	<u>A084908</u> <u>003</u>	

TRIAMCINOLONE ACETONIDE

<u>AT</u>	ALTANA	<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>	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>0.025%</u>	<u>A086277</u> <u>001</u>	
<u>AT</u>		<u>0.1%</u>	<u>A040039</u> <u>001</u>	Nov 26, 1997
<u>AT</u>		<u>0.1%</u>	<u>A086276</u> <u>001</u>	
<u>AT</u>		<u>0.5%</u>	<u>A086275</u> <u>001</u>	
<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>0.1%</u>	<u>A088042</u> <u>001</u>	Mar 19, 1984

PRESCRIPTION DRUG PRODUCT LIST

3 - 407 (of 424)

TRIAMCINOLONE ACETONIDE

INJECTABLE; INJECTION			
<u>KENALOG-10</u>			
AB	APOTHECON	<u>10MG/ML</u>	N012041 001
<u>KENALOG-40</u>			
AB	+ APOTHECON	<u>40MG/ML</u>	N014901 001
<u>TRIAMCINOLONE ACETONIDE</u>			
AB	SANDOZ	<u>10MG/ML</u>	A090166 001 May 27, 2009
AB		<u>40MG/ML</u>	A090164 001 May 27, 2009
INJECTABLE; INTRA-ARTICULAR, INTRAMUSCULAR, INTRAVITREAL			
TRIVARIS			
+ ALLERGAN		8MG/0.1ML (8MG/0.1ML)	N022220 001 Jun 16, 2008
INJECTABLE; INTRAVITREAL			
TRIESENCE			
+ ALCON		40MG/ML (40MG/ML)	N022048 001 Nov 29, 2007
LOTION; TOPICAL			
<u>TRIAMCINOLONE ACETONIDE</u>			
AT	ALTANA	<u>0.025%</u>	A040467 001 Apr 21, 2003
AT		<u>0.1%</u>	A040467 002 Apr 21, 2003
AT	MORTON GROVE	<u>0.025%</u>	A088450 001 Apr 01, 1985
AT	TARO	<u>0.1%</u>	A089129 001 Aug 14, 1986
AT	VINTAGE	<u>0.1%</u>	A040672 002 Dec 13, 2006
AT	+ WOCKHARDT	<u>0.1%</u>	A088451 001 Apr 03, 1985
OINTMENT; TOPICAL			
<u>KENALOG</u>			
AT	APOTHECON	<u>0.025%</u>	N011600 003
AT		<u>0.1%</u>	N011600 001
<u>TRIAMCINOLONE ACETONIDE</u>			
AT	NYCOMED US	<u>0.025%</u>	A085691 001
AT		<u>0.1%</u>	A085691 003
AT		<u>0.5%</u>	A085691 002
AT	+ PERRIGO NEW YORK	<u>0.025%</u>	A087356 001
AT	+	<u>0.1%</u>	A087357 001
AT	+	<u>0.5%</u>	A087385 001
AT	TARO	<u>0.025%</u>	A040040 001 Sep 30, 1994
AT		<u>0.025%</u>	A040374 001 Jun 05, 2001
AT		<u>0.1%</u>	A040037 001 Sep 30, 1994
AT		<u>0.1%</u>	A087902 001 Dec 27, 1982
AT		<u>0.5%</u>	A040386 001 Jun 05, 2001
TRIAMCINOLONE ACETONIDE IN ABSORBASE			
+ CAROLINA MEDCL		0.05%	A089595 001 Mar 23, 1995
PASTE; DENTAL			
<u>TRIAMCINOLONE ACETONIDE</u>			
AT	LYNE	<u>0.1%</u>	A040771 001 Jul 01, 2010
AT	+ TARO	<u>0.1%</u>	A070730 001 Oct 01, 1986
SPRAY; TOPICAL			
KENALOG			
+ RANBAXY		0.147MG/GM	N012104 001
SPRAY, METERED; NASAL			
<u>NASACORT AQ</u>			
AB	+ SANOFI AVENTIS US	<u>0.055MG/SPRAY</u>	N020468 001 May 20, 1996
<u>TRIAMCINOLONE ACETONIDE</u>			
AB	TEVA BRANDED PHARM	<u>0.055MG/SPRAY</u>	A078104 001 Jul 30, 2009
<u>TRIAMCINOLONE HEXACETONIDE</u>			
INJECTABLE; INJECTION			
ARISTOSPAN			
+ SANDOZ		5MG/ML	N016466 001
+		20MG/ML	N016466 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 408 (of 424)

TRIAMTERENE

CAPSULE; ORAL			
DYRENIUM			
WELLSPRING PHARM	50MG	N013174 001	
+	100MG	N013174 002	

TRIAZOLAM

TABLET; ORAL			
<u>HALCION</u>			
AB PHARMACIA AND UPJOHN	<u>0.125MG</u>	<u>N017892 003</u>	Apr 26, 1985
AB +	<u>0.25MG</u>	<u>N017892 001</u>	Nov 15, 1982
<u>TRIAZOLAM</u>			
AB ALPHAPHARM	<u>0.125MG</u>	<u>A074031 001</u>	Mar 25, 1994
AB	<u>0.25MG</u>	<u>A074031 002</u>	Mar 25, 1994
AB ROXANE	<u>0.125MG</u>	<u>A074224 001</u>	Jun 01, 1994
AB	<u>0.25MG</u>	<u>A074224 002</u>	Jun 01, 1994
AB WATSON LABS	<u>0.125MG</u>	<u>A074445 001</u>	Oct 20, 1995
AB	<u>0.25MG</u>	<u>A074445 002</u>	Oct 20, 1995

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL			
SYPRINE			
+ ATON	250MG	N019194 001	Nov 08, 1985

TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL			
<u>TRIFLUOPERAZINE HYDROCHLORIDE</u>			
AB MYLAN	<u>EQ 1MG BASE</u>	<u>A040209 001</u>	Jul 07, 1997
AB	<u>EQ 2MG BASE</u>	<u>A040209 002</u>	Jul 07, 1997
AB	<u>EQ 5MG BASE</u>	<u>A040209 003</u>	Jul 07, 1997
AB +	<u>EQ 10MG BASE</u>	<u>A040209 004</u>	Jul 07, 1997
AB SANDOZ	<u>EQ 1MG BASE</u>	<u>A085785 001</u>	
AB	<u>EQ 2MG BASE</u>	<u>A085786 001</u>	
AB	<u>EQ 5MG BASE</u>	<u>A085789 001</u>	
AB	<u>EQ 10MG BASE</u>	<u>A085788 001</u>	

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC			
<u>TRIFLURIDINE</u>			
AT ALCON	<u>1%</u>	<u>A074311 001</u>	Oct 06, 1995
<u>VIROPTIC</u>			
AT + MONARCH PHARMS	<u>1%</u>	<u>N018299 001</u>	

TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL			
<u>TRIHEXYPHENIDYL HYDROCHLORIDE</u>			
AA MIKART	<u>2MG/5ML</u>	<u>A040251 001</u>	Sep 27, 1999
AA + PHARM ASSOC	<u>2MG/5ML</u>	<u>A040177 001</u>	Apr 17, 1997

TABLET; ORAL

<u>TRIHEXYPHENIDYL HYDROCHLORIDE</u>			
AA NATCO PHARMA LTD	<u>2MG</u>	<u>A091630 001</u>	Nov 17, 2010
AA	<u>5MG</u>	<u>A091630 002</u>	Nov 17, 2010
AA VINTAGE PHARMS	<u>2MG</u>	<u>A040254 001</u>	Dec 24, 1998
AA	<u>5MG</u>	<u>A040254 002</u>	Dec 24, 1998
AA WATSON LABS	<u>2MG</u>	<u>A040184 001</u>	Feb 06, 1998
AA +	<u>2MG</u>	<u>A084363 001</u>	
AA	<u>5MG</u>	<u>A040184 002</u>	Feb 06, 1998
AA +	<u>5MG</u>	<u>A084364 001</u>	
AA WEST WARD	<u>2MG</u>	<u>A040337 002</u>	Feb 16, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 409 (of 424)

TRIHEXYPHENIDYL HYDROCHLORIDE

TABLET; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

<u>AA</u>	WEST WARD	<u>5MG</u>	<u>A040337 001</u>	Feb 16, 2000
-----------	-----------	------------	--------------------	--------------

TRIMETHADIONE

TABLET; ORAL

TRIDIIONE

+ ABBOTT 150MG

N005856 009

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TIGAN

<u>AB</u>	+ KING PHARMS	<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>	MUTUAL PHARMA	<u>300MG</u>	<u>A076570 001</u>	Aug 28, 2003
-----------	---------------	--------------	--------------------	--------------

INJECTABLE; INJECTION

TIGAN

<u>AP</u>	+ JHP PHARMS	<u>100MG/ML</u>	<u>N017530 001</u>
-----------	--------------	-----------------	--------------------

TRIMETHOBENZAMIDE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>100MG/ML</u>	<u>A088804 001</u>	Apr 03, 1987
-----------	---------	-----------------	--------------------	--------------

<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>	NOVEL LABS INC	<u>100MG</u>	<u>A091437 001</u>	Jun 15, 2011
-----------	----------------	--------------	--------------------	--------------

<u>AB</u>	TEVA	<u>100MG</u>	<u>N018679 001</u>	Jul 30, 1982
-----------	------	--------------	--------------------	--------------

<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>A070049 001</u>	Jun 06, 1985
-----------	-------------	--------------	--------------------	--------------

TRIMETHOPRIM

+ TEVA	200MG		A071259 001	Jun 18, 1987
--------	-------	--	-------------	--------------

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

+ FSC	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>
-----------	---	----------------------	--------------------

TRIMIPRAMINE MALEATE

<u>AB</u>	MIKAH PHARMA	<u>EQ 25MG BASE</u>	<u>A077361 001</u>
-----------	--------------	---------------------	--------------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077361 002</u>
-----------	--	---------------------	--------------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077361 003</u>
-----------	--	----------------------	--------------------

TRIPTORELIN PAMOATE

INJECTABLE; INTRAMUSCULAR

TRELSTAR

+ WATSON LABS	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
---	---------------------	--	-------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 410 (of 424)

TROMETHAMINE

INJECTABLE; INJECTION
 THAM
 + HOSPIRA 3.6GM/100ML N013025 002

TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC
MYDRIACYL
AT + ALCON 0.5% A084305 001
AT + 1% A084306 001
TROPICACYL
AT AKORN 0.5% A040314 001 Sep 29, 2000
AT 1% A040315 001 Sep 29, 2000
TROPICAMIDE
AT BAUSCH AND LOMB 0.5% A040067 001 Jul 27, 1994
AT 1% A040064 001 Jul 27, 1994

TROSPiUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
 SANCTURA XR
 + ALLERGAN 60MG N022103 001 Aug 03, 2007
 TABLET; ORAL
SANCTURA
AB + ALLERGAN 20MG N021595 001 May 28, 2004
TROSPiUM CHLORIDE
AB APOTEX 20MG A091513 001 Dec 06, 2011
AB GLENMARK GENERICS 20MG A091575 001 Aug 13, 2010
AB PADDICK LLC 20MG A091573 001 Nov 17, 2010

TRYPAN BLUE

SOLUTION; OPHTHALMIC
 MEMBRANEBLUE
 + DORC 0.15% N022278 001 Feb 20, 2009
 VISIONBLUE
 + DORC 0.06% N021670 001 Dec 16, 2004

ULIPRISTAL ACETATE

TABLET; ORAL
 ELLA
 + LAB HRA PHARMA 30MG N022474 001 Aug 13, 2010

UREA C-13

FOR SOLUTION; ORAL
 BREATHTEK UBT FOR H-PYLORI
 + OTSUKA AMERICA EQ 75MG /POUCH N020586 002 May 10, 2001

UREA, C-14

CAPSULE; ORAL
 PYTEST
 + AVENT 1uCi N020617 001 May 09, 1997
 PYTEST KIT
 + AVENT 1uCi N020617 002 May 09, 1997

UROFOLLITROPIN

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS
 BRAVELLE
 + FERRING 75 IU/VIAL N021289 001 May 06, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 411 (of 424)

URSODIOL

CAPSULE; ORAL

ACTIGALL

<u>AB</u>	+ WATSON PHARMS	<u>300MG</u>	<u>N019594</u>	<u>002</u>	Dec 31, 1987
<u>URSODIOL</u>					
<u>AB</u>	COREPHARMA	<u>300MG</u>	<u>A077895</u>	<u>001</u>	Jul 27, 2006
<u>AB</u>	EPIC PHARMA	<u>300MG</u>	<u>A075517</u>	<u>001</u>	Mar 14, 2000
<u>AB</u>	LANNETT	<u>300MG</u>	<u>A079082</u>	<u>001</u>	Dec 15, 2008
<u>AB</u>	MYLAN	<u>300MG</u>	<u>A090530</u>	<u>001</u>	Feb 17, 2010
<u>AB</u>	TEVA PHARMS	<u>300MG</u>	<u>A075592</u>	<u>001</u>	May 25, 2000

TABLET; ORAL

URSO 250

<u>AB</u>	APITALIS PHARMA US	<u>250MG</u>	<u>N020675</u>	<u>001</u>	Dec 10, 1997
<u>URSO FORTE</u>					
<u>AB</u>	+ APTALIS PHARMA US	<u>500MG</u>	<u>N020675</u>	<u>002</u>	Jul 21, 2004
<u>URSODIOL</u>					
<u>AB</u>	GLENMARK GENERICS	<u>250MG</u>	<u>A090801</u>	<u>001</u>	Jul 12, 2011
<u>AB</u>		<u>500MG</u>	<u>A090801</u>	<u>002</u>	Jul 12, 2011
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A079184</u>	<u>001</u>	May 13, 2009
<u>AB</u>		<u>500MG</u>	<u>A079184</u>	<u>002</u>	May 13, 2009
<u>AB</u>	WATSON LABS INC	<u>250MG</u>	<u>A200826</u>	<u>001</u>	Dec 23, 2011
<u>AB</u>		<u>500MG</u>	<u>A200826</u>	<u>002</u>	Dec 23, 2011

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

<u>AB</u>	ACTAVIS PHARMA	<u>EQ 500MG BASE</u>	<u>A090370</u>	<u>001</u>	Mar 16, 2011
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090370</u>	<u>002</u>	Mar 16, 2011
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A090682</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090682</u>	<u>002</u>	May 24, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 500MG BASE</u>	<u>A079012</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A079012</u>	<u>002</u>	May 24, 2010
<u>AB</u>	MATRIX LABS LTD	<u>EQ 500MG BASE</u>	<u>A078518</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A078518</u>	<u>002</u>	May 24, 2010
<u>AB</u>	MYLAN	<u>EQ 500MG BASE</u>	<u>A078070</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A078070</u>	<u>002</u>	May 24, 2010
<u>AB</u>	RANBAXY	<u>EQ 500MG BASE</u>	<u>A076588</u>	<u>001</u>	Jan 31, 2007
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A076588</u>	<u>002</u>	Jan 31, 2007
<u>AB</u>	ROXANE	<u>EQ 500MG BASE</u>	<u>A078656</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A078656</u>	<u>002</u>	May 24, 2010
<u>AB</u>	SANDOZ	<u>EQ 500MG BASE</u>	<u>A077478</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A077478</u>	<u>002</u>	May 24, 2010
<u>AB</u>	TEVA PHARMS	<u>EQ 500MG BASE</u>	<u>A077655</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A077655</u>	<u>002</u>	May 24, 2010
<u>AB</u>	WATSON LABS	<u>EQ 500MG BASE</u>	<u>A077135</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A077135</u>	<u>002</u>	May 24, 2010
<u>AB</u>	WOCKHARDT	<u>EQ 500MG BASE</u>	<u>A090216</u>	<u>001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090216</u>	<u>002</u>	May 24, 2010
<u>VALTREX</u>					
<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 500MG BASE</u>	<u>N020487</u>	<u>001</u>	Jun 23, 1995
<u>AB</u>	+	<u>EQ 1GM BASE</u>	<u>N020487</u>	<u>002</u>	Jun 23, 1995

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION; ORAL

VALCYTE

<u>AB</u>	+ ROCHE PALO	<u>50MG/ML</u>	<u>N022257</u>	<u>001</u>	Aug 28, 2009
-----------	--------------	----------------	----------------	------------	--------------

TABLET; ORAL

VALCYTE

<u>AB</u>	+ ROCHE PALO	<u>EQ 450MG BASE</u>	<u>N021304</u>	<u>001</u>	Mar 29, 2001
-----------	--------------	----------------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 412 (of 424)

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPACON

<u>AP</u>	+ ABBOTT	<u>EQ 100MG BASE/ML</u>	<u>N020593 001</u>	Dec 30, 1996
<u>VALPROATE SODIUM</u>				
<u>AP</u>	APP PHARMS	<u>EQ 100MG BASE/ML</u>	<u>A076539 001</u>	Jun 26, 2003
<u>AP</u>	BEDFORD	<u>EQ 100MG BASE/ML</u>	<u>A076295 001</u>	Nov 14, 2002

HIKMA FARMACEUTICAEQ 100MG BASE/MLA078523 001

Feb 17, 2010

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

<u>AB</u>	+ ABBOTT	<u>250MG</u>	<u>N018081 001</u>	
<u>VALPROIC ACID</u>				
<u>AB</u>	BANNER PHARMACAPS	<u>250MG</u>	<u>A073484 001</u>	Jun 29, 1993

ABCATALENTA073229 001

Oct 29, 1991

CAPSULE, DELAYED RELEASE; ORAL

STAVZOR

BANNER PHARMACAPS	125MG	
	250MG	
+	500MG	

N022152 001 Jul 29, 2008

N022152 002 Jul 29, 2008

N022152 003 Jul 29, 2008

SYRUP; ORAL

DEPAKENE

<u>AA</u>	+ ABBOTT	<u>250MG/5ML</u>	<u>N018082 001</u>	
<u>VALPROIC ACID</u>				
<u>AA</u>	ALPHARMA	<u>250MG/5ML</u>	<u>A075782 001</u>	Dec 22, 2000
<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>	SUN PHARM IND'S INC	<u>250MG/5ML</u>	<u>A090517 001</u>	May 28, 2010
<u>AA</u>	TEVA PHARMS	<u>250MG/5ML</u>	<u>A073178 001</u>	Aug 25, 1992
<u>AA</u>	VINTAGE	<u>250MG/5ML</u>	<u>A077960 001</u>	Oct 13, 2006
<u>AA</u>	WOCKHARDT	<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

TABLET; ORAL

DIOVAN

NOVARTIS	40MG	
	80MG	
	160MG	
+	320MG	

N021283 004 Aug 14, 2002

N021283 001 Jul 18, 2001

N021283 002 Jul 18, 2001

N021283 003 Jul 18, 2001

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOCIN HYDROCHLORIDE

VIROPHARMA	<u>EQ 125MG BASE</u>	<u>N050606 001</u>	Apr 15, 1986
+	<u>EQ 250MG BASE</u>	<u>N050606 002</u>	Apr 15, 1986

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	+ APP PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A062663 001</u>	Mar 17, 1987
<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>	BIONICHE PHARMA USA	<u>EQ 500MG BASE/VIAL</u>	<u>A065401 001</u>	Jun 30, 2008
		<u>EQ 1GM BASE/VIAL</u>	<u>A065401 002</u>	Jun 30, 2008

PRESCRIPTION DRUG PRODUCT LIST

3 - 413 (of 424)

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	+ HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A062911</u>	<u>001</u>	Aug 04, 1988
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>A062931</u>	<u>001</u>	Oct 29, 1992
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062912</u>	<u>002</u>	Jan 07, 2009
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062933</u>	<u>002</u>	May 27, 2009
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>A062912</u>	<u>001</u>	Aug 04, 1988
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>A062933</u>	<u>001</u>	Oct 29, 1992
<u>AP</u>	+	<u>EQ 5GM BASE/VIAL</u>	<u>A063076</u>	<u>001</u>	Dec 21, 1990
<u>AP</u>	HOSPIRA INC	<u>EQ 10GM BASE/VIAL</u>	<u>A065455</u>	<u>001</u>	Apr 29, 2009
<u>AP</u>	PFIZER	<u>EQ 500MG BASE/VIAL</u>	<u>A065397</u>	<u>001</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065397</u>	<u>002</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A065432</u>	<u>001</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A091469</u>	<u>001</u>	Jul 01, 2011
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A090250</u>	<u>001</u>	Apr 27, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090250</u>	<u>002</u>	Apr 27, 2010
<u>AP</u>	STRIDES ARCOLAB LTD	<u>EQ 10GM BASE/VIAL</u>	<u>A091554</u>	<u>001</u>	Sep 19, 2011
	VANCOCIN HYDROCHLORIDE	IN PLASTIC CONTAINER			
+ BAXTER HLTHCARE		EQ 500MG BASE/100ML	N050671	001	Apr 29, 1993
+		EQ 750MG BASE/150ML	N050671	002	Dec 20, 2010

VANDETANIB

TABLET; ORAL

VANDETANIB

IPR PHARMS INC	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

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>	+ BEDFORD	<u>10MG/VIAL</u>	<u>A075549</u>	<u>001</u>	Jun 13, 2000
<u>AP</u>	+	<u>20MG/VIAL</u>	<u>A075549</u>	<u>002</u>	Jun 13, 2000
<u>AP</u>	HOSPIRA	<u>10MG/VIAL</u>	<u>A075164</u>	<u>001</u>	Oct 21, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075164</u>	<u>002</u>	Oct 21, 1999
<u>AP</u>	MUSTAFA NEVZAT	<u>10MG/VIAL</u>	<u>A078274</u>	<u>001</u>	Dec 29, 2008
<u>AP</u>		<u>20MG/VIAL</u>	<u>A078274</u>	<u>002</u>	Dec 29, 2008
<u>AP</u>	PFIZER	<u>10MG/VIAL</u>	<u>A090243</u>	<u>001</u>	May 11, 2010
<u>AP</u>		<u>20MG/VIAL</u>	<u>A090243</u>	<u>002</u>	May 11, 2010
<u>AP</u>	SUN PHARMA GLOBAL	<u>10MG/VIAL</u>	<u>A079001</u>	<u>001</u>	Jun 17, 2009
<u>AP</u>		<u>20MG/VIAL</u>	<u>A079001</u>	<u>002</u>	Jun 17, 2009
<u>AP</u>	TEVA PARENTERAL	<u>10MG/VIAL</u>	<u>A074688</u>	<u>001</u>	Aug 25, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 414 (of 424)

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>	TEVA PARENTERAL	<u>20MG/VIAL</u>	<u>A074688</u>	<u>002</u>	Aug 25, 1999
<u>AP</u>	WATSON LABS	<u>10MG/VIAL</u>	<u>A074334</u>	<u>001</u>	Aug 31, 1995
<u>AP</u>		<u>20MG/VIAL</u>	<u>A074334</u>	<u>002</u>	Aug 31, 1995

VELAGLUCERASE ALFA

INJECTABLE; IV (INFUSION)

SHIRE HUMAN GENETIC 400 UNITS/VIAL

N022575 001 Feb 26, 2010

VEMURAFENIB

TABLET; ORAL

ZELBORAF

+ HOFFMANN LA ROCHE 240MG

N202429 001 Aug 17, 2011

VENLAFAKINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

<u>AB</u>	WYETH PHARMS INC	<u>EQ 37.5MG BASE</u>	<u>N020699</u>	<u>001</u>	Oct 20, 1997
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N020699</u>	<u>002</u>	Oct 20, 1997
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N020699</u>	<u>004</u>	Oct 20, 1997
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 37.5MG BASE</u>	<u>A200834</u>	<u>001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A200834</u>	<u>002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A200834</u>	<u>003</u>	Apr 14, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 37.5MG BASE</u>	<u>A078421</u>	<u>001</u>	May 06, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078421</u>	<u>002</u>	May 06, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078421</u>	<u>003</u>	May 06, 2011
<u>AB</u>	MYLAN	<u>EQ 37.5MG BASE</u>	<u>A078789</u>	<u>001</u>	Jun 01, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078789</u>	<u>002</u>	Jun 01, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078789</u>	<u>003</u>	Jun 01, 2011
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 37.5MG BASE</u>	<u>A091123</u>	<u>001</u>	Jul 11, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091123</u>	<u>002</u>	Jul 11, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091123</u>	<u>003</u>	Jul 11, 2011
<u>AB</u>	TEVA	<u>EQ 37.5MG BASE</u>	<u>A076565</u>	<u>001</u>	Jun 28, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A076565</u>	<u>002</u>	Jun 28, 2010
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A076565</u>	<u>003</u>	Jun 28, 2010
<u>AB</u>	TORRENT PHARMS LLC	<u>EQ 37.5MG BASE</u>	<u>A090899</u>	<u>001</u>	Jun 01, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090899</u>	<u>002</u>	Jun 01, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090899</u>	<u>003</u>	Jun 01, 2011
<u>AB</u>	VALEANT INTL	<u>EQ 37.5MG BASE</u>	<u>A090071</u>	<u>001</u>	Apr 15, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090071</u>	<u>002</u>	Apr 15, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090071</u>	<u>003</u>	Apr 15, 2011
<u>AB</u>	WOCKHARDT	<u>EQ 37.5MG BASE</u>	<u>A078865</u>	<u>001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078865</u>	<u>002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A078865</u>	<u>003</u>	Apr 14, 2011
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 37.5MG BASE</u>	<u>A090174</u>	<u>001</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090174</u>	<u>002</u>	Apr 14, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090174</u>	<u>003</u>	Apr 14, 2011

TABLET; ORAL

VENLAFAKINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS TOTOWA	<u>EQ 25MG BASE</u>	<u>A078554</u>	<u>001</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078554</u>	<u>002</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078554</u>	<u>003</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078554</u>	<u>004</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078554</u>	<u>005</u>	Jan 09, 2009
<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<u>A078932</u>	<u>001</u>	Dec 14, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078932</u>	<u>002</u>	Dec 14, 2010

PRESCRIPTION DRUG PRODUCT LIST

3 - 415 (of 424)

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

AB	ALEMBIC PHARMS LTD	<u>EQ 50MG BASE</u>	<u>A078932</u>	<u>003</u>	Dec 14, 2010
AB		<u>EQ 75MG BASE</u>	<u>A078932</u>	<u>004</u>	Dec 14, 2010
AB		<u>EQ 100MG BASE</u>	<u>A078932</u>	<u>005</u>	Dec 14, 2010
AB	AMNEAL PHARMS	<u>EQ 25MG BASE</u>	<u>A079098</u>	<u>001</u>	May 11, 2010
AB		<u>EQ 37.5MG BASE</u>	<u>A079098</u>	<u>002</u>	May 11, 2010
AB		<u>EQ 50MG BASE</u>	<u>A079098</u>	<u>003</u>	May 11, 2010
AB		<u>EQ 75MG BASE</u>	<u>A079098</u>	<u>004</u>	May 11, 2010
AB		<u>EQ 100MG BASE</u>	<u>A079098</u>	<u>005</u>	May 11, 2010
AB	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A090555</u>	<u>001</u>	Apr 07, 2010
AB		<u>EQ 37.5MG BASE</u>	<u>A090555</u>	<u>002</u>	Apr 07, 2010
AB		<u>EQ 50MG BASE</u>	<u>A090555</u>	<u>003</u>	Apr 07, 2010
AB		<u>EQ 75MG BASE</u>	<u>A090555</u>	<u>004</u>	Apr 07, 2010
AB		<u>EQ 100MG BASE</u>	<u>A090555</u>	<u>005</u>	Apr 07, 2010
AB	CARACO	<u>EQ 25MG BASE</u>	<u>A090555</u>	<u>001</u>	Jun 13, 2008
AB		<u>EQ 37.5MG BASE</u>	<u>A090555</u>	<u>002</u>	Jun 13, 2008
AB		<u>EQ 50MG BASE</u>	<u>A090555</u>	<u>003</u>	Jun 13, 2008
AB		<u>EQ 75MG BASE</u>	<u>A090555</u>	<u>004</u>	Jun 13, 2008
AB		<u>EQ 100MG BASE</u>	<u>A090555</u>	<u>005</u>	Jun 13, 2008
AB	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A078627</u>	<u>001</u>	Jun 13, 2008
AB		<u>EQ 37.5MG BASE</u>	<u>A078627</u>	<u>002</u>	Jun 13, 2008
AB		<u>EQ 50MG BASE</u>	<u>A078627</u>	<u>003</u>	Jun 13, 2008
AB		<u>EQ 75MG BASE</u>	<u>A078627</u>	<u>004</u>	Jun 13, 2008
AB		<u>EQ 100MG BASE</u>	<u>A078627</u>	<u>005</u>	Jun 13, 2008
AB	MYLAN	<u>EQ 25MG BASE</u>	<u>A078301</u>	<u>001</u>	Jun 13, 2008
AB		<u>EQ 37.5MG BASE</u>	<u>A078301</u>	<u>002</u>	Jun 13, 2008
AB		<u>EQ 50MG BASE</u>	<u>A078301</u>	<u>003</u>	Jun 13, 2008
AB		<u>EQ 75MG BASE</u>	<u>A078301</u>	<u>004</u>	Jun 13, 2008
AB		<u>EQ 100MG BASE</u>	<u>A078301</u>	<u>005</u>	Jun 13, 2008
AB	TEVA	<u>EQ 25MG BASE</u>	<u>A077166</u>	<u>001</u>	Jun 13, 2008
AB		<u>EQ 37.5MG BASE</u>	<u>A077166</u>	<u>002</u>	Jun 13, 2008
AB		<u>EQ 50MG BASE</u>	<u>A077166</u>	<u>003</u>	Jun 13, 2008
AB		<u>EQ 75MG BASE</u>	<u>A077166</u>	<u>004</u>	Jun 13, 2008
AB		<u>EQ 100MG BASE</u>	<u>A077166</u>	<u>005</u>	Jun 13, 2008
AB	+ VINTAGE	<u>EQ 25MG BASE</u>	<u>A076690</u>	<u>001</u>	Aug 03, 2006
AB		<u>EQ 37.5MG BASE</u>	<u>A076690</u>	<u>002</u>	Aug 03, 2006
AB		<u>EQ 50MG BASE</u>	<u>A076690</u>	<u>003</u>	Aug 03, 2006
AB		<u>EQ 75MG BASE</u>	<u>A076690</u>	<u>004</u>	Aug 03, 2006
AB		<u>EQ 100MG BASE</u>	<u>A076690</u>	<u>005</u>	Aug 03, 2006
AB	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A090027</u>	<u>001</u>	Aug 04, 2010
AB		<u>EQ 37.5MG BASE</u>	<u>A090027</u>	<u>002</u>	Aug 04, 2010
AB		<u>EQ 50MG BASE</u>	<u>A090027</u>	<u>003</u>	Aug 04, 2010
AB		<u>EQ 75MG BASE</u>	<u>A090027</u>	<u>004</u>	Aug 04, 2010
AB		<u>EQ 100MG BASE</u>	<u>A090027</u>	<u>005</u>	Aug 04, 2010
AB		<u>EQ 25MG BASE</u>	<u>A077653</u>	<u>001</u>	Jun 13, 2008
AB		<u>EQ 37.5MG BASE</u>	<u>A077653</u>	<u>002</u>	Jun 13, 2008
AB		<u>EQ 50MG BASE</u>	<u>A077653</u>	<u>003</u>	Jun 13, 2008
AB		<u>EQ 75MG BASE</u>	<u>A077653</u>	<u>004</u>	Jun 13, 2008
AB		<u>EQ 100MG BASE</u>	<u>A077653</u>	<u>005</u>	Jun 13, 2008

TABLET, EXTENDED RELEASE; ORAL

VENLAFAXINE HYDROCHLORIDE

AB	OSMOTICA PHARM	<u>EQ 37.5MG BASE</u>	<u>N022104</u>	<u>001</u>	May 20, 2008
AB		<u>EQ 75MG BASE</u>	<u>N022104</u>	<u>002</u>	May 20, 2008
AB	+	<u>EQ 150MG BASE</u>	<u>N022104</u>	<u>003</u>	May 20, 2008
AB	SUN PHARMA GLOBAL	<u>EQ 37.5MG BASE</u>	<u>A091272</u>	<u>001</u>	Aug 18, 2010
AB		<u>EQ 75MG BASE</u>	<u>A091272</u>	<u>002</u>	Aug 18, 2010
AB		<u>EQ 150MG BASE</u>	<u>A091272</u>	<u>003</u>	Aug 18, 2010

VENLAFAXINE HYDROCHLORIDE

OSMOTICA PHARM	EQ 225MG BASE	<u>N022104</u>	<u>004</u>	May 20, 2008
----------------	---------------	----------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 416 (of 424)

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>100MG</u>	<u>A078306</u> <u>001</u>	Aug 09, 2007
<u>AB</u>		<u>120MG</u>	<u>A075138</u> <u>001</u>	Apr 20, 1999
<u>AB</u>		<u>180MG</u>	<u>A075138</u> <u>002</u>	Apr 20, 1999
<u>AB</u>		<u>200MG</u>	<u>A078306</u> <u>002</u>	Aug 09, 2007
<u>AB</u>		<u>240MG</u>	<u>A075138</u> <u>003</u>	Apr 20, 1999
<u>AB</u>		<u>300MG</u>	<u>A078306</u> <u>003</u>	Aug 09, 2007

VERELAN

<u>AB</u>	ALKERMES GAINESVILLE	<u>120MG</u>	<u>N019614</u> <u>001</u>	May 29, 1990
<u>AB</u>		<u>180MG</u>	<u>N019614</u> <u>003</u>	Jan 09, 1992
<u>AB</u>		<u>240MG</u>	<u>N019614</u> <u>002</u>	May 29, 1990

VERELAN PM

<u>AB</u>	ELAN DRUG	<u>100MG</u>	<u>N020943</u> <u>001</u>	Nov 25, 1998
<u>AB</u>		<u>200MG</u>	<u>N020943</u> <u>002</u>	Nov 25, 1998
<u>AB</u>	+	<u>300MG</u>	<u>N020943</u> <u>003</u>	Nov 25, 1998

VERELAN

+ ALKERMES GAINESVILLE	360MG		N019614 004	May 10, 1996
------------------------	-------	--	-------------	--------------

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>2.5MG/ML</u>	<u>A070737</u> <u>001</u>	May 06, 1987
<u>AP</u>		<u>2.5MG/ML</u>	<u>A070738</u> <u>001</u>	May 06, 1987
<u>AP</u>		<u>2.5MG/ML</u>	<u>A075136</u> <u>001</u>	Oct 20, 1998
<u>AP</u>	INTL MEDICATION	<u>2.5MG/ML</u>	<u>A070451</u> <u>001</u>	Dec 16, 1985

TABLET; ORAL

CALAN

<u>AB</u>	GD SEARLE LLC	<u>40MG</u>	<u>N018817</u> <u>003</u>	Feb 23, 1988
<u>AB</u>		<u>80MG</u>	<u>N018817</u> <u>001</u>	Sep 10, 1984
<u>AB</u>	+	<u>120MG</u>	<u>N018817</u> <u>002</u>	Sep 10, 1984

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>80MG</u>	<u>A071482</u> <u>001</u>	Feb 15, 1989
<u>AB</u>		<u>120MG</u>	<u>A071483</u> <u>001</u>	Feb 15, 1989
<u>AB</u>	WATSON LABS	<u>40MG</u>	<u>A072923</u> <u>001</u>	Jun 29, 1993
<u>AB</u>		<u>40MG</u>	<u>A072924</u> <u>001</u>	Jun 29, 1993
<u>AB</u>		<u>80MG</u>	<u>A070855</u> <u>001</u>	Sep 24, 1986
<u>AB</u>		<u>80MG</u>	<u>A070995</u> <u>001</u>	Oct 01, 1986
<u>AB</u>		<u>80MG</u>	<u>A071366</u> <u>001</u>	Oct 01, 1986
<u>AB</u>		<u>120MG</u>	<u>A070856</u> <u>001</u>	Sep 24, 1986
<u>AB</u>		<u>120MG</u>	<u>A070994</u> <u>001</u>	Oct 01, 1986
<u>AB</u>		<u>120MG</u>	<u>A071367</u> <u>001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE; ORAL

I索托平 SR

<u>AB</u>	PFIZER	<u>120MG</u>	<u>N019152</u> <u>003</u>	Mar 06, 1991
<u>AB</u>	+	<u>180MG</u>	<u>N019152</u> <u>002</u>	Dec 15, 1989
<u>AB</u>	+	<u>240MG</u>	<u>N019152</u> <u>001</u>	Dec 16, 1986

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	GLENMARK GENERICS	<u>120MG</u>	<u>A090700</u> <u>001</u>	Aug 03, 2011
<u>AB</u>		<u>180MG</u>	<u>A090700</u> <u>002</u>	Aug 03, 2011
<u>AB</u>		<u>240MG</u>	<u>A078906</u> <u>001</u>	Sep 17, 2009
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>120MG</u>	<u>A073568</u> <u>002</u>	Oct 10, 1997
<u>AB</u>		<u>180MG</u>	<u>A074330</u> <u>001</u>	Jan 31, 1994
<u>AB</u>		<u>240MG</u>	<u>A073568</u> <u>001</u>	Jul 31, 1992
<u>AB</u>	MYLAN	<u>120MG</u>	<u>A074587</u> <u>002</u>	Feb 21, 1997
<u>AB</u>		<u>180MG</u>	<u>A074587</u> <u>003</u>	Sep 09, 1997
<u>AB</u>		<u>240MG</u>	<u>A074587</u> <u>001</u>	Mar 23, 1996
<u>AB</u>	PAR PHARM	<u>120MG</u>	<u>A075072</u> <u>001</u>	May 25, 1999
<u>AB</u>		<u>240MG</u>	<u>A075072</u> <u>003</u>	May 25, 1999
<u>AB</u>	SUN PHARM INDs INC	<u>120MG</u>	<u>A090529</u> <u>001</u>	Dec 30, 2011
<u>AB</u>		<u>180MG</u>	<u>A090529</u> <u>002</u>	Dec 30, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 417 (of 424)

VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	SUN PHARM INDNS INC	<u>240MG</u>	<u>A090529</u>	<u>003</u>	Dec 30, 2011
	COVERA-HS				
BC +	GD SEARLE LLC	180MG	N020552	001	Feb 26, 1996
BC +		240MG	N020552	002	Feb 26, 1996

VERTEPORFIN

INJECTABLE; INJECTION

VISUDYNE

+ QLT 15MG/VIAL

N021119 001 Apr 12, 2000

VIGABATRIN

FOR SOLUTION; ORAL

SABRIL

+ LUNDBECK INC 500MG/PACKET

N022006 001 Aug 21, 2009

TABLET; ORAL

SABRIL

+ LUNDBECK INC 500MG

N020427 001 Aug 21, 2009

VILAZODONE HYDROCHLORIDE

TABLET; ORAL

VIIBRYD

FOREST LABS INC	10MG	N022567	001	Jan 21, 2011
	20MG	N022567	002	Jan 21, 2011
+	40MG	N022567	003	Jan 21, 2011

VINBLASTINE SULFATE

INJECTABLE; INJECTION

VINBLASTINE SULFATE

+ APP PHARMS 1MG/ML
+ BEDFORD 10MG/VIALA089515 001 Apr 29, 1987
A089395 001 Apr 09, 1987VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE PFS

<u>AP</u> + HOSPIRA	<u>1MG/ML</u>	<u>A071484</u>	<u>001</u>	Apr 19, 1988
<u>AP</u> TEVA PARENTERAL	<u>1MG/ML</u>	<u>A075493</u>	<u>001</u>	Sep 01, 1999

VINORELBINE TARTRATE

INJECTABLE; INJECTION

NAVELBINE

<u>AP</u> + PIERRE FABRE	<u>EQ 10MG BASE/ML</u>	<u>N020388</u>	<u>001</u>	Dec 23, 1994
--------------------------	------------------------	----------------	------------	--------------

VINORELBINE TARTRATE

<u>AP</u> ACTAVIS TOTOWA	<u>EQ 10MG BASE/ML</u>	<u>A078011</u>	<u>001</u>	Jul 22, 2009
<u>AP</u> APP PHARMS	<u>EQ 10MG BASE/ML</u>	<u>A076849</u>	<u>001</u>	Apr 18, 2005
<u>AP</u> BAXTER HLTHCARE	<u>EQ 10MG BASE/ML</u>	<u>A075992</u>	<u>001</u>	Jun 10, 2003
<u>AP</u> BEDFORD	<u>EQ 10MG BASE/ML</u>	<u>A076461</u>	<u>001</u>	Dec 11, 2003
<u>AP</u> EBEWE PHARMA	<u>EQ 10MG BASE/ML</u>	<u>A078408</u>	<u>001</u>	Feb 13, 2008
<u>AP</u> HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A076827</u>	<u>001</u>	Jun 02, 2005
<u>AP</u> TEVA PARENTERAL	<u>EQ 10MG BASE/ML</u>	<u>A076028</u>	<u>001</u>	Feb 03, 2003

VITAMIN A PALMITATE

INJECTABLE; INJECTION

AQUASOL A

+ HOSPIRA EQ 50,000 UNITS BASE/ML

N006823 001

PRESCRIPTION DRUG PRODUCT LIST

3 - 418 (of 424)

VORICONAZOLE

FOR SUSPENSION; ORAL

VFEND

+ PFIZER 200MG/5ML

N021630 001 Dec 19, 2003

INJECTABLE; IV (INFUSION)

+ PFIZER 200MG/VIAL

N021267 001 May 24, 2002

TABLET; ORAL

VFEND

<u>AB</u>	PFIZER	<u>50MG</u>	<u>N021266 001</u>	May 24, 2002
<u>AB</u>	+	<u>200MG</u>	<u>N021266 002</u>	May 24, 2002
VORICONAZOLE				
<u>AB</u>	MATRIX LABS LTD	<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>	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

VORINOSTAT

CAPSULE; ORAL

ZOLINZA

+ MERCK 100MG

N021991 001 Oct 06, 2006

WARFARIN SODIUM

INJECTABLE; INJECTION

COUMADIN

+ BRISTOL MYERS SQUIBB 5MG/VIAL

N009218 024 Feb 07, 1995

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>	BARR	<u>1MG</u>	<u>A040145 001</u>	Mar 26, 1997
<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>	INVAGEN PHARMS	<u>1MG</u>	<u>A090935 001</u>	May 25, 2011
<u>AB</u>		<u>2MG</u>	<u>A090935 002</u>	May 25, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 419 (of 424)

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A090935</u> <u>003</u>	May 25, 2011
<u>AB</u>		<u>3MG</u>	<u>A090935</u> <u>004</u>	May 25, 2011
<u>AB</u>		<u>4MG</u>	<u>A090935</u> <u>005</u>	May 25, 2011
<u>AB</u>		<u>5MG</u>	<u>A090935</u> <u>006</u>	May 25, 2011
<u>AB</u>		<u>6MG</u>	<u>A090935</u> <u>007</u>	May 25, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A090935</u> <u>008</u>	May 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A090935</u> <u>009</u>	May 25, 2011
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A040415</u> <u>001</u>	Sep 27, 2004
<u>AB</u>		<u>2MG</u>	<u>A040415</u> <u>002</u>	Sep 27, 2004
<u>AB</u>		<u>2.5MG</u>	<u>A040415</u> <u>003</u>	Sep 29, 2004
<u>AB</u>		<u>3MG</u>	<u>A040415</u> <u>004</u>	Sep 27, 2004
<u>AB</u>		<u>4MG</u>	<u>A040415</u> <u>005</u>	Sep 27, 2004
<u>AB</u>		<u>5MG</u>	<u>A040415</u> <u>006</u>	Sep 27, 2004
<u>AB</u>		<u>6MG</u>	<u>A040415</u> <u>007</u>	Sep 27, 2004
<u>AB</u>		<u>7.5MG</u>	<u>A040415</u> <u>008</u>	Sep 27, 2004
<u>AB</u>		<u>10MG</u>	<u>A040415</u> <u>009</u>	Sep 27, 2004
<u>AB</u>	PLIVA	<u>1MG</u>	<u>A040616</u> <u>009</u>	Jul 05, 2006
<u>AB</u>		<u>2MG</u>	<u>A040616</u> <u>001</u>	Jul 05, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040616</u> <u>002</u>	Jul 05, 2006
<u>AB</u>		<u>3MG</u>	<u>A040616</u> <u>003</u>	Jul 05, 2006
<u>AB</u>		<u>4MG</u>	<u>A040616</u> <u>004</u>	Jul 05, 2006
<u>AB</u>		<u>5MG</u>	<u>A040616</u> <u>005</u>	Jul 05, 2006
<u>AB</u>		<u>6MG</u>	<u>A040616</u> <u>006</u>	Jul 05, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040616</u> <u>007</u>	Jul 05, 2006
<u>AB</u>		<u>10MG</u>	<u>A040616</u> <u>008</u>	Jul 05, 2006
<u>AB</u>	TARO	<u>1MG</u>	<u>A040301</u> <u>002</u>	Jul 15, 1999
<u>AB</u>		<u>2MG</u>	<u>A040301</u> <u>003</u>	Jul 15, 1999
<u>AB</u>		<u>2.5MG</u>	<u>A040301</u> <u>004</u>	Jul 15, 1999
<u>AB</u>		<u>3MG</u>	<u>A040301</u> <u>005</u>	Jul 15, 1999
<u>AB</u>		<u>4MG</u>	<u>A040301</u> <u>006</u>	Jul 15, 1999
<u>AB</u>		<u>5MG</u>	<u>A040301</u> <u>007</u>	Jul 15, 1999
<u>AB</u>		<u>6MG</u>	<u>A040301</u> <u>008</u>	Jul 15, 1999
<u>AB</u>		<u>7.5MG</u>	<u>A040301</u> <u>009</u>	Jul 15, 1999
<u>AB</u>		<u>10MG</u>	<u>A040301</u> <u>001</u>	Jul 15, 1999
<u>AB</u>	ZYDUS PHARMS USA	<u>1MG</u>	<u>A040663</u> <u>001</u>	May 30, 2006
<u>AB</u>		<u>2MG</u>	<u>A040663</u> <u>002</u>	May 30, 2006
<u>AB</u>		<u>2.5MG</u>	<u>A040663</u> <u>003</u>	May 30, 2006
<u>AB</u>		<u>3MG</u>	<u>A040663</u> <u>004</u>	May 30, 2006
<u>AB</u>		<u>4MG</u>	<u>A040663</u> <u>005</u>	May 30, 2006
<u>AB</u>		<u>5MG</u>	<u>A040663</u> <u>006</u>	May 30, 2006
<u>AB</u>		<u>6MG</u>	<u>A040663</u> <u>007</u>	May 30, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A040663</u> <u>008</u>	May 30, 2006
<u>AB</u>		<u>10MG</u>	<u>A040663</u> <u>009</u>	May 30, 2006

XENON XE-133

GAS; INHALATION

XENON XE 133

<u>AA</u>	LANTHEUS MEDCL	<u>20mCi/VIAL</u>	<u>N017284</u> <u>002</u>
<u>AB</u>	XENON XE 133		
	LANTHEUS MEDCL	10mCi/VIAL	N017284 001

ZAFIRLUKAST

TABLET; ORAL

ACCOLATE

<u>AB</u>	ASTRAZENECA	<u>10MG</u>	<u>N020547</u> <u>003</u>
<u>AB</u>	+	<u>20MG</u>	Sep 17, 1999
			N020547 001
			Sep 26, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 420 (of 424)

ZAFIRLUKAST

TABLET; ORAL

ZAFIRLUKAST

<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A090372 001</u>	Nov 18, 2010
<u>AB</u>		<u>20MG</u>	<u>A090372 002</u>	Nov 18, 2010

ZALEPLON

CAPSULE; ORAL

SONATA

<u>AB</u>	KING PHARMS	<u>5MG</u>	<u>N020859 001</u>	Aug 13, 1999
<u>AB</u>	+	<u>10MG</u>	<u>N020859 002</u>	Aug 13, 1999

ZALEPLON

<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>	MYLAN	<u>5MG</u>	<u>A077238 001</u>	Jun 06, 2008
<u>AB</u>		<u>10MG</u>	<u>A077238 002</u>	Jun 06, 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>	ROXANE	<u>5MG</u>	<u>A077237 001</u>	Jun 06, 2008
<u>AB</u>		<u>10MG</u>	<u>A077237 002</u>	Jun 06, 2008
<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>	UPSHER SMITH	<u>5MG</u>	<u>A078706 001</u>	Jun 06, 2008
<u>AB</u>		<u>10MG</u>	<u>A078706 002</u>	Jun 06, 2008
<u>AB</u>	WEST WARD	<u>5MG</u>	<u>A078147 001</u>	Nov 25, 2008
<u>AB</u>		<u>10MG</u>	<u>A078147 002</u>	Nov 25, 2008

ZANAMIVIR

POWDER; INHALATION

RELENZA

+	GLAXOSMITHKLINE	5MG	<u>N021036 001</u>	Jul 26, 1999
---	-----------------	-----	--------------------	--------------

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

+	AZUR PHARMA II	100MCG/1ML (100MCG/ML)	<u>N021060 002</u>	Dec 28, 2004
+		500MCG/20ML (25MCG/ML)	<u>N021060 001</u>	Dec 28, 2004
+		500MCG/5ML (100MCG/ML)	<u>N021060 004</u>	Dec 28, 2004

ZIDOVUDINE

CAPSULE; ORAL

RETROVIR

<u>AB</u>	+	VIIV HLTHCARE	<u>100MG</u>	<u>N019655 001</u>	Mar 19, 1987
-----------	---	---------------	--------------	--------------------	--------------

ZIDOVUDINE

<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

RETROVIR

<u>AP</u>	+	VIIV HLTHCARE	<u>10MG/ML</u>	<u>N019951 001</u>	Feb 02, 1990
-----------	---	---------------	----------------	--------------------	--------------

ZIDOVUDINE

<u>AP</u>	LUITPOLD	<u>10MG/ML</u>	<u>A091457 001</u>	May 06, 2010
-----------	----------	----------------	--------------------	--------------

SYRUP; ORAL

RETROVIR

<u>AA</u>	+	VIIV HLTHCARE	<u>50MG/5ML</u>	<u>N019910 001</u>	Sep 28, 1989
-----------	---	---------------	-----------------	--------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 421 (of 424)

ZIDOVUDINE

SYRUP; ORAL

ZIDOVUDINE

<u>AA</u>	AUROBINDO	<u>50MG/5ML</u>	<u>A077268</u>	<u>001</u>	Sep 19, 2005
<u>AA</u>	CIPLA LTD	<u>50MG/5ML</u>	<u>A077981</u>	<u>001</u>	Jun 26, 2008

TABLET; ORAL

RETROVIR

<u>AB</u>	+ VIIV HLTHCARE	<u>300MG</u>	<u>N020518</u>	<u>002</u>	Oct 04, 1996
<u>AB</u>	<u>ZIDOVUDINE</u>				
<u>AB</u>	AUROBINDO	<u>300MG</u>	<u>A077267</u>	<u>001</u>	Sep 19, 2005
<u>AB</u>	CIPLA	<u>300MG</u>	<u>A090561</u>	<u>001</u>	Oct 27, 2010
<u>AB</u>	HEC PHARM INC	<u>300MG</u>	<u>A202058</u>	<u>001</u>	Oct 07, 2011
<u>AB</u>	HETERO DRUGS LTD	<u>300MG</u>	<u>A090092</u>	<u>001</u>	Apr 25, 2008
<u>AB</u>	MATRIX LABS LTD	<u>300MG</u>	<u>A078922</u>	<u>001</u>	Feb 14, 2008
<u>AB</u>	RANBAXY	<u>300MG</u>	<u>A077327</u>	<u>001</u>	Sep 19, 2005
<u>AB</u>	ROXANE	<u>300MG</u>	<u>A076844</u>	<u>001</u>	Sep 19, 2005

ZILEUTON

TABLET; ORAL

ZYFLO

+ CORNERSTONE THERAP 600MG

N020471 003 Dec 09, 1996

TABLET, EXTENDED RELEASE; ORAL

ZYFLO CR

+ CORNERSTONE THERAP 600MG

N022052 001 May 30, 2007

ZINC ACETATE

CAPSULE; ORAL

GALZIN

TEVA

EQ 25MG ZINC

N020458 001 Jan 28, 1997

+ EQ 50MG ZINC

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

+ PFIZER EQ 20MG BASE

N020825 001 Feb 05, 2001

EQ 40MG BASE

N020825 002 Feb 05, 2001

EQ 60MG BASE

N020825 003 Feb 05, 2001

EQ 80MG BASE

N020825 004 Feb 05, 2001

SUSPENSION; ORAL

GEODON

+ PFIZER INC EQ 10MG BASE/ML

N021483 001 Mar 29, 2006

ZIPRASIDONE MESYLATE

INJECTABLE; INTRAMUSCULAR

GEODON

+ PFIZER EQ 20MG BASE/ML

N020919 001 Jun 21, 2002

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

+ NOVARTIS EQ 5MG BASE/100ML

N021817 001 Apr 16, 2007

ZOMETA

+ NOVARTIS EQ 4MG BASE/5ML

N021223 002 Mar 07, 2003

+ EQ 4MG BASE/100ML

N021223 003 Jun 17, 2011

PRESCRIPTION DRUG PRODUCT LIST

3 - 422 (of 424)

ZOLMITRIPTAN

SPRAY; NASAL ZOMIG				
+ ASTRazeneca	5MG/SPRAY		N021450 004	Sep 30, 2003
TABLET; ORAL ZOMIG				
IPR	2.5MG		N020768 001	Nov 25, 1997
+	5MG		N020768 002	Nov 25, 1997
TABLET, ORALLY DISINTEGRATING; ORAL ZOMIG-ZMT				
ASTRAZeneca	2.5MG		N021231 001	Feb 13, 2001
+	5MG		N021231 002	Sep 17, 2001

ZOLPIDEM TARTRATE

SPRAY, METERED; ORAL ZOLPIMIST				
+ NOVADEL	5MG/SPRAY		N022196 001	Dec 19, 2008
TABLET; ORAL <u>AMBIEN</u>				
AB SANOFI AVENTIS US	<u>5MG</u>		<u>N019908 001</u>	Dec 16, 1992
AB +	<u>10MG</u>		<u>N019908 002</u>	Dec 16, 1992
<u>ZOLPIDEM TARTRATE</u>				
AB APOTEX INC	<u>5MG</u>		<u>A077884 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077884 002</u>	Apr 23, 2007
AB AUROBINDO PHARMA	<u>5MG</u>		<u>A078413 001</u>	May 04, 2007
AB	<u>10MG</u>		<u>A078413 002</u>	May 04, 2007
AB CARACO	<u>5MG</u>		<u>A077359 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077359 002</u>	Apr 23, 2007
AB CARLSBAD	<u>5MG</u>		<u>A077990 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077990 002</u>	Apr 23, 2007
AB DR REDDYS LABS LTD	<u>5MG</u>		<u>A077985 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077985 002</u>	Apr 23, 2007
AB HIKMA	<u>5MG</u>		<u>A078129 001</u>	Apr 30, 2008
AB	<u>10MG</u>		<u>A078129 002</u>	Apr 30, 2008
AB INVAGEN PHARMS	<u>5MG</u>		<u>A078184 001</u>	Sep 07, 2007
AB	<u>10MG</u>		<u>A078184 002</u>	Sep 07, 2007
AB LEK PHARMS DD	<u>5MG</u>		<u>A077322 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077322 002</u>	Apr 23, 2007
AB MYLAN	<u>5MG</u>		<u>A076578 001</u>	Apr 23, 2007
AB	<u>5MG</u>		<u>A078016 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A076578 002</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A078016 002</u>	Apr 23, 2007
AB RANBAXY	<u>5MG</u>		<u>A078055 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A078055 002</u>	Apr 23, 2007
AB ROXANE	<u>5MG</u>		<u>A077214 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077214 002</u>	Apr 23, 2007
AB TEVA	<u>5MG</u>		<u>A076410 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A076410 002</u>	Apr 23, 2007
AB TORRENT PHARMS	<u>5MG</u>		<u>A077903 001</u>	Aug 17, 2007
AB	<u>10MG</u>		<u>A077903 002</u>	Aug 17, 2007
AB VINTAGE	<u>5MG</u>		<u>A078616 001</u>	Nov 21, 2008
AB	<u>10MG</u>		<u>A078616 002</u>	Nov 21, 2008
AB WATSON LABS	<u>5MG</u>		<u>A077773 001</u>	Apr 23, 2007
AB	<u>10MG</u>		<u>A077773 002</u>	Apr 23, 2007
AB WOCKHARDT	<u>5MG</u>		<u>A078426 001</u>	May 15, 2007
AB	<u>10MG</u>		<u>A078426 002</u>	May 15, 2007
TABLET; SUBLINGUAL EDLUAR				
MEDA PHARMS	5MG		N021997 001	Mar 13, 2009

PRESCRIPTION DRUG PRODUCT LIST

3 - 423 (of 424)

ZOLPIDEM TARTRATE

TABLET; SUBLINGUAL EDLUAR			
+ MEDA PHARMS	10MG	N021997 002	Mar 13, 2009
INTERMEZZO			
PURDUE PHARMA	1.75MG	N022328 001	Nov 23, 2011
+	3.5MG	N022328 002	Nov 23, 2011
TABLET, EXTENDED RELEASE; ORAL			
<u>AMBIEN CR</u>			
AB SANOFI AVENTIS US	<u>6.25MG</u>	<u>N021774 002</u>	Sep 02, 2005
AB +	<u>12.5MG</u>	<u>N021774 001</u>	Sep 02, 2005
<u>ZOLPIDEM TARTRATE</u>			
AB ACTAVIS S ATLANTIC	<u>6.25MG</u>	<u>A078179 002</u>	Oct 13, 2010
AB ANCHEN PHARMS	<u>6.25MG</u>	<u>A078148 002</u>	Apr 14, 2011
AB	<u>12.5MG</u>	<u>A078148 001</u>	Dec 03, 2010
AB SANDOZ	<u>6.25MG</u>	<u>A090107 001</u>	Jul 01, 2011
AB	<u>12.5MG</u>	<u>A090107 002</u>	Jul 01, 2011
AB SYNTHON PHARMS	<u>6.25MG</u>	<u>A078483 001</u>	Apr 12, 2011
AB	<u>12.5MG</u>	<u>A078483 002</u>	Jun 06, 2011
<u>ZOLPIDEM TATRATE</u>			
AB ACTAVIS S ATLANTIC	<u>12.5MG</u>	<u>A078179 001</u>	Jun 06, 2011

ZONISAMIDE

CAPSULE; ORAL			
<u>ZONEGRAN</u>			
AB EISAI INC	<u>25MG</u>	<u>N020789 003</u>	Aug 22, 2003
AB	<u>50MG</u>	<u>N020789 002</u>	Aug 22, 2003
AB +	<u>100MG</u>	<u>N020789 001</u>	Mar 27, 2000
<u>ZONISAMIDE</u>			
AB ALPHAPHARM	<u>25MG</u>	<u>A077647 001</u>	Dec 22, 2005
AB	<u>50MG</u>	<u>A077647 002</u>	Dec 22, 2005
AB	<u>100MG</u>	<u>A077647 003</u>	Dec 22, 2005
AB APOTEX INC	<u>25MG</u>	<u>A077642 001</u>	Dec 22, 2005
AB	<u>50MG</u>	<u>A077642 002</u>	Dec 22, 2005
AB	<u>100MG</u>	<u>A077642 003</u>	Dec 22, 2005
AB BANNER PHARMACAPS	<u>25MG</u>	<u>A077813 001</u>	Aug 16, 2006
AB	<u>50MG</u>	<u>A077813 002</u>	Aug 16, 2006
AB	<u>100MG</u>	<u>A077813 003</u>	Aug 16, 2006
AB BARR	<u>25MG</u>	<u>A077639 001</u>	Dec 22, 2005
AB	<u>50MG</u>	<u>A077639 002</u>	Dec 22, 2005
AB	<u>100MG</u>	<u>A077639 003</u>	Dec 22, 2005
AB COREPHARMA	<u>25MG</u>	<u>A077876 001</u>	Feb 21, 2007
AB	<u>50MG</u>	<u>A077876 002</u>	Feb 21, 2007
AB	<u>100MG</u>	<u>A077876 003</u>	Feb 21, 2007
AB DR REDDYS LABS LTD	<u>25MG</u>	<u>A077645 002</u>	Sep 29, 2006
AB	<u>50MG</u>	<u>A077645 003</u>	Sep 29, 2006
AB	<u>100MG</u>	<u>A077645 001</u>	Dec 22, 2005
AB GLENMARK GENERICS	<u>25MG</u>	<u>A077651 001</u>	Jan 30, 2006
AB	<u>50MG</u>	<u>A077651 002</u>	Jan 30, 2006
AB	<u>100MG</u>	<u>A077651 003</u>	Jan 30, 2006
AB INVAGEN PHARMS	<u>25MG</u>	<u>A077869 001</u>	May 31, 2006
AB	<u>50MG</u>	<u>A077869 002</u>	May 31, 2006
AB	<u>100MG</u>	<u>A077869 003</u>	May 31, 2006
AB MYLAN	<u>25MG</u>	<u>A077637 001</u>	Dec 22, 2005
AB	<u>50MG</u>	<u>A077637 002</u>	Dec 22, 2005
AB	<u>100MG</u>	<u>A077637 003</u>	Dec 22, 2005
AB SANDOZ	<u>25MG</u>	<u>A077644 001</u>	Dec 22, 2005
AB	<u>50MG</u>	<u>A077644 002</u>	Dec 22, 2005
AB	<u>100MG</u>	<u>A077644 003</u>	Dec 22, 2005
AB SUN PHARM INDs (IN)	<u>25MG</u>	<u>A077634 001</u>	Mar 17, 2006

PRESCRIPTION DRUG PRODUCT LIST

3 - 424 (of 424)

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

<u>AB</u>	SUN PHARM INDs (IN)	<u>50MG</u>	<u>A077634</u>	<u>002</u>	Mar 17, 2006
<u>AB</u>		<u>100MG</u>	<u>A077634</u>	<u>003</u>	Mar 17, 2006
<u>AB</u>	WOCKHARDT	<u>25MG</u>	<u>A077636</u>	<u>003</u>	Jul 27, 2006
<u>AB</u>		<u>50MG</u>	<u>A077636</u>	<u>002</u>	Jul 27, 2006
<u>AB</u>		<u>100MG</u>	<u>A077636</u>	<u>001</u>	Dec 22, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077625</u>	<u>001</u>	Oct 16, 2006
<u>AB</u>		<u>50MG</u>	<u>A077625</u>	<u>002</u>	Oct 16, 2006
<u>AB</u>		<u>100MG</u>	<u>A077625</u>	<u>003</u>	Oct 16, 2006

OTC DRUG PRODUCT LIST

4 - 1 (of 19)

ACETAMINOPHEN

SUPPOSITORY; RECTAL				
ACEPHEN				
G AND W LABS	120MG	N018060	001	
	325MG	A072344	001	Mar 27, 1992
	325MG	N018060	003	Dec 18, 1986
	650MG	A072237	001	Mar 27, 1992
	650MG	N018060	002	
ACETAMINOPHEN				
ACTAVIS MID ATLANTIC	120MG	N018337	003	Sep 12, 1983
	325MG	N018337	002	
+ PERRIGO NEW YORK	650MG	N018337	001	
	120MG	A070607	001	Apr 06, 1987
	650MG	A070608	001	Dec 01, 1986
INFANTS' FEVERALL				
ACTAVIS MID ATLANTIC	80MG	N018337	004	Aug 26, 1992
NEOPAP				
POLYMEDICA	120MG	N016401	001	
TABLET, EXTENDED RELEASE; ORAL				
ACETAMINOPHEN				
OHM LABS	650MG	A076200	001	Mar 19, 2002
PERRIGO	650MG	A075077	001	Feb 25, 2000
RANBAXY LABS LTD	650MG	A078569	001	Dec 14, 2011
TYLENOL (CAPLET)				
+ MCNEIL CONS	650MG	N019872	001	Jun 08, 1994
TYLENOL (GELTAB)				
+ MCNEIL CONS	650MG	N019872	002	Jan 11, 2001

ACETAMINOPHEN; ASPIRIN; CAFFEINE

TABLET; ORAL				
ACETAMINOPHEN, ASPIRIN AND CAFFFEINE				
PERRIGO	250MG;250MG;65MG	A075794	001	Nov 26, 2001
EXCEDRIN (MIGRAINE)				
+ NOVARTIS	250MG;250MG;65MG	N020802	001	Jan 14, 1998

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL				
TAVIST ALLERGY/SINUS/HEADACHE				
+ NOVARTIS	500MG;EQ 0.25MG BASE;30MG	N021082	001	Mar 01, 2001

ACETAMINOPHEN; DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL				
DRIXORAL PLUS				
+ SCHERING PLOUGH	500MG;3MG;60MG	N019453	001	May 22, 1987

ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL				
AVAGARD				
+ 3M	61%;1%	N021074	001	Jun 07, 2001

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL				
FOAMCOAT				
GUARDIAN DRUG	80MG;20MG	A071793	001	Sep 04, 1987
GAVISCON				
SANOFI AVENTIS US	80MG;20MG	N018685	001	Dec 09, 1983
+	160MG;40MG	N018685	002	Dec 09, 1983

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

OTC DRUG PRODUCT LIST

4 - 2 (of 19)

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
+	2%;3%;10%;5%	N022009	002 Oct 29, 2009

AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL SHADE UVAGUARD			
+ SCHERING PLOUGH	3%;7.5%;3%	N020045	001 Dec 07, 1992

BENTOQUATAM

LOTION; TOPICAL IVY BLOCK			
+ STAND HOMEOPATH	5%	N020532	001 Aug 26, 1996

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL LOTRIMIN ULTRA			
+ SCHERING PLOUGH	1%	N021307	001 Dec 07, 2001

BUTOCONAZOLE NITRATE

CREAM; VAGINAL FEMSTAT 3			
+ BAYER	2%	N020421	001 Dec 21, 1995

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL CALCIUM CARBONATE, FAMOTIDINE AND MAGNESIUM HYDROXIDE			
PERRIGO R AND D	800MG;10MG;165MG	A077355	001 Feb 06, 2008
PEPCID COMPLETE			
+ MERCK SHARP DOHME	800MG;10MG;165MG	N020958	001 Oct 16, 2000

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL CETIRIZINE HYDROCHLORIDE ALLERGY			
BANNER PHARMACAPS	5MG	N022429	001 Jul 23, 2009
+	10MG	N022429	004 Jul 23, 2009
CETIRIZINE HYDROCHLORIDE HIVES RELIEF			
BANNER PHARMACAPS	5MG	N022429	003 Jul 23, 2009
+	10MG	N022429	002 Jul 23, 2009

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY			
AMNEAL PHARMS	5MG/5ML	A090765	002 Oct 07, 2009
APOTEX	5MG/5ML	A090188	002 Apr 22, 2008
AUROBINDO PHARMA	5MG/5ML	A090750	002 Feb 02, 2010
CYPRESS PHARM	5MG/5ML	A090300	001 Oct 10, 2008
DR REDDYS LABS LTD	5MG/5ML	A090474	002 Mar 30, 2009
PERRIGO R AND D	5MG/5ML	A090254	002 Apr 09, 2008
RANBAXY	5MG/5ML	A090183	002 Apr 24, 2008
SILARX	5MG/5ML	A091130	001 Apr 22, 2011
SUN PHARM INDs INC	5MG/5ML	A091327	001 Oct 17, 2011
TARO	5MG/5ML	A090182	002 Apr 22, 2008
	5MG/5ML	A201546	001 May 20, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS	5MG/5ML	A090765	001 Oct 07, 2009
APOTEX	5MG/5ML	A090188	001 Apr 22, 2008
AUROBINDO PHARMA	5MG/5ML	A090750	001 Feb 02, 2010
CYPRESS PHARM	5MG/5ML	A090300	002 Oct 10, 2008
DR REDDYS LABS LTD	5MG/5ML	A090474	001 Mar 30, 2009
PERRIGO R AND D	5MG/5ML	A090254	001 Apr 09, 2008
RANBAXY	5MG/5ML	A090183	001 Apr 24, 2008

OTC DRUG PRODUCT LIST

4 - 3 (of 19)

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SILARX	5MG/5ML	A091130	002	Apr 22, 2011
SUN PHARM INDs INC	5MG/5ML	A091327	002	Oct 17, 2011
TARO	5MG/5ML	A090182	001	Apr 22, 2008
	5MG/5ML	A201546	002	May 20, 2011

CHILDREN'S ZYRTEC ALLERGY

+ MCNEIL CONSUMER	5MG/5ML	N022155	002	Nov 16, 2007
+ MCNEIL CONSUMER	5MG/5ML	N022155	001	Nov 16, 2007

TABLET, CHEWABLE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

CARACO	5MG	A077631	004	Jan 11, 2008
	10MG	A077631	003	Jan 11, 2008
SANDOZ	5MG	A078692	001	Feb 14, 2008
	10MG	A078692	002	Feb 14, 2008

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

CARACO	5MG	A077631	001	Jan 11, 2008
	10MG	A077631	002	Jan 11, 2008

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

SUN PHARMA GLOBAL	5MG	A090142	001	Aug 30, 2011
	10MG	A090142	002	Aug 30, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARMA GLOBAL	5MG	A090142	003	Aug 30, 2011
	10MG	A090142	004	Aug 30, 2011

CHILDREN'S ZYRTEC ALLERGY

Pfizer	5MG	N021621	003	Nov 16, 2007
+ MCNEIL CONSUMER	10MG	N021621	004	Nov 16, 2007
CHILDREN'S ZYRTEC HIVES RELIEF				
Pfizer	5MG	N021621	005	Nov 16, 2007

+ 10MG

TABLET, ORALLY DISINTEGRATING; ORAL

ZYRTEC ALLERGY

+ MCNEIL CONSUMER	10MG	N022578	001	Sep 03, 2010
-------------------	------	---------	-----	--------------

TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

ACTAVIS ELIZABETH	5MG	A078615	003	Dec 28, 2007
	10MG	A078615	004	Dec 28, 2007

AMNEAL PHARMS NY	5MG	A078780	001	Jan 21, 2010
	10MG	A078780	004	Jan 21, 2010

APOTEX INC	5MG	A078317	001	Dec 27, 2007
	10MG	A078317	002	Dec 27, 2007

CADISTA PHARMS	5MG	A078933	001	Jun 15, 2010
	10MG	A078933	002	Jun 15, 2010

CARACO	5MG	A077499	001	Dec 27, 2007
	10MG	A077499	002	Dec 27, 2007

CONTRACT PHARMA	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

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

RANBAXY	5MG	A077498	001	Dec 27, 2007
	10MG	A077498	002	Dec 27, 2007

SANDOZ	5MG	A077946	001	Dec 27, 2007
	10MG	A077946	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

OTC DRUG PRODUCT LIST

4 - 4 (of 19)

CETIRIZINE HYDROCHLORIDE

TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

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				
CADISTA PHARMS	5MG	A078933	003	Jun 15, 2010
	10MG	A078933	004	Jun 15, 2010
CARACO	5MG	A077499	003	Dec 27, 2007
	10MG	A077499	004	Dec 27, 2007
DR REDDYS LABS LTD	5MG	A078343	001	Jan 15, 2008
	10MG	A078343	002	Jan 15, 2008
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
RANBAXY	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
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
ZYRTEC ALLERGY				
MCNEIL CONSUMER	5MG	N019835	003	Nov 16, 2007
+	10MG	N019835	004	Nov 16, 2007
ZYRTEC HIVES RELIEF				
MCNEIL CONSUMER	5MG	N019835	005	Nov 16, 2007
+	10MG	N019835	006	Nov 16, 2007

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA PHARMS	5MG;120MG	A077170	001	Feb 25, 2008
SANDOZ	5MG;120MG	A077991	001	Mar 05, 2008
ZYRTEC-D 12 HOUR				
+ MCNEIL	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

ECOLAB	2%	N019258	001	Jul 22, 1986
--------	----	---------	-----	--------------

OTC DRUG PRODUCT LIST

4 - 5 (of 19)

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL					
DYNA-HEX					
XTTRIUM	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
PHARMASEAL SCRUB CARE					
+ CAREFUSION	4%		N019793	001	Dec 02, 1988

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SPONGE; TOPICAL					
CHLORAPREP ONE-STEP					
+ CAREFUSION	2%;70% (3ML)		N020832	001	Jul 14, 2000
	2%;70% (10.5ML)		N020832	004	Aug 20, 2003
	2%;70% (26ML)		N020832	006	Nov 21, 2006
CHLORAPREP ONE-STEP FREPP					
+ CAREFUSION	2%;70% (1.5ML)		N020832	003	Apr 26, 2002
CHLORAPREP WITH TINT					
+ CAREFUSION	2%;70% (26ML)		N020832	002	May 03, 2005
	2%;70% (10.5ML)		N020832	005	Apr 03, 2006
	2%;70% (3ML)		N020832	007	Oct 10, 2006
SWAB; TOPICAL					
CHLORAPREP ONE-STEP SEPP					
+ CARDINAL HLTH	2%;70% (0.67ML)		N021555	001	Oct 07, 2002
CHLORAPREP SINGLE SWABSTICK					
+ CARDINAL HLTH	2%;70% (1.75ML)		N021555	002	May 10, 2005
CHLORAPREP TRIPLE SWABSTICK					
+ CARDINAL HLTH	2%;70% (5.25ML)		N021555	003	Jun 10, 2009
CHLORASCRUB MAXI SWABSTICK					
+ 3M INFECTION	3.15%;70% (5.1ML)		N021524	003	Jun 03, 2005
CHLORASCRUB SWAB					
+ 3M INFECTION	3.15%;70% (1ML)		N021524	001	Jun 03, 2005
CHLORASCRUB SWABSTICK					
+ 3M INFECTION	3.15%;70% (1.6ML)		N021524	002	Jun 03, 2005

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL					
CHLORPHENIRAMINE MALEATE					
AVANTHI INC	12MG		A040829	001	May 13, 2009
CHLOR-TRIMETON					
+ SCHERING PLOUGH	12MG		N007638	002	

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL					
ADVIL ALLERGY AND CONGESTION RELIEF					
+ PFIZER CONS HLTHCARE	4MG;200MG;10MG		N022113	001	Dec 21, 2011

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL					
CHILDREN'S ADVIL ALLERGY SINUS					
+ WYETH CONS	1MG/5ML;100MG/5ML;15MG/5ML		N021587	001	Feb 24, 2004
TABLET; ORAL					
ADVIL ALLERGY SINUS					
+ WYETH CONS	2MG;200MG;30MG		N021441	001	Dec 19, 2002

OTC DRUG PRODUCT LIST

4 - 6 (of 19)

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL
 CHLOR-TRIMETON
 + SCHERING PLOUGH 8MG;120MG N018397 001

CIMETIDINE

TABLET; ORAL
 CIMETIDINE
 APOTEX 100MG A074948 001 Jun 19, 1998
 200MG A074948 002 Jul 26, 2002
 CONTRACT PHARMACAL 200MG A074961 001 Jun 19, 1998
 200MG A074963 001 Jun 19, 1998
 IVAX SUB TEVA PHARMS 200MG A075345 001 Jun 16, 1999
 PERRIGO 200MG A075285 001 Oct 29, 1998
 WATSON LABS 200MG A075425 001 Jul 29, 1999
 TAGAMET HB
 + GLAXOSMITHKLINE 200MG N020238 002 Aug 21, 1996

CLEMASTINE FUMARATE

TABLET; ORAL
 CLEMASTINE FUMARATE
 PERRIGO 1.34MG A074512 001 Nov 22, 1995
 SANDOZ 1.34MG A073458 001 Oct 31, 1993
 TAVIST-1
 + NOVARTIS 1.34MG N020925 001 Aug 21, 1992

CLOTTRIMAZOLE

CREAM, TABLET; TOPICAL, VAGINAL
 GYNE-LOTRIMIN 3 COMBINATION PACK
 + SCHERING PLOUGH 1%,200MG N020526 002 Jul 29, 1996
 GYNE-LOTRIMIN COMBINATION PACK
 + SCHERING PLOUGH 1%,100MG N020289 002 Apr 26, 1993
 MYCELEX-7 COMBINATION PACK
 BAYER PHARMS 1%,100MG N020389 002 Jun 23, 1994
 CREAM; VAGINAL
 CLOTTRIMAZOLE
 ACTAVIS MID ATLANTIC 1% A074165 001 Jul 16, 1993
 TARO 1% A072641 001 Dec 04, 1995
 GYNE-LOTRIMIN
 + SCHERING PLOUGH 1% N018052 002 Nov 30, 1990
 GYNE-LOTRIMIN 3
 + SCHERING PLOUGH 2% N020574 001 Nov 24, 1998
 MYCELEX-7
 BAYER PHARMS 1% N018230 002 Dec 26, 1991
 TRIVAGIZOLE 3
 TARO 2% N021143 001 Apr 12, 2000
 TABLET; VAGINAL
 GYNE-LOTRIMIN
 + SCHERING PLOUGH 100MG N017717 002 Nov 30, 1990
 GYNE-LOTRIMIN 3
 + SCHERING PLOUGH 200MG N020525 001 Jul 29, 1996
 MYCELEX-7
 BAYER PHARMS 100MG N018182 002 Dec 26, 1991

CROMOLYN SODIUM

SPRAY, METERED; NASAL
 CROMOLYN SODIUM
 + BAUSCH AND LOMB 5.2MG/SPRAY A075702 001 Jul 03, 2001
 HH AND P 5.2MG/SPRAY A077976 001 Sep 07, 2007
 PERRIGO 5.2MG/SPRAY A075427 001 Dec 12, 2001

OTC DRUG PRODUCT LIST

4 - 7 (of 19)

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL DISOPHROL				
SCHERING PLOUGH	6MG;120MG	N013483	004	Sep 13, 1982
DRIXORAL + SCHERING PLOUGH	6MG;120MG	N013483	003	Sep 13, 1982

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL 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

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET; ORAL ADVIL PM				
+ PFIZER CONS HLTHCARE	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				
+ WYETH CONS	25MG;EQ 200MG FREE ACID AND POTASSIUM SALT	N021393	001	Dec 21, 2005
IBUPROFEN AND DIPHENHYDRAMINE				
BANNER PHARMACAPS	25MG;EQ 200MG FREE ACID AND POTASSIUM SALT	A090397	001	Nov 22, 2010

DOCOSANOL

CREAM; TOPICAL ABREVA				
+ GLAXOSMITHKLINE	10%	N020941	001	Jul 25, 2000

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	

EPINEPHRINE

AEROSOL, METERED; INHALATION EPINEPHRINE				
ARMSTRONG PHARMS	0.2MG/INH	A087907	001	May 23, 1984

FAMOTIDINE

TABLET, CHEWABLE; ORAL FAMOTIDINE				
PERRIGO	10MG	A075715	001	Aug 22, 2003
PEPCID AC				
+ MERCK SHARP DOHME	20MG	N020801	002	Dec 17, 2007
TABLET; ORAL FAMOTIDINE				
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

OTC DRUG PRODUCT LIST

4 - 8 (of 19)

FAMOTIDINE

TABLET; ORAL					
FAMOTIDINE					
MYLAN	10MG	A075674	001	Dec 21, 2001	
PERRIGO	10MG	A075400	001	Mar 18, 2005	
	20MG	A077351	001	Sep 25, 2006	
RANBAXY	10MG	A090283	001	Nov 17, 2009	
	20MG	A090283	002	Nov 17, 2009	
SANDOZ	10MG	A076101	001	Oct 21, 2002	
TEVA	10MG	A075312	001	May 31, 2001	
WATSON LABS	10MG	A075404	001	Nov 28, 2001	
WOCKHARDT	10MG	A077146	001	Mar 07, 2005	
	20MG	A090837	001	Aug 04, 2010	
PEPCID AC					
MERCK SHARP DOHME	10MG	N020325	001	Apr 28, 1995	
+	20MG	N020325	002	Sep 23, 2003	
PEPCID AC (GELTAB)					
MERCK SHARP DOHME	10MG	N020902	001	Aug 05, 1999	

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	
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	
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	
CHILDREN'S ALLEGRA HIVES					
SANOFI AVENTIS US	30MG	N020872	006	Jan 24, 2011	
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY					
DR REDDYS LABS LTD	30MG	A076502	004	Apr 12, 2011	
MYLAN	30MG	A077081	004	Jul 21, 2011	
TEVA	30MG	A076447	004	Apr 13, 2011	
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES					
DR REDDYS LABS LTD	30MG	A076502	005	Apr 12, 2011	
MYLAN	30MG	A077081	005	Jul 21, 2011	
TEVA	30MG	A076447	005	Apr 13, 2011	
FEXOFENADINE HYDROCHLORIDE ALLERGY					
DR REDDYS LABS LTD	60MG	A076502	006	Apr 12, 2011	
	180MG	A076502	008	Apr 12, 2011	
MYLAN	60MG	A077081	006	Jul 21, 2011	
	180MG	A077081	008	Jul 21, 2011	
TEVA	60MG	A076447	006	Apr 13, 2011	
	180MG	A076447	008	Apr 13, 2011	
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	
TEVA	60MG	A076447	007	Apr 13, 2011	
	180MG	A076447	009	Apr 13, 2011	

OTC DRUG PRODUCT LIST

4 - 9 (of 19)

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE 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 PSEUDOEPHEDRINE HYDROCHLORIDE DR REDDYS LABS LTD 180MG;240MG	A079043	002	Jun 22, 2011

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL GUAIFENESIN			
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 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			
+ WYETH CONS EQ 200MG FREE ACID AND POTASSIUM SALT	N020402	001	Apr 20, 1995
ADVIL MIGRAINE LIQUI-GELS			
+ WYETH CONS 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
BANNER PHARMACAPS EQ 200MG FREE ACID AND POTASSIUM SALT	A078682	001	Mar 24, 2009
+ CONTRACT PHARMACAL 200MG	A074782	001	Jul 06, 1998
DR REDDYS LABS LTD EQ 200MG FREE ACID AND POTASSIUM SALT	A077338	001	Jul 10, 2009
MARKSANS PHARMA EQ 200MG FREE ACID AND POTASSIUM SALT	A079205	001	Jun 26, 2009
MIDOL LIQUID GELS			
+ BANNER PHARMACAPS 200MG	N021472	001	Oct 18, 2002
SUSPENSION/DROPS; ORAL CHILDREN'S MOTRIN			
+ MCNEIL CONS 40MG/ML	N020603	001	Jun 10, 1996
IBUPROFEN			
PERRIGO 40MG/ML	A075217	001	Dec 16, 1998
TRIS PHARMA INC 40MG/ML	A079058	001	Aug 31, 2009
PEDIATRIC ADVIL			
+ WYETH CONS 100MG/2.5ML	N020812	001	Jan 30, 1998
SUSPENSION; ORAL CHILDREN'S ADVIL			
WYETH CONS 100MG/5ML	N020589	001	Jun 27, 1996
CHILDREN'S ADVIL-FLAVORED			
WYETH CONS 100MG/5ML	N020589	002	Nov 07, 1997
CHILDREN'S ELIXSURE			
ALTERNA TCHP LLC 100MG/5ML	N021604	001	Jan 07, 2004
CHILDREN'S IBUPROFEN			
PERRIGO 100MG/5ML	A074937	001	Dec 22, 1998
CHILDREN'S MOTRIN			
+ MCNEIL 100MG/5ML	N020516	001	Jun 16, 1995
IBUPROFEN			
ACTAVIS MID ATLANTIC 100MG/5ML	A074916	001	Apr 30, 1999
AMNEAL PHARMS 100MG/5ML	A200457	001	Aug 18, 2011
TABLET, CHEWABLE; ORAL CHILDREN'S ADVIL			
WYETH CONS 50MG	N020944	001	Dec 18, 1998
CHILDREN'S MOTRIN			
MCNEIL CONS 50MG	N020601	001	Nov 15, 1996

OTC DRUG PRODUCT LIST

4 - 10 (of 19)

IBUPROFEN

TABLET, CHEWABLE; ORAL

IBUPROFEN

PERRIGO	50MG	A076359	001	Jan 16, 2004
	100MG	A076359	002	Jan 16, 2004
JUNIOR STRENGTH ADVIL				
WYETH CONS	100MG	N020944	002	Dec 18, 1998
JUNIOR STRENGTH MOTRIN				
+ MCNEIL CONS	100MG	N020601	003	Nov 15, 1996
TABLET; ORAL				
ADVIL				
WYETH CONS	200MG	N018989	001	May 18, 1984
IBUPROFEN				
AMNEAL PHARMS NY	200MG	A071333	001	Feb 17, 1987
	200MG	A072199	001	May 23, 1988
AVEMA PHARMA	200MG	A076460	001	Nov 26, 2003
CONTRACT PHARMACAL	200MG	A071732	001	Sep 10, 1987
	200MG	A071735	001	Sep 10, 1987
	200MG	A072299	001	Jul 01, 1988
	200MG	A073691	001	Feb 25, 1994
	200MG	A074931	001	Jul 20, 1998
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
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
SVADS HOLDINGS SA	200MG	A079129	001	Mar 28, 2011
	200MG	A091355	001	Apr 04, 2011
VINTAGE PHARMS	200MG	A071229	001	Apr 01, 1987
	200MG	A071639	001	Feb 02, 1988
WATSON LABS	200MG	A070435	001	Mar 05, 1986
IBUPROFEN				
OHM LABS	200MG	A071214	001	Dec 01, 1986
IBU-TAB 200				
ALRA	200MG	A071057	001	Aug 11, 1988
JUNIOR STRENGTH ADVIL				
WYETH CONS	100MG	N020267	002	Dec 13, 1996
JUNIOR STRENGTH IBUPROFEN				
PERRIGO	100MG	A075367	001	Apr 22, 1999
JUNIOR STRENGTH MOTRIN				
MCNEIL CONS	100MG	N020602	001	Jun 10, 1996
MOTRIN IB				
+ MCNEIL	200MG	N019012	003	Dec 17, 1990
MOTRIN MIGRAINE PAIN				
+ MCNEIL	200MG	N019012	004	Feb 25, 2000
PROFEN				
CONTRACT PHARMACAL	200MG	A071265	001	Oct 15, 1986
TAB-PROFEN				
PERRIGO	200MG	A072095	001	Dec 08, 1987

IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET; ORAL

ADVIL CONGESTION RELIEF

+ WYETH CONS	200MG;10MG	N022565	001	May 27, 2010
--------------	------------	---------	-----	--------------

OTC DRUG PRODUCT LIST

4 - 11 (of 19)

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL ADVIL COLD AND SINUS				
+ WYETH CONS	EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG	N021374	001	May 30, 2002
SUSPENSION; ORAL CHILDREN'S ADVIL COLD				
WYETH CONS	100MG/5ML; 15MG/5ML	N021373	001	Apr 18, 2002
CHILDREN'S MOTRIN COLD				
+ MCNEIL CONS	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				
+ WYETH CONS	200MG; 30MG	N019771	001	Sep 19, 1989
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE				
CONTRACT PHARMACAL	200MG; 30MG	A075588	001	Apr 08, 2002
DR REDDYS LABS LTD	200MG; 30MG	A077628	001	Aug 14, 2006
IBUPROPHM COLD AND SINUS				
OHM LABS	200MG; 30MG	A074567	001	Apr 17, 2001
SINE-AID IB				
MCNEIL CONS	200MG; 30MG	N019899	001	Dec 31, 1992

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION HUMULIN R				
+ LILLY	100 UNITS/ML	N018780	001	Oct 28, 1982
HUMULIN R PEN				
+ LILLY	100 UNITS/ML	N018780	005	Aug 06, 1998
NOVOLIN R				
+ NOVO NORDISK INC	100 UNITS/ML	N019938	001	Jun 25, 1991

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION HUMULIN 70/30				
+ LILLY	30 UNITS/ML; 70 UNITS/ML	N019717	001	Apr 25, 1989
HUMULIN 70/30 PEN				
+ LILLY	30 UNITS/ML; 70 UNITS/ML	N019717	002	Aug 06, 1998
NOVOLIN 70/30				
+ NOVO NORDISK INC	30 UNITS/ML; 70 UNITS/ML	N019991	001	Jun 25, 1991

INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION HUMULIN N				
+ LILLY	100 UNITS/ML	N018781	001	Oct 28, 1982
NOVOLIN N				
+ NOVO NORDISK INC	100 UNITS/ML	N019959	001	Jul 01, 1991

IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE; TOPICAL DURAPREP				
+ 3M	EQ 0.7% IODINE; 74% (6ML)	N021586	001	Sep 29, 2006
+	EQ 0.7% IODINE; 74% (26ML)	N021586	002	Sep 29, 2006

KETOCONAZOLE

SHAMPOO; TOPICAL NIZORAL A-D				
+ MCNEIL CONS	1%	N020310	001	Oct 10, 1997

KETOPROFEN

FILM; ORAL NEXCEDE				
+ NOVARTIS	12.5MG	N022470	001	Nov 25, 2009

OTC DRUG PRODUCT LIST

4 - 12 (of 19)

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC ALAWAY				
+ BAUSCH AND LOMB	EQ 0.025% BASE	N021996	001	Dec 01, 2006
KETOTIFEN FUMARATE				
AKORN	EQ 0.025% BASE	A077958	001	Jul 26, 2007
ALCON PHARMS LTD	EQ 0.025% BASE	A077200	001	Sep 02, 2008
APOTEX INC	EQ 0.025% BASE	A077354	001	May 09, 2006
ZADITOR				
+ ALCON PHARMA	EQ 0.025% BASE	N021066	002	Oct 19, 2006

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL PREVACID 24 HR				
+ NOVARTIS	15MG	N022327	001	May 18, 2009

LEVONORGESTREL

TABLET; ORAL LEVONORGESTREL				
PERRIGO R AND D	0.75MG	A090740	001	Dec 30, 2010
+ WATSON LABS	0.75MG	A078665	001	Aug 28, 2009
PLAN B				
+ TEVA WOMENS	0.75MG	N021045	002	Aug 24, 2006
PLAN B ONE-STEP				
+ DURAMED	1.5MG	N021998	001	Jul 10, 2009

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL LOPERAMIDE HYDROCHLORIDE				
BANNER PHARMACAPS	1MG	N021855	001	Aug 04, 2005
+	2MG	N021855	002	Aug 04, 2005
SOLUTION; ORAL IMODIUM A-D				
+ MCNEIL CONS	1MG/5ML	N019487	001	Mar 01, 1988
LOPERAMIDE HYDROCHLORIDE				
HI TECH PHARMA	1MG/5ML	A074352	001	Nov 17, 1995
PERRIGO	1MG/5ML	A073243	001	Jan 21, 1992
ROXANE	1MG/5ML	A073079	001	Apr 30, 1992
WOCKHARDT	1MG/5ML	A074730	001	Aug 28, 1997
SUSPENSION; ORAL IMODIUM A-D				
+ MCNEIL CONS	1MG/7.5ML	N019487	002	Jul 08, 2004
LOPERAMIDE HYDROCHLORIDE				
PERRIGO R AND D	1MG/7.5ML	A091292	001	May 20, 2011
TABLET, CHEWABLE; ORAL IMODIUM A-D EZ CHEWS				
+ MCNEIL	2MG	N020448	001	Jul 24, 1997
TABLET; ORAL IMODIUM A-D				
+ MCNEIL CONS	2MG	N019860	001	Nov 22, 1989
LOPERAMIDE HYDROCHLORIDE				
CONTRACT PHARMACAL	2MG	A073254	001	Jul 30, 1993
LNK	2MG	A076497	001	Jun 10, 2003
OHM LABS	2MG	A074091	001	Dec 10, 1992
PERRIGO	2MG	A075232	001	Jan 06, 2000

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET, CHEWABLE; ORAL IMODIUM MULTI-SYMPOTOM RELIEF				
+ MCNEIL	2MG;125MG	N020606	001	Jun 26, 1996
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE				
PERRIGO	2MG;125MG	A076029	001	Aug 30, 2002

OTC DRUG PRODUCT LIST

4 - 13 (of 19)

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET; ORAL				
IMODIUM MULTI-SYMPOTM RELIEF				
+ MCNEIL CONS	2MG;125MG	N021140	001	Nov 30, 2000
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE				
RANBAXY	2MG;125MG	A077500	001	Sep 06, 2006

LORATADINE

CAPSULE; ORAL				
CLARITIN				
+ SCHERING PLOUGH	10MG	N021952	001	Jun 16, 2008
SUSPENSION; ORAL				
LORATADINE				
+ TARO	1MG/ML	N021734	001	Oct 04, 2005
SYRUP; ORAL				
CLARITIN				
+ SCHERING PLOUGH	1MG/ML	N020641	002	Nov 27, 2002
LORATADINE				
APOTEX INC	1MG/ML	A075565	001	Oct 05, 2004
PERRIGO	1MG/ML	A075728	001	Aug 20, 2004
RANBAXY	1MG/ML	A076529	001	Aug 20, 2004
SILARX	1MG/ML	A077421	001	Jun 29, 2006
TARO	1MG/ML	A076805	001	Aug 20, 2004
TEVA	1MG/ML	A075505	001	Nov 07, 2003
WOCKHARDT	1MG/ML	A075815	001	Aug 20, 2004
TABLET, CHEWABLE; ORAL				
CHILDREN'S CLARITIN				
+ SCHERING PLOUGH	5MG	N021891	001	Aug 23, 2006
TABLET, ORALLY DISINTEGRATING; ORAL				
ALAVERT				
WYETH CONS	10MG	N021375	001	Dec 19, 2002
CLARITIN HIVES RELIEF REDITAB				
+ SCHERING PLOUGH	10MG	N020704	003	Nov 19, 2003
CLARITIN REDITABS				
+ SCHERING PLOUGH	5MG	N021993	001	Dec 12, 2006
+	10MG	N020704	002	Nov 27, 2002
LORATADINE				
IMPAK LABS	10MG	A076011	001	Sep 29, 2003
WATSON LABS FLORIDA	10MG	A075990	001	Nov 03, 2003
WYETH CONS	10MG	A075822	001	Feb 10, 2003
LORATADINE REDIDOSE				
RANBAXY	10MG	A077153	001	Apr 11, 2007
TABLET; ORAL				
CLARITIN				
+ SCHERING PLOUGH	10MG	N019658	002	Nov 27, 2002
CLARITIN HIVES RELIEF				
+ SCHERING PLOUGH	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
RANBAXY	10MG	A076134	001	Aug 18, 2003
SANDOZ	10MG	A075209	001	Jan 21, 2003

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL				
CLARITIN-D				
+ SCHERING PLOUGH	5MG;120MG	N019670	002	Nov 27, 2002
CLARITIN-D 24 HOUR				
+ SCHERING PLOUGH	10MG;240MG	N020470	002	Nov 27, 2002
LORATADINE AND PSEUDOEPHEDRINE SULFATE				
IMPAK LABS	5MG;120MG	A076050	001	Jan 30, 2003
	10MG;240MG	A075989	001	Mar 04, 2004

OTC DRUG PRODUCT LIST

4 - 14 (of 19)

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

LORATADINE AND PSEUDOEPHEDRINE SULFATE
 RANBAXY 10MG;240MG
 WATSON LABS FLORIDA 5MG;120MG
 10MG;240MG

A076557 001 Sep 22, 2004
 A076208 001 Jan 28, 2004
 A075706 001 Feb 21, 2003

MENTHOL; METHYL SALICYLATE

PATCH; TOPICAL

SALONPAS

+ HISAMITSU 3%;10%

N022029 001 Feb 20, 2008

MICONAZOLE NITRATE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

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

+ INSIGHT PHARMS 2%,1.2GM

N021308 001 Jun 29, 2001

MONISTAT 7 COMBINATION PACK

+ INSIGHT PHARMS 2%,100MG

N020288 002 Apr 26, 1993

MONISTAT-3 COMBINATION PACK

+ INSIGHT PHARMS 2%,200MG

N020670 002 Apr 16, 1996

M-ZOLE 3 COMBINATION PACK

ACTAVIS MID ATLANTIC 2%,200MG

A074926 001 Apr 16, 1999

CREAM; TOPICAL, VAGINAL

MICONAZOLE 3 COMBINATION PACK

PERRIGO 2%,4%

A076357 001 Mar 30, 2004

MONISTAT 3 COMBINATION PACK

INSIGHT PHARMS 2%,4%

N021261 003 Jun 17, 2003

MONISTAT 3 COMBINATION PACK (PREFILLED)

+ INSIGHT PHARMS 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 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

+ INSIGHT PHARMS 4%

N020827 001 Mar 30, 1998

MONISTAT 7

+ INSIGHT PHARMS 2%

N017450 002 Feb 15, 1991

SUPPOSITORIY; VAGINAL

MICONAZOLE NITRATE

ACTAVIS MID ATLANTIC 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

+ INSIGHT PHARMS 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

OTC DRUG PRODUCT LIST

4 - 15 (of 19)

MINOXIDIL

SOLUTION; TOPICAL				
MINOXIDIL (FOR MEN)				
ACTAVIS MID ATLANTIC	2%	A074588	001	Apr 05, 1996
HI TECH PHARMA	2%	A074731	001	Dec 24, 1996
NOVEX	2%	A074924	001	Apr 29, 1998
PERRIGO	2%	A075357	001	Jul 30, 1999
WOCKHARDT	2%	A074767	001	Feb 28, 1997
MINOXIDIL (FOR WOMEN)				
HI TECH PHARMA	2%	A074731	002	May 11, 2005
NOVEX	2%	A074924	002	Apr 29, 1998
PERRIGO	2%	A075357	002	Jul 30, 1999
MINOXIDIL EXTRA STRENGTH (FOR MEN)				
ACTAVIS MID ATLANTIC	5%	A075518	001	Nov 17, 2000
AVACOR PRODS	5%	A075619	001	Nov 17, 2000
NOVEX	5%	A075839	001	Oct 01, 2001
PERRIGO	5%	A075598	001	Jun 13, 2001
PERRIGO NEW YORK	5%	A075737	001	Mar 15, 2002
WOCKHARDT	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
THEROXIDIL				
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				
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				
+ BANNER PHARMACAPS	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
CONTRACT PHARMACAL	EQ 200MG BASE	A074635	001	Jan 13, 1997
	EQ 200MG BASE	A074789	001	Feb 27, 1997
DR REDDYS LABS INC	EQ 200MG BASE	A075168	001	Jul 28, 1998
GRANULES INDIA	EQ 200MG BASE	A091353	001	Sep 20, 2011
MARKSANS PHARMA	EQ 200MG BASE	A090545	001	Mar 16, 2011
PERRIGO	EQ 200MG BASE	A074661	001	Jan 13, 1997
RANBAXY LABS LTD	EQ 200MG BASE	A091183	001	May 20, 2011
SANDOZ	EQ 200MG BASE	A074646	001	Jan 13, 1997

NAPROXEN SODIUM; PSEUDOEPHENDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL				
ALEVE-D SINUS & COLD				
+ BAYER	200MG;120MG	N021076	001	Nov 29, 1999
NAPROXEN SODIUM AND PSEUDOEPHENDRINE HYDROCHLORIDE				
DR REDDYS LABS INC	EQ 220MG BASE;120MG	A077381	001	Sep 27, 2006

OTC DRUG PRODUCT LIST

4 - 16 (of 19)

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE
PERRIGO EQ 200MG BASE;120MG

A076518 001 Mar 17, 2004

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

HABITROL

+ NOVARTIS	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	A074645	001	Oct 20, 1997
	14MG/24HR	A074611	001	Oct 20, 1997
	21MG/24HR	A074612	001	Oct 20, 1997

NICOTINE POLACRILEXGUM, 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

IVAX SUB TEVA PHARMS	EQ 2MG BASE	A076880	001	Feb 18, 2009
	EQ 4MG BASE	A077850	001	Feb 18, 2009
PERRIGO	EQ 2MG BASE	A076775	001	Sep 16, 2004
	EQ 2MG BASE	A076776	001	Sep 16, 2004
	EQ 4MG 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

PERRIGO R AND D	EQ 2MG 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 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	A078325	001	Oct 30, 2006

WATSON LABS	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	A079044	001	Jul 08, 2009
	EQ 2MG BASE	A079216	001	Jul 08, 2009

	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

THRIVE				
NOVARTIS	EQ 2MG BASE	A077658	001	Jun 19, 2007
	EQ 4MG BASE	A077656	001	Jun 19, 2007

TROCHE/LOZENGE; ORAL				
COMMIT				
GLAXOSMITHKLINE CONS	EQ 2MG BASE	N021330	001	Oct 31, 2002

OTC DRUG PRODUCT LIST

4 - 17 (of 19)

NICOTINE POLACRILEX

TROCHE/LOZENGE; ORAL					
NICORETTE					
+ GLAXOSMITHKLINE CONS	EQ 2MG BASE		N022360	001	May 18, 2009
	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 4MG BASE		A077007	002	Jan 31, 2006
	EQ 4MG BASE		A090711	002	Jul 10, 2009
	EQ 4MG BASE		A090821	002	Jul 10, 2009

NIZATIDINE

TABLET; ORAL					
AXID AR					
+ WYETH CONS	75MG		N020555	001	May 09, 1996

NONOXYNOL-9

SPONGE; VAGINAL					
TODAY					
+ AZTIQ PHARMA	1GM		N018683	001	Apr 01, 1983

OMEPRAZOLE

TABLET, DELAYED RELEASE; ORAL					
OMEPRAZOLE					
+ DEXCEL PHARMA	20MG		N022032	001	Dec 04, 2007

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE; ORAL					
OMEPRAZOLE MAGNESIUM					
+ DR REDDYS LABS LTD	EQ 20MG BASE		A078878	001	Jun 05, 2009
TABLET, DELAYED RELEASE; ORAL					
+ PRILOSEC OTC					
+ ASTRazeneca	EQ 20MG BASE		N021229	001	Jun 20, 2003

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE; ORAL					
ZEGERID OTC					
+ MSD CONSUMER	20MG; 1.1GM		N022281	001	Dec 01, 2009

ORLISTAT

CAPSULE; ORAL					
ALLI					
+ GLAXOSMITHKLINE CONS	60MG		N021887	001	Feb 07, 2007

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC					
OCUCLEAR					
+ SCHERING PLOUGH	0.025%		N018471	001	May 30, 1986
VISINE L.R.					
+ JOHNSON AND JOHNSON	0.025%		N019407	001	Mar 31, 1989

PERMETHRIN

LOTION; TOPICAL					
NIX					
+ INSIGHT PHARMS	1%		N019918	001	May 02, 1990
PERMETHRIN					
+ ACTAVIS MID ATLANTIC	1%		A075014	001	Mar 28, 2000
PERRIGO NEW YORK	1%		A076090	001	Dec 20, 2001

OTC DRUG PRODUCT LIST

4 - 18 (of 19)

PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL; TOPICAL RID MOUSSE + BAYER HLTHCARE	4%;EQ 0.33% BASE	N021043 001	Mar 07, 2000
--	------------------	-------------	--------------

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL GLYCOLAX			
KREMERS URBAN PHARMS	17GM/PACKET 17GM/SCOOPFUL	A090600 001 A090600 002	Oct 06, 2009 Oct 06, 2009
MIRALAX			
+ SCHERRING PLOUGH	17GM/SCOOPFUL	N022015 001	Oct 06, 2006
POLYETHYLENE GLYCOL 3350			
MYLAN	17GM/PACKET 17GM/SCOOPFUL	A078915 001 A078915 002	Oct 06, 2009 Oct 06, 2009
NEXGEN PHARMA	17GM/SCOOPFUL	A090812 001	Oct 07, 2009
NOVEL LABS INC	17GM/SCOOPFUL	A091077 001	Oct 06, 2009
PADDOCK LLC	17GM/SCOOPFUL	A090567 001	Oct 15, 2009
PERRIGO R AND D	17GM/PACKET 17GM/SCOOPFUL	A090685 001 A090685 002	Oct 06, 2009 Oct 06, 2009

POTASSIUM IODIDE

SOLUTION; ORAL THYROSHIELD			
+ FLEMING	65MG/ML	A077218 001	Jan 12, 2005
TABLET; ORAL IOSAT			
ANBEX	65MG	N018664 002	May 12, 2011
+ THYROSafe	130MG	N018664 001	Oct 14, 1982
+ 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			
PERRIGO	120MG	A075153 001	Feb 26, 1999
RANBAXY	120MG	A077442 001	Sep 28, 2005
SUDAFED 12 HOUR			
+ MCNEIL CONS	120MG	A073585 001	Oct 31, 1991
SUDAFED 24 HOUR			
+ MCNEIL CONS	240MG	N020021 002	Dec 15, 1992

PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL AFRINOL			
+ SCHERRING PLOUGH	120MG	N018191 001	

PURIFIED WATER

SOLUTION; OPHTHALMIC PUR-WASH			
+ NIAGARA PHARMS	98.3%	N022305 001	Sep 01, 2011

OTC DRUG PRODUCT LIST

4 - 19 (of 19)

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

CONTRACT PHARMACAL	EQ 75MG BASE	A075094	001	Jun 21, 1999
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
SVADS HOLDINGS SA	EQ 150MG BASE	A200536	001	Jun 28, 2011
TORPHARM	EQ 75MG BASE	A075167	001	May 04, 2000
WATSON LABS	EQ 75MG BASE	A075212	001	Jan 14, 2000
WOCKHARDT	EQ 75MG BASE	A076760	001	Feb 24, 2006
	EQ 75MG BASE	A078884	001	Jul 31, 2008
	EQ 150MG BASE	A078653	001	Nov 26, 2007
ZANTAC 150				
+ BOEHRINGER INGELHEIM	EQ 150MG BASE	N021698	001	Aug 31, 2004
	EQ 150MG BASE	N021698	002	Mar 13, 2007
ZANTAC 75				
BOEHRINGER INGELHEIM	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

+ NOVARTIS	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

+ NOVARTIS	6 .5%	N020676	001	Feb 11, 1997
------------	-------	---------	-----	--------------

**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		
MEDSEP CORPORATION	N760305	Jun 30, 1978

ANTICOAGULANT CITRATE DEXTROSE SOLUTION (ACD)

INJECTABLE; INJECTION		
CITRA LABS LLC	N020037	Aug 26, 2003
ACD-A SOLUTION		
GAMBRO BCT INC	A010228	ANDA Feb 25, 2002
ADSOL WITH ACD-A		
FENWAL INC	N000922	Aug 29, 2002
ANTICOAGULANT CITRATE DEXTROSE SOLUTION FORMULA A		
HAEMONETICS CORP	A980728	ANDA Feb 06, 2002
AS3 SOLUTION/ACD-A		
GAMBRO BCT INC	N001214	May 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION		
NONE		
FENWAL INC	N100855	Jun 03, 1959
	N160918	Mar 17, 1978
FRESENIUS MEDICAL CARE NORTH AMERICA	N110912	Sep 02, 1959
MEDSEP CORPORATION	A710497	ANDA
MILES INC	N100102	Dec 14, 1961

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		
FRESENIUS MEDICAL CARE NORTH AMERICA	N780519	Apr 23, 1980
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

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

INJECTABLE; INJECTION		
NONE		
MEDSEP CORPORATION	N800077	Nov 06, 1980

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION

INJECTABLE; INJECTION		
ADSOL IN PLASTIC CONTAINER		
FENWAL INC	N900223	Dec 27, 1991

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

INJECTABLE; INJECTION

MACOPRODUCTIONS SAS CPD/AS-1: MACOPHARMA LEUCOFLEX MTL1 LEUKOREDUCTION SYSTEM FOR BLOOD
COMPONENTS KNOWN AS MTL1-WB

MACOPRODUCTIONS SAS	N040083	Nov 21, 2005
NONE		
GAMBRO BCT INC	A070025	Jan 06, 2008

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC	N170401	Dec 06, 1977
FRESENIUS MEDICAL CARE NORTH AMERICA	N811012	Jun 28, 1983
MEDSEP CORPORATION	N160907	May 16, 1973
MILES INC	N800222	Aug 23, 1982
TERUMO MEDICAL CORP	N160527	Jun 22, 1970
	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
MEDSEP CORP 0.042GM/100ML; 0.276GM/100ML;
0.410GM/100ML; 0.30GM/100ML;
1.10GM/100ML; 0.588GM/100ML N820915 Oct 19, 1984

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 ETHYLENEDIAMINE TETRAACETIC ACID (EDTA)

INJECTABLE; INJECTION

NONE
FENWAL INC N090480 Mar 07, 1955

ANTICOAGULANT HEPARIN SOLUTION USP

INJECTABLE; INJECTION

NONE
FRESENIUS MEDICAL
CARE NORTH AMERICA N770822 May 17, 1978

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

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
FRESENIUS MEDICAL
CARE NORTH AMERICA
TERUMO MEDICAL CORP

N770923

Jan 20, 1978

N160702

Dec 28, 1970

N781214

Feb 08, 1980

DEXTRAN 1 IN SODIUM CHLORIDE 0.6%

INJECTABLE; INJECTION

PROMIT

MEDA AB

N830715

Oct 30, 1984

DEXTRAN 40, 10% IN DEXTROSE 5%

INJECTABLE; INJECTION

GENTRAN 40

BAXTER HLTHCARE CORP	10GM/100ML;5GM/100ML	N840619	Feb 22, 1985
LMD IN PLASTIC CONTAINER			
HOSPIRA INC	10GM/100ML;5GM/100ML	A720563	ANDA Oct 30, 1992
NONE			
B BRAUN MEDICAL INC	10GM/100ML;5GM/100ML	N160767	Apr 06, 1970
HOSPIRA INC	10GM/100ML;5GM/100ML	N160375	Jul 25, 1967
MEDA AB	10GM/100ML;5GM/100ML	N140716	Jan 18, 1967
MILES INC	10GM/100ML;5GM/100ML	N160653	Sep 23, 1969
PHARMACHEM CORP	10GM/100ML;5GM/100ML	N160836	Nov 14, 1972

DEXTRAN 40, 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

GENTRAN 40

BAXTER HLTHCARE CORP	10GM/100ML;0.9GM/100ML	N840620	Feb 22, 1985
LMD IN PLASTIC CONTAINER			
HOSPIRA INC	10GM/100ML;0.9GM/100ML	A720562	ANDA Oct 30, 1992
NONE			
B BRAUN MEDICAL INC	10GM/100ML;0.9GM/100ML	N160767	Apr 06, 1970
HOSPIRA INC	10GM/100ML;0.9GM/100ML	N160375	Jul 25, 1967
MEDA AB	10GM/100ML;0.9GM/100ML	N140716	Jan 18, 1967
MILES INC	10GM/100ML;0.9GM/100ML	N160653	Sep 23, 1969
PHARMACHEM CORP	10GM/100ML;0.9GM/100ML	N160836	Nov 14, 1972

DEXTRAN 40, 10% W/V DEXTRAN 40 IN 0.8% NACL 500 ML PVC BAGS

INJECTABLE; INJECTION

RHEOMACRODEX

MEDA AB

N830527

Mar 27, 1985

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

DEXTRAN 40, 10% W/V DEXTRAN 40 IN 5% DEXTROSE 500 ML PVC BAGS

INJECTABLE; INJECTION
 RHEOMACRODEX
 MEDA AB N830627 Mar 27, 1985

DEXTRAN 70, 6% IN DEXTROSE 5%

INJECTABLE; INJECTION
 MACRODEX
 MEDA AB 6GM/100ML;5GM/100ML N060826 Jun 08, 1954

DEXTRAN 70, 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION
 GENTRAIN 70
 BAXTER HLTHCARE CORP N160607 Jan 26, 1970
 MACRODEX
 MEDA AB 6GM/100ML;5GM/100ML N060826 Jun 08, 1954
 NONE
 B BRAUN MEDICAL INC 6GM/100ML;0.9 GM/100ML N090024 Aug 18, 1969
 MILES INC 6GM/100ML;0.9 GM/100ML N080716 Mar 13, 1953

DEXTRAN 70, 6% W/V DEXTRAN 70 IN 0.9% NAACL IN 500 ML PVC BAGS

INJECTABLE; INJECTION
 MACRODEX
 MEDA AB N830613 Mar 27, 1985

DEXTRAN 70, 6% W/V DEXTRAN 70 IN 5% DEXTROSE

INJECTABLE; INJECTION
 MACRODEX
 MEDA AB N830629 Mar 27, 1985

DEXTRAN 75, 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION
 NONE
 PHARMACHEM CORP 6GM/100ML;0.9GM/100ML N080564 Sep 19, 1952
 6GM/100ML;0.9GM/100ML N160759 Aug 19, 1970

HETASTARCH 6% IN LACTATED ELECTROLYTE INJECTION

INJECTABLE; INJECTION
 HEXTEND
 BIOTIME INC 6GM/100ML N200952 Mar 31, 1999

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 ANDA Jan 30, 1995
 HESPA
 B BRAUN MEDICAL INC 6GM/100ML;0.9GM/100ML N160889 Jul 17, 1972
 HESPA IN PLASTIC CONTAINER
 B BRAUN MEDICAL INC 6GM/100ML;0.9GM/100ML N890105 Apr 04, 1991
 NONE
 B BRAUN MEDICAL INC 6GM/100ML;0.9GM/100ML A740283 ANDA Oct 21, 1998
 TEVA PARENTERAL 6GM/100ML;0.9GM/100ML A740592 ANDA Nov 12, 1998
 MEDICINES INC

HETASTARCH 6% IN SODIUM CHLORIDE 0.9% NAACL 500 ML GLASS BOTTLES

INJECTABLE; INJECTION
 NONE
 HOSPIRA INC 6GM/100ML;0.9 GM/100ML A720746 ANDA Feb 07, 1996

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

HYDROXYETHYL STARCH 130/0.4 IN 6% SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

VOLUVEN

FRESENIUS KABI DEUTSCHLAND GMBH 6GM/100ML;0.9GM/100ML

N070012 NDA Dec 27, 2007

PENTASTARCH 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

PENTASPAK

B BRAUN MEDICAL INC 10GM/100ML;0.9GM/100ML
PENTASPAK IN PLASTIC CONTAINER

N841207 May 19, 1987

B BRAUN MEDICAL INC 10GM/100ML;0.9GM/100ML

N890104 Apr 04, 1991

PERFLUORODECALIN; PERFLUOROTRI-N-PROPYLAMINE

INJECTABLE; INJECTION

FLUOSOL

ALPHA THERAPEUTIC CORP 17.5GM/100ML;7.5GM/100ML

N860909 Dec 26, 1989

RED BLOOD CELL PROCESSING SOLUTION

INJECTABLE; INJECTION

REJUVESOL

CITRA LABS LLC

N950522 Feb 26, 1997

SODIUM CHLORIDE; SODIUM ACETATE; SODIUM CITRATE DIHYDRATE; SODIUM PHOSPHATE,
DIABASIC ANHYDROUS; SODIUM PHOSPHATE MONOBASIC, MONOHYDRATE

INJECTABLE; INJECTION

INTERSOL SOLUTION

FENWAL INC. 2.26G/500ML; 2.21G/500ML; 1.59G/500ML; N080041 NDA Dec 09, 2009
1.53G/500ML; 0.465G/500ML

DISCONTINUED DRUG PRODUCT LIST

6 - 1 (of 346)

ABARELIX

INJECTABLE; INTRAMUSCULAR
 PLENAXIS
 SPECIALITY EUROPEAN 100MG/VIAL N021320 001 Nov 25, 2003

ACETAMINOPHEN

INJECTABLE; INJECTION INJECTAPAP ORTHO MCNEIL PHARM	100MG/ML N017785 001 Mar 07, 1986
SUPPOSITORY; RECTAL ACEPHEN G AND W LABS	120MG A072218 001 Mar 27, 1992
ACETAMINOPHEN ABLE	120MG A073106 001 Feb 27, 1995
	325MG A073107 001 Feb 27, 1995
	650MG A073108 001 Feb 27, 1995
ROXANE	120MG A071010 001 May 12, 1987
	650MG A071011 001 May 12, 1987
TYLENOL MCNEIL CONS	120MG N017756 002
	650MG N017756 001
TABLET, EXTENDED RELEASE; ORAL ACETAMINOPHEN RANBAXY	650MG A090205 001 Nov 18, 2009

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE MIKART	150MG;180MG;15MG A081095 001 Oct 26, 1990
	150MG;180MG;30MG A081096 001 Oct 26, 1990
	150MG;180MG;60MG A081097 001 Oct 26, 1990
CODEINE, ASPIRIN, APAP FORMULA NO. 2 SCHERER LABS	150MG;180MG;15MG A085640 001
CODEINE, ASPIRIN, APAP FORMULA NO. 3 SCHERER LABS	150MG;180MG;30MG A085639 001
CODEINE, ASPIRIN, APAP FORMULA NO. 4 SCHERER LABS	150MG;180MG;60MG A085638 001

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL BANCAP FOREST PHARMS	325MG;50MG A088889 001 Jan 16, 1986
BUCET MALLINCKRODT	650MG;50MG A088991 001 Jun 28, 1985
TENCON MALLINCKRODT	650MG;50MG A089405 001 May 15, 1990
TRIAPRIN DUNHALL	325MG;50MG A089268 001 Jul 02, 1987
TABLET; ORAL BUTALBITAL AND ACETAMINOPHEN HALSEY	325MG;50MG A089568 001 Oct 05, 1988
WATSON LABS	325MG;50MG A087550 001 Oct 19, 1984
SEDAPAP MAYRAND	650MG;50MG A088944 001 Oct 17, 1985

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

DISCONTINUED DRUG PRODUCT LIST

6 - 2 (of 346)

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL					
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE					
MALLINCKRODT	325MG;50MG;40MG		A088758	001	Mar 27, 1985
BUTALIBITAL, ACETAMINOPHEN AND CAFFEINE					
GILBERT LABS	325MG;50MG;40MG		A088825	001	Dec 05, 1984
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
MUTUAL PHARM	325MG;50MG;40MG		A040601	001	Jul 29, 2005
WATSON LABS	325MG;50MG;40MG		A089536	001	Feb 16, 1988
ESGIC					
FOREST PHARMS	325MG;50MG;40MG		A089660	001	Dec 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

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL					
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					
WEST-WARD PHARM CORP	712.8MG;60MG;32MG		A040637	001	Sep 22, 2006

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	
CLONMEL	120MG/5ML;12MG/5ML		A040098	001	Sep 20, 1996
ROXANE	120MG/5ML;12MG/5ML		A086366	001	
TYLENOL W/ CODEINE					
ORTHO MCNEIL PHARM	120MG/5ML;12MG/5ML		A085057	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 3 (of 346)

ACETAMINOPHEN; CODEINE PHOSPHATE

SUSPENSION; ORAL				
CAPITAL AND CODEINE				
ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A085883	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
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;60MG	A089244	001	Feb 25, 1986
MUTUAL PHARM	300MG;15MG	A085795	001	
	300MG;15MG	A089671	001	Feb 10, 1988
	300MG;30MG	A085794	001	
	300MG;30MG	A089672	001	Feb 10, 1988
	300MG;60MG	A087653	001	Apr 13, 1982
	300MG;60MG	A089673	001	Feb 10, 1988
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	
	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 4 (of 346)

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

WARNER CHILCOTT	300MG;30MG 300MG;60MG	A085218 002 A087306 001	
WATSON LABS	300MG;15MG 300MG;15MG 300MG;30MG 300MG;30MG 300MG;60MG 300MG;60MG	A087277 001 A089997 001 A087276 001 A089998 001 A087275 001 A089999 001	May 26, 1982 Dec 28, 1994 May 26, 1982 Dec 28, 1994 May 26, 1982 Dec 28, 1994
WATSON LABS FLORIDA	300MG;15MG 300MG;30MG 300MG;60MG	A040443 001 A040443 002 A040443 003	Jan 22, 2003 Jan 22, 2003 Jan 22, 2003
WHITEWORTH TOWN PLSN	300MG;30MG 300MG;60MG	A084360 001 A085607 001	
CAPITAL AND CODEINE			
CARNRICK	325MG;30MG	A083643 001	
CODRIX			
WATSON LABS FLORIDA	500MG;15MG 500MG;30MG 500MG;60MG	A040447 001 A040441 001 A040488 001	Feb 26, 2003 Mar 27, 2003 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 325MG;15MG 325MG;30MG 325MG;60MG	A085056 001 A085056 002 A085056 003 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	

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 500MG;5MG	A088956 001 A089006 001	Jul 19, 1985 Aug 09, 1985
MIKART	500MG;5MG 500MG;5MG 500MG;5MG	A081068 001 A081069 001 A081070 001	Nov 30, 1989 Nov 30, 1989 Nov 30, 1989
LORCET-HD			
MALLINCKRODT	500MG;5MG	A087336 001	Jul 08, 1982
SOLUTION; ORAL			
HYDROCODONE BITARTRATE AND ACETAMINOPHEN			
MALLINCKRODT INC	500MG/15ML;10MG/15ML	A040508 001	Aug 29, 2003

DISCONTINUED DRUG PRODUCT LIST

6 - 5 (of 346)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MIKART	500MG/15ML;5MG/15ML	A081226	001	Oct 27, 1992
	500MG/15ML;5MG/15ML	A089557	001	Apr 29, 1992

TABLET; ORAL

DURADYNE DHC

FOREST PHARMS	500MG;5MG	A087809	001	Mar 17, 1983
---------------	-----------	---------	-----	--------------

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

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
	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

ENDO PHARMS	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

HALSEY	500MG;5MG	A089554	001	Jun 12, 1987
--------	-----------	---------	-----	--------------

IVAX PHARMS	500MG;5MG	A089696	001	Apr 21, 1988
-------------	-----------	---------	-----	--------------

MIKART	500MG;5MG	A089271	001	Jul 16, 1986
--------	-----------	---------	-----	--------------

MUTUAL PHARM	500MG;5MG	A089697	001	Jan 28, 1992
	650MG;7.5MG	A040236	001	Sep 25, 1997
	650MG;10MG	A040240	002	Nov 26, 1997
	750MG;7.5MG	A040240	001	Nov 26, 1997
	750MG;5MG	A040236	002	Sep 25, 1997

RANBAXY	500MG;10MG	A040825	001	Aug 16, 2007
---------	------------	---------	-----	--------------

RANBAXY LABS LTD	325MG;10MG	A040824	001	Aug 16, 2007
	750MG;7.5MG	A040826	001	Aug 16, 2007

SANDOZ	500MG;5MG	A040822	001	Aug 16, 2007
--------	-----------	---------	-----	--------------

	750MG;7.5MG	A040149	001	Jan 27, 1997
	750MG;5MG	A040149	002	Jan 27, 1997

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;5MG	A089831	001	Sep 07, 1988
----------------	-----------	---------	-----	--------------

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;5MG	A040122	001	Mar 04, 1996

	500MG;7.5MG	A040123	004	Mar 04, 1996
	650MG;7.5MG	A040123	001	Mar 04, 1996

	650MG;10MG	A040123	002	Mar 04, 1996
	750MG;7.5MG	A040122	002	Mar 04, 1996

WATSON LABS FLORIDA	500MG;5MG	A040493	001	May 28, 2003
	750MG;7.5MG	A040494	001	May 28, 2003

HY-PHEN	500MG;5MG	A087677	001	May 03, 1982
---------	-----------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 6 (of 346)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL					
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	

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL					
OXYCODONE AND ACETAMINOPHEN					
ACTAVIS TOTOWA	500MG;5MG		A040199	001	Dec 30, 1998
BARR	500MG;5MG		A040304	001	Oct 02, 2000
ENDO PHARMS	500MG;5MG		A040303	001	Dec 30, 1999
HALSEY	500MG;5MG		A089994	001	May 04, 1989
MUTUAL PHARM	500MG;5MG		A040219	001	Jan 22, 1998
TYLOX-325					
ORTHO MCNEIL PHARM	325MG;5MG		A088246	001	Nov 08, 1984
SOLUTION; ORAL					
OXYCODONE AND ACETAMINOPHEN					
MALLINCKRODT INC	325MG/5ML;5MG/5ML		A040680	001	Sep 29, 2006
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 TOTOWA	325MG;5MG		A040203	001	Mar 15, 1999
BARR	325MG;5MG		A087406	001	
DURAMED PHARMS BARR	325MG;5MG		A040272	001	Jun 30, 1998
PEROCET					
ENDO PHARMS	325MG;5MG		A085106	002	

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

CAPSULE; ORAL					
TYLOX					
ORTHO MCNEIL PHARM	500MG;4.5MG;0.38MG		A085375	001	

ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL					
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					
LEITNER PHARMS	650MG;65MG		A084999	001	

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL					
DARVOCET A500					
XANODYNE PHARM	500MG;100MG		A076429	001	Sep 10, 2003

DISCONTINUED DRUG PRODUCT LIST

6 - 7 (of 346)

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL					
DARVOSET-N 100					
XANODYNE PHARM	650MG;100MG		N017122	002	
DARVOSET-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		A077821	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
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

ACETAZOLAMIDE

TABLET; ORAL					
ACETAZOLAMIDE					
ALRA	250MG		A083320	001	
ASCOT	250MG		A087686	001	Oct 20, 1982
MUTUAL PHARM	250MG		A089753	001	Jun 22, 1988
VANGARD	250MG		A087654	001	Feb 05, 1982
WATSON LABS	250MG		A084498	002	
DIAMOX					
DURAMED PHARMS BARR	125MG		N008943	001	
	250MG		N008943	002	

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION					
ACETAZOLAMIDE SODIUM					
HOSPIRA	EQ 500MG BASE/VIAL		A040108	001	Oct 30, 1995

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	

DISCONTINUED DRUG PRODUCT LIST

6 - 8 (of 346)

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC 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
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			
BARR	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
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%	A072621	001 Sep 30, 1992
	20%	A072622	001 Sep 30, 1992
MUCOMYST			
APOTHECON	10%	N013601	002
	20%	N013601	001

DISCONTINUED DRUG PRODUCT LIST

6 - 9 (of 346)

ACETYLCYSTEINE

SOLUTION; INHALATION, ORAL 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

ACRISORCIN

CREAM; TOPICAL AKRINOL			
SCHERING	2MG/GM	N012470	001

ACYCLOVIR

CAPSULE; ORAL ACYCLOVIR			
ACTAVIS ELIZABETH	200MG	A074906	001 Aug 26, 1997
BELCHER PHARMS	200MG	A074889	001 Oct 31, 1997
DAVA PHARMS INC	200MG	A074872	001 Apr 22, 1997
IVAX SUB TEVA PHARMS	200MG	A074674	001 Apr 22, 1997
LEK PHARM	200MG	A074750	001 Apr 22, 1997
ROXANE	200MG	A074570	002 Apr 22, 1997
TEVA	200MG	A074828	001 Apr 22, 1997
TEVA PHARMS	200MG	A074914	001 Nov 26, 1997
TABLET; ORAL ACYCLOVIR			
ACTAVIS ELIZABETH	400MG	A074870	001 Jun 05, 1997
	800MG	A074870	002 Jun 05, 1997
BELCHER PHARMS	400MG	A074891	001 Oct 31, 1997
	800MG	A074891	002 Oct 31, 1997
DAVA PHARMS INC	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
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			
ABBOTT	EQ 50MG BASE/ML	A075114	001 Jul 26, 1999
ACYCLOVIR SODIUM APOTHECON	EQ 500MG BASE/VIAL	A074897	001 Feb 27, 1998
	EQ 1GM BASE/VIAL	A074897	002 Feb 27, 1998
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
TEVA PARENTERAL	EQ 50MG BASE/ML	A075627	001 Mar 28, 2001

DISCONTINUED DRUG PRODUCT LIST

6 - 10 (of 346)

ACYCLOVIR SODIUM

INJECTABLE; INJECTION ACYCLOVIR SODIUM TEVA PARENTERAL	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A074969 001 Aug 26, 1997 A074969 002 Aug 26, 1997
ZOVIRAX GLAXOSMITHKLINE	EQ 250MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	N018603 003 Aug 30, 1983 N018603 001 Oct 22, 1982 N018603 002 Jun 29, 1989

ADAPALENE

SOLUTION; TOPICAL DIFFERIN GALDERMA LABS LP	0.1%	N020338 001 May 31, 1996
---	------	--------------------------

ADENOSINE

INJECTABLE; INJECTION ADENOSINE BAXTER HLTHCARE	3MG/ML	A076501 001 Jun 16, 2004
---	--------	--------------------------

ALATROFLOXACIN MESYLATE

INJECTABLE; INJECTION TROVAN PRESERVATIVE FREE PFIZER	EQ 200MG BASE/VIAL EQ 300MG BASE/VIAL	N020760 001 Dec 18, 1997 N020760 002 Dec 18, 1997
---	--	--

ALBUMIN CHROMATED CR-51 SERUM

INJECTABLE; INJECTION CHROMALBIN ISO TEX	100uCi/VIAL 250uCi/VIAL 500uCi/VIAL	N017835 001 N017835 002 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 10uCi/ML 100uCi/ML	N017844 003 N017844 001 N017844 002

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION MEGATOPE ISO TEX	2mCi/VIAL 5uCi/AMP 20uCi/AMP	N017837 003 N017837 004 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 SCHERING	0.09MG/INH	N017559 001
VENTOLIN GLAXOSMITHKLINE	0.09MG/INH	N018473 001

DISCONTINUED DRUG PRODUCT LIST

6 - 11 (of 346)

ALBUTEROL SULFATE

CAPSULE; INHALATION					
VENTOLIN ROTACAPS					
GLAXOSMITHKLINE	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
BAUSCH AND LOMB	EQ 0.083% BASE		A075358	001	Mar 29, 2000
COPLEY PHARM	EQ 0.083% BASE		A073495	001	May 28, 1993
	EQ 0.5% BASE		A073307	001	Nov 27, 1991
ROXANE	EQ 0.083% BASE		A075129	001	Feb 13, 2001
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		A072859	001	Dec 20, 1989
	EQ 4MG BASE		A072860	001	Dec 20, 1989
PLIVA	EQ 2MG BASE		A072316	001	Dec 05, 1989
	EQ 4MG BASE		A072317	001	Dec 05, 1989
SANDOZ	EQ 2MG BASE		A072151	001	Dec 05, 1989
	EQ 4MG BASE		A072152	001	Dec 05, 1989
TEVA	EQ 2MG BASE		A072619	001	Dec 05, 1989
	EQ 2MG BASE		A072779	001	Jun 25, 1993
	EQ 2MG 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 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 - 12 (of 346)

ALCOHOL

INJECTABLE; INJECTION
 ALCOHOL 5% IN DEXTROSE 5%
 MILES 5ML/100ML A083483 001

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION
 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

TABLET; ORAL
 ALENDRONATE SODIUM
 SANDOZ 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
 TEVA PHARMS EQ 35MG BASE A076184 002 Aug 04, 2008
 EQ 70MG BASE A076184 001 Feb 06, 2008

ALGLUCERASE

INJECTABLE; INJECTION
 CEREDASE
 GENZYME 10 UNITS/ML N020057 004 May 08, 1992

ALKAVERVIR

TABLET; ORAL
 VERILOID
 3M 2MG N007336 002
 3MG N007336 003

ALLOPURINOL

TABLET; 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

ALPRAZOLAM

SOLUTION; ORAL
 ALPRAZOLAM
 ROXANE 0.5MG/5ML A074314 001 Oct 31, 1993
 TABLET; ORAL
 ALPRAZOLAM
 IVAX SUB TEVA PHARMS 0.25MG A074294 001 Jul 29, 1994
 0.5MG A074294 002 Jul 29, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 13 (of 346)

ALPRAZOLAMTABLET; ORAL
ALPRAZOLAM

IVAX SUB TEVA PHARMS	1MG	A074294	003	Jul 29, 1994
	2MG	A074294	004	Jul 29, 1994
ROXANE	0.25MG	A074199	001	Oct 19, 1993
	0.5MG	A074199	002	Oct 19, 1993
	1MG	A074199	003	Oct 19, 1993
TEVA	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
WATSON LABS	0.25MG	A074456	001	Aug 31, 1995
	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

ALPROSTADILINJECTABLE; 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				
SCHWARZ PHARMA	0.005MG/VIAL	N020649	001	Jun 12, 1997

ALSEROXYLONTABLET; ORAL
RAUTENSIN
NOVARTIS
RAUWILOID
3M

NOVARTIS	2MG	N009215	001
RAUWILOID	2MG	N008867	001

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL ALUMINUM HYDROXIDE AND MAGNESIUM TRISILICATE			
PENNEX	80MG;20MG	A089449	001
FOAMICON		A072687	001

NOVARTIS 80MG;20MG Nov 27, 1987

NOVARTIS 80MG;20MG Jun 28, 1989

AMANTADINE HYDROCHLORIDECAPSULE; ORAL
AMANTADINE HYDROCHLORIDE

ACTAVIS TOTOWA	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				
ACTAVIS MID ATLANTIC	50MG/5ML	A072655	001	Oct 30, 1990
TEVA PHARMS	50MG/5ML	A073115	001	Aug 23, 1991

ENDO PHARMS 50MG/5ML N016023 002

TABLET; ORAL

SYMMETREL				
ENDO PHARMS	50MG/5ML	N016023	002	
ENDO PHARMS	100MG	N018101	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 14 (of 346)

AMCINONIDE

CREAM; TOPICAL CYCLOCORT ASTELLAS	0.025% 0.1%	N018116 001 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 500MG/VIAL 1GM/VIAL	N050565 001 Dec 21, 1984 N050565 002 Dec 21, 1984 N050565 003 Dec 21, 1984
---	--------------------------------------	--

AMIFOSTINE

INJECTABLE; INJECTION ETHYOL MEDIMMUNE	375MG/VIAL	N020221 002 Sep 10, 1999
--	------------	--------------------------

AMIKACIN SULFATE

INJECTABLE; INJECTION AMIKACIN SULFATE ABBOTT	EQ 250MG BASE/ML EQ 250MG BASE/ML	A063265 001 Nov 30, 1994 A063266 001 Oct 31, 1994
ASTRAZENECA	EQ 50MG BASE/ML EQ 250MG BASE/ML	A063167 001 Dec 14, 1995 A063169 001 Dec 14, 1995
BAXTER HLTHCARE	EQ 50MG BASE/ML EQ 250MG BASE/ML	A063274 001 May 18, 1992 A063275 001 May 18, 1992
HOSPIRA	EQ 50MG BASE/ML EQ 62.5MG BASE/ML EQ 250MG BASE/ML EQ 250MG BASE/ML EQ 250MG BASE/ML	A063350 001 Jul 30, 1993 A063283 001 Oct 31, 1994 A063350 002 Jul 30, 1993 A064098 001 Jun 26, 1995 A064099 001 Jun 20, 1995
TEVA PARENTERAL	EQ 50MG BASE/ML	A064045 001 Sep 28, 1993
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 EQ 50MG BASE/ML EQ 50MG BASE/ML EQ 250MG BASE/ML EQ 250MG BASE/ML EQ 250MG BASE/ML	A062311 001 A062562 001 Sep 20, 1984 N050495 001 A062311 002 A062562 002 Sep 20, 1984 N050495 002
AMIKIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER APOTHECON	EQ 5MG BASE/ML EQ 10MG BASE/ML	N050618 002 Nov 30, 1987 N050618 001 Nov 30, 1987

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE SANDOZ	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

DISCONTINUED DRUG PRODUCT LIST

6 - 15 (of 346)

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%					
HOSPIRA	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%					
HOSPIRA	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
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		
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% (15GM/100ML)	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

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 2.4MG/100ML;261MG/100ML;205MG/100ML	N019714	003	Sep 12,	1988
HOSPIRA INC	5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2 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

DISCONTINUED DRUG PRODUCT LIST

6 - 16 (of 346)

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT	3.5%;25GM/100ML 3.5%;25GM/100ML	N019505 002 N019713 006	Nov 07, 1986 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 3.5%;5GM/100ML	N019506 001 N019713 002	Nov 07, 1986 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 4.25%;10GM/100ML	N019713 001 N019681 004	Sep 09, 1988 Nov 01, 1988
HOSPIRA	4.25%;10GM/100ML	N019681 005	Nov 01, 1988

AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

ABBOTT	4.25%;20GM/100ML 4.25%;20GM/100ML	N019713 004 N019681 005	Sep 09, 1988 Nov 01, 1988
HOSPIRA	4.25%;20GM/100ML	N019681 003	Nov 01, 1988

AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT	4.25%;25GM/100ML 4.25%;25GM/100ML	N019504 002 N019713 005	Nov 07, 1986 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 5%;25GM/100ML	N019565 001 N019713 003	Dec 17, 1986 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
-----------------	------------------	-------------	--------------

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	4.25%;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML	N019564 004	Dec 16, 1986
--	--	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 17 (of 346)

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;12 0MG/100ML;49.3MG/100ML	N019564	001	Dec 16,	1986	
	3.5%;5GM/100ML;30MG/100ML;97MG/100ML;12 0MG/100ML;49.3MG/100ML	N019712	001	Sep 08,	1988	
HOSPIRA INC	3.5%;5GM/100ML;30MG/100ML;97MG/100ML;12 0MG/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;1 20MG/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	
TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER						
BAXTER HLTHCARE	4.25%;25GM/100ML;51MG/100ML;261MG/100ML ;297MG/100ML;77MG/100ML	N020147	010	Oct 23,	1995	
TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER						
BAXTER HLTHCARE	4.25%;5GM/100ML;51MG/100ML;261MG/100ML; 297MG/100ML;77MG/100ML	N020147	006	Oct 23,	1995	

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; 234MG/100ML	N018804	002	May 15,	1984	
	3.5%;21MG/100ML;40MG/100ML;128MG/100ML; 234MG/100ML	N018875	002	Aug 08,	1984	

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M	
HOSPIRA	3.5%;21MG/100ML;128MG/100ML;234MG/100ML

DISCONTINUED DRUG PRODUCT LIST

6 - 18 (of 346)

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/100MLAMINO 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/100MLAMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 7% W/ ELECTROLYTES

HOSPIRA 7%;102MG/100ML;45MG/100ML;522MG/100ML;4 N019437 006 Apr 03, 1986
10MG/100MLAMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M

HOSPIRA 3.5%;30MG/100ML;97MG/100ML;120MG/100ML; N019437 007 Apr 03, 1986
49MG/100MLAMINOCAPROIC ACID

INJECTABLE; INJECTION

AMICAR

XANODYNE PHARM 250MG/ML N015229 002

AMINOCAPROIC ACID

ABRAXIS PHARM 250MG/ML A070522 001 Jun 17, 1986

BAXTER HLTHCARE 250MG/ML N018590 001 Oct 29, 1982

HOSPIRA 250MG/ML A070888 001 Jun 16, 1988

AMINOGLUTETHIMIDE

TABLET; ORAL

CYTADREN

NOVARTIS 250MG N018202 001

AMINOPHYLLINE

ENEMA; RECTAL

SOMOPHYLLIN

FISON'S 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 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

SMITH AND NEPHEW 25MG/ML A088429 001 May 30, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 19 (of 346)

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

SMITH AND NEPHEW	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

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	
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

SANDOZ	100MG	A085261	003	
	100MG	A085262	002	
	200MG	A085261	002	

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	

DISCONTINUED DRUG PRODUCT LIST

6 - 20 (of 346)

AMINOPHYLLINE

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
SODIUM AMINOSALICYLATE			
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

AMINOSALICYLIC ACID RESIN COMPLEX

POWDER; ORAL			
REZIPAS			
BRISTOL MYERS SQUIBB	EQ 500MG BASE/GM	N009052	001

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION			
AMIODARONE HYDROCHLORIDE			
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
CORDARONE			
WYETH PHARMS INC	50MG/ML	N020377	001 Aug 03, 1995
TABLET; ORAL			
AMIODARONE HYDROCHLORIDE			
TEVA	200MG	A074895	001 Apr 16, 1999

AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE; ORAL			
ENDEP			
ROCHE	40MG/ML	A085749	001
INJECTABLE; INJECTION			
AMITRIPTYLINE HYDROCHLORIDE			
WATSON LABS	10MG/ML	A085594	001

DISCONTINUED DRUG PRODUCT LIST

6 - 21 (of 346)

AMITRIPTYLINE HYDROCHLORIDE

INJECTABLE; INJECTION

ELAVIL

ASTRAZENECA 10MG/ML

N012704 001

TABLET; ORAL

AMITID

BRISTOL MYERS SQUIBB 10MG
25MG
50MG
75MG
100MGA086454 001
A086454 002
A086454 003
A086454 004
A086454 005

AMITRIL

WARNER CHILCOTT 10MG
25MG
50MG
75MG
100MG
150MGA083939 001
A083937 001
A083938 002
A084957 001
A085093 001
A086295 001

AMITRIPTYLINE HYDROCHLORIDE

AM THERAP 25MG
50MG
75MG
100MGA088672 001 Nov 20, 1984
A088673 001 Nov 20, 1984
A088674 001 Nov 20, 1984
A088675 001 Nov 20, 1984COPELEY PHARM 10MG
25MG
50MG
75MG
100MG
150MGA088421 001 Apr 30, 1984
A088422 001 Apr 30, 1984
A088423 001 Apr 30, 1984
A088424 001 Apr 30, 1984
A088425 001 Apr 30, 1984
A088426 001 Apr 30, 1984HALSEY 10MG
25MG
50MG
50MG
75MG
100MGA085923 001
A085922 001
A085925 001
A087557 001 Mar 05, 1982
A085926 001 May 20, 1983
A085927 001 May 20, 1983LEDERLE 10MG
10MG
25MG
25MG
50MG
50MG
75MG
75MG
100MG
100MG
150MGA086744 001
A087366 001 Jan 04, 1982
A086746 001
A087367 001 May 03, 1982
A086743 001
A087181 001 Jan 04, 1982
A086745 001
A087369 001 Jan 04, 1982
A086747 001
A087368 001 May 03, 1982
A087370 001 Jan 04, 1982MUTUAL PHARM 10MG
25MG
50MG
75MG
100MG
150MGA085744 001
A085627 001
A085745 001
A085743 001
A085742 002 May 11, 1982
A089423 001 Feb 17, 1987PAR PHARM 10MG
25MG
50MG
75MG
100MG
150MGA088697 001 Sep 25, 1984
A088698 001 Sep 25, 1984
A088699 001 Sep 25, 1984
A088700 001 Sep 25, 1984
A088701 001 Sep 25, 1984
A088702 001 Sep 25, 1984PLIVA 10MG
25MG
50MG
75MG
100MGA088883 001 Sep 26, 1984
A088884 001 Sep 26, 1984
A088885 001 Sep 26, 1984
A088886 001 Sep 26, 1984
A088887 001 Sep 26, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 22 (of 346)

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

PLIVA	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	
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
TEVA	10MG	A084910	003	
	10MG	A086610	001	
	25MG	A085031	001	
	25MG	A086859	001	
	50MG	A085032	001	
	50MG	A086857	001	
	75MG	A085030	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

DISCONTINUED DRUG PRODUCT LIST

6 - 23 (of 346)

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL		
AMITRIPTYLINE HYDROCHLORIDE		
WATSON LABS	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		
MUTUAL PHARM	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 12.5MG BASE;5MG	A072052 001 Dec 16, 1988
	EQ 25MG BASE;10MG	A072053 001 Dec 16, 1988

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
MUTUAL PHARM	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
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

DISCONTINUED DRUG PRODUCT LIST

6 - 24 (of 346)

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

SANDOZ	50MG;4MG	A071863	001	Dec 21, 1987
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
	10MG;4MG	A073009	001	Oct 17, 1991
	25MG;2MG	A070374	001	Aug 25, 1986
	25MG;2MG	A072541	001	Feb 15, 1989
	25MG;2MG	A073008	001	Oct 17, 1991
	25MG;4MG	A070376	001	Aug 25, 1986
	25MG;4MG	A072134	001	Feb 15, 1989
	25MG;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

PATCH; TOPICAL

AMLEXANOX

ULURU

2MG

N021727 001 Sep 29, 2004

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

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
MUTUAL PHARMA	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
SYNTTHON 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

SYNTTHON 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

DISCONTINUED DRUG PRODUCT LIST

6 - 25 (of 346)

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

AMODIAQUINE HYDROCHLORIDE

TABLET; ORAL		
CAMOQUIN HYDROCHLORIDE		
PARKE DAVIS	EQ 200MG BASE	N006441 001

AMOXAPINE

TABLET; ORAL		
AMOXAPINE		
SANDOZ	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 LABS	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

AMOXICILLIN

CAPSULE; ORAL		
AMOXICILLIN		
LABS ATRAL	250MG	A062528 001 Aug 07, 1985
	500MG	A062528 002 Aug 07, 1985
MYLAN	250MG	A062067 001
	500MG	A062067 002
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

DISCONTINUED DRUG PRODUCT LIST

6 - 26 (of 346)

AMOXICILLINFOR SUSPENSION; ORAL
AMOXICILLIN

AM ANTIBIOTICS	125MG/5ML 250MG/5ML	A062059 001 A062059 002
MYLAN	125MG/5ML 250MG/5ML	A062090 001 A062090 002
TEVA	125MG/5ML 250MG/5ML	A062946 001 Nov 01, 1988 A063001 001 Jan 06, 1989
AMOXIL		
GLAXOSMITHKLINE	50MG/ML 125MG/5ML 250MG/5ML	N050460 005 N050460 001 N050460 002
LAROTID		
GLAXOSMITHKLINE	50MG/ML	N050460 006
POLYMOX		
APOTHECON	125MG/5ML 125MG/5ML 250MG/5ML 250MG/5ML	A061851 001 A062323 001 A061851 002 A062323 002
TRIMOX		
APOTHECON	50MG/ML 125MG/5ML 125MG/5ML 125MG/5ML 125MG/5ML 250MG/5ML 250MG/5ML 250MG/5ML 250MG/5ML	A061886 001 A061886 002 A062099 001 A062154 001 A062885 001 Mar 08, 1988 A061886 003 A062099 002 A062154 002 A062885 002 Mar 08, 1988
UTIMOX		
PARKE DAVIS	125MG/5ML 250MG/5ML	A062127 001 A062127 002
WYMOX		
WYETH AYERST	125MG/5ML 250MG/5ML	A062131 001 A062131 002
TABLET, CHEWABLE; ORAL		
AMOXICILLIN		
APOTHECON	125MG 250MG	A064131 001 May 06, 1996 A064131 002 May 06, 1996
TEVA	125MG 250MG	A064031 001 Dec 19, 1996 A064031 002 Dec 19, 1996
TABLET, FOR SUSPENSION; ORAL		
DISPERMOX		
RANBAXY LABS LTD	200MG 400MG	A065080 002 Aug 11, 2003 A065080 001 Aug 11, 2003

AMPHETAMINE ADIPATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE ADIPATE;
DEXTROAMPHETAMINE SULFATECAPSULE; 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

DISCONTINUED DRUG PRODUCT LIST

6 - 27 (of 346)

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE;
DEXTROAMPHETAMINE SULFATE

TABLET; ORAL ADDERALL 10				
TEVA WOMENS ADDERALL 12.5	2.5MG;2.5MG;2.5MG;2.5MG	N011522	007	Feb 13, 1996
TEVA WOMENS ADDERALL 15	3.125MG;3.125MG;3.125MG;3.125MG	N011522	012	Aug 31, 2000
TEVA WOMENS ADDERALL 20	3.75MG;3.75MG;3.75MG;3.75MG	N011522	013	Aug 31, 2000
TEVA WOMENS ADDERALL 30	5MG;5MG;5MG;5MG	N011522	008	Feb 13, 1996
TEVA WOMENS ADDERALL 5	7.5MG;7.5MG;7.5MG;7.5MG	N011522	010	May 12, 1997
TEVA WOMENS ADDERALL 7.5	1.25MG;1.25MG;1.25MG;1.25MG	N011522	009	May 12, 1997
TEVA WOMENS DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE	1.875MG;1.875MG;1.875MG;1.875MG	N011522	011	Aug 31, 2000
MALLINCKRODT INC	1.25MG;1.25MG;1.25MG;1.25MG 1.875MG;1.875MG;1.875MG;1.875MG 2.5MG;2.5MG;2.5MG;2.5MG 3.125MG;3.125MG;3.125MG;3.125MG 3.75MG;3.75MG;3.75MG;3.75MG 5MG;5MG;5MG;5MG 7.5MG;7.5MG;7.5MG;7.5MG	A040440	001 002 003 004 005 006 007	Oct 07, 2003
MUTUAL PHARM	1.25MG;1.25MG;1.25MG;1.25MG 1.875MG;1.875MG;1.875MG;1.875MG 2.5MG;2.5MG;2.5MG;2.5MG 3.125MG;3.125MG;3.125MG;3.125MG 3.75MG;3.75MG;3.75MG;3.75MG 5MG;5MG;5MG;5MG 7.5MG;7.5MG;7.5MG;7.5MG	A040480	001 002 003 004 005 006 007	Sep 09, 2003
WATSON LABS	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	A040456	001 002 003 004	May 06, 2003 May 06, 2003 May 06, 2003 May 06, 2003

AMPHETAMINE RESIN COMPLEX; DEXTROAMPHETAMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL BIPHETAMINE 12.5				
UCB INC BIPHETAMINE 20	EQ 6.25MG BASE;EQ 6.25MG BASE	N010093	007	
UCB INC BIPHETAMINE 7.5	EQ 10MG BASE;EQ 10MG BASE	N010093	003	
UCB INC	EQ 3.75MG BASE;EQ 3.75MG BASE	N010093	009	

AMPHETAMINE SULFATE

TABLET; ORAL AMPHETAMINE SULFATE				
LANNETT	5MG	A083901	001	Aug 31, 1984
	10MG	A083901	002	Aug 31, 1984

AMPHOTERICIN B

CREAM; TOPICAL FUNGIZONE APOTHECON INJECTABLE; INJECTION AMPHOTERICIN B	3%	N050314	001	
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 28 (of 346)

AMPHOTERICIN B

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
BAXTER HLTHCARE	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
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	A063143	001	Apr 15, 1993
	EQ 500MG BASE/VIAL	A063147	001	Apr 15, 1993
	EQ 1GM BASE/VIAL	A063139	001	Apr 15, 1993
	EQ 2GM BASE/VIAL	A063141	001	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
MARSAM PHARMS LLC	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
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
	EQ 500MG BASE/VIAL	A060626	003	
	EQ 500MG BASE/VIAL	A062718	003	Dec 16, 1986
	EQ 1GM BASE/VIAL	A060626	004	
	EQ 1GM BASE/VIAL	A062718	004	Dec 16, 1986
	EQ 2GM BASE/VIAL	A060626	005	
	EQ 2GM BASE/VIAL	A062718	005	Dec 16, 1986
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 29 (of 346)

AMPICILLIN SODIUMINJECTABLE; INJECTION
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 1GM BASE/VIAL	A062727	001	Dec 19, 1986
	EQ 2GM BASE/VIAL	A060677	005	
	EQ 2GM BASE/VIAL	A062727	002	Dec 19, 1986
	EQ 10GM BASE/VIAL	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	

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	

DISCONTINUED DRUG PRODUCT LIST

6 - 30 (of 346)

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

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 - 31 (of 346)

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

CAPSULE; ORAL PRINCIPEN W/ PROBENECID		
APOTHECON	EQ 389MG BASE;111MG	A062150 001
	EQ 389MG BASE;111MG	N050488 001
FOR SUSPENSION; ORAL POLYCILLIN-PRB		
APOTHECON	EQ 3.5GM BASE/BOT;1GM/BOT	A061898 001
BRISTOL	EQ 3.5GM BASE/BOT;1GM/BOT	N050457 001
PROBAMPACIN		
TEVA	EQ 3.5GM BASE/BOT;1GM/BOT	A061741 001

AMPRENAVIR

CAPSULE; ORAL AGENERASE			
GLAXOSMITHKLINE	50MG	N021007 001	Apr 15, 1999
	150MG	N021007 002	Apr 15, 1999
SOLUTION; ORAL AGENERASE			
GLAXOSMITHKLINE	15MG/ML	N021039 001	Apr 15, 1999

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL AGRILYN			
SHIRE	EQ 1MG BASE	N020333 002	Mar 14, 1997
ANAGRELIDE HYDROCHLORIDE			
ROXANE	EQ 0.5MG BASE	A076489 001	Apr 18, 2005
	EQ 1MG BASE	A076489 002	Apr 18, 2005

ANASTROZOLE

TABLET; ORAL ANASTROZOLE			
SYNTTHON PHARMS	1MG	A078322 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	

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC VASOCON-A			
NOVARTIS	0.5%;0.05%	N018746 002	Jul 11, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 32 (of 346)

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS
 APOKYN
 US WORLDMEDS 20MG/2ML (10MG/ML) N021264 001 Apr 20, 2004

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

TABLET, ORALLY DISINTEGRATING; ORAL
 ABILIFY
 OTSUKA 20MG N021729 004 Jun 07, 2006
 30MG N021729 005 Jun 07, 2006

ARMODAFINIL

TABLET; ORAL
 NUvigil
 CEPHALON 100MG N021875 002 Mar 26, 2009
 200MG N021875 005 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;
 1.5MG/ML; 1,650 IU/ML; 5 IU/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/
 ML; 0.3MG/ML; 330 UNITS/ML; 1 IU/ML

M.V.I.-12
 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/
 ML; 0.3MG/ML; 330 UNITS/ML; 1 IU/ML

MVC PLUS
 WATSON LABS 10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2 N018439 002 Aug 08, 1985
 0
 IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/
 ML; 0.3MG/ML; 330 UNITS/ML; 1 IU/ML

DISCONTINUED DRUG PRODUCT LIST

6 - 33 (of 346)

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12

HOSPIRA

20MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;2 IU/ML;0.6MG/ML;4MG/ML;0.4MG/ML;0.36MG/M L;0.6MG/ML;330 UNITS/ML;1 IU/ML

Apr 22, 2004

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

ASTRAZENECA

100MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15MG G/VIAL;5MCG/VIAL;0.4MG/VIAL;40MG/VIAL;4 MG/VIAL;3.6MG/VIAL;3MG/VIAL;1MG/VIAL;10 MG/VIAL

Aug 08, 1985

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A; PALMITATE; VITAMIN E

INJECTABLE; INJECTION

VITAPED

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

Dec 29, 1993

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

MUTUAL PHARM 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

FIORINAL

WATSON LABS INC 325MG;50MG;40MG

N017534 003 Apr 16, 1986

LANORINAL

LANNETT 325MG;50MG;40MG

A086986 002 Oct 18, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 34 (of 346)

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL
 BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE
 ENDO PHARMS 325MG;50MG;40MG;30MG A075351 001 Mar 05, 1999

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL
 ORPHENGESIC
 SOLCO HLTHCARE 385MG;30MG;25MG A075141 001 May 29, 1998
 ORPHENGESIC FORTE
 SOLCO HLTHCARE 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

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL
 AZDONE
 SCHWARZ PHARMA 500MG;5MG A089420 001 Jan 25, 1988
 VICOPRIN
 ABBOTT 500MG;5MG A086333 001 Sep 14, 1983

ASPIRIN; MEPROBAMATE

TABLET; ORAL
 EQUAGESIC
 CARACO 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

DISCONTINUED DRUG PRODUCT LIST

6 - 35 (of 346)

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL CODOXY					
HALSEY	325MG;4.5MG;0.38MG		A087464	001	Jul 01, 1982
OXYCODONE AND ASPIRIN					
MUTUAL PHARM	325MG;4.5MG;0.38MG		A040260	001	Jul 17, 1998
	325MG;4.5MG;0.38MG		A087794	001	May 26, 1982
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	

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
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

DISCONTINUED DRUG PRODUCT LIST

6 - 36 (of 346)

ATENOLOL

TABLET; ORAL

ATENOLOL

SANDOZ	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

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

ACTAVIS ELIZABETH	10MG	A078940	001	Aug 30, 2010
	18MG	A078940	002	Aug 30, 2010
	25MG	A078940	003	Aug 30, 2010
	40MG	A078940	004	Aug 30, 2010
	60MG	A078940	005	Aug 30, 2010
	80MG	A078940	006	Aug 30, 2010
	100MG	A078940	007	Aug 30, 2010
STRATTERA				
LILLY	5MG	N021411	001	Nov 26, 2002

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
MARSAM PHARMS LLC	10MG/ML	A074945	001	Jul 28, 1998
TEVA PARENTERAL	10MG/ML	A074784	001	Jun 11, 1997
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
MARSAM PHARMS LLC	10MG/ML	A074741	001	Mar 28, 1997
TRACRIUM		A074944	001	Jul 28, 1998
HOSPIRA	10MG/ML	N018831	002	Jun 20, 1985
TRACRIUM PRESERVATIVE FREE		N018831	001	Nov 23, 1983
HOSPIRA	10MG/ML			

ATROPINE

INJECTABLE; INJECTION

ATROPINE

SOLVAY	EQ 2MG SULFATE/0.7ML	A071295	001	Jan 30, 1987
--------	----------------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 37 (of 346)

ATROPINE SULFATE

AEROSOL, METERED; INHALATION
ATROPINE SULFATE
US ARMY EQ 0.36MG BASE/INH N020056 001 Sep 19, 1990

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL
MOTOFEN HALF-STRENGTH
VALEANT 0.025MG; 0.5MG N017744 001

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL				
DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE				
SCHERER RP	0.025MG;2.5MG		A086440	001
SOLUTION; ORAL				
COLONAIID				
MEDPOINTE PHARM HLC	0.025MG/5ML;2.5MG/5ML		A085735	001
LOMANATE				
ALPHARMA US PHARMS	0.025MG/5ML;2.5MG/5ML		A085746	001
TABLET; ORAL				
COLONAIID				
MEDPOINTE PHARM HLC	0.025MG;2.5MG		A085737	001
DI-ATRO				
MD PHARM	0.025MG;2.5MG		A085266	001
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE 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
INWOOD LABS	0.025MG;2.5MG		A085509	001
IVAX PHARMS	0.025MG;2.5MG		A086727	001
KV PHARM	0.025MG;2.5MG		A085659	001
LEDERLE	0.025MG;2.5MG		A086950	001
MUTUAL PHARM	0.025MG;2.5MG		A085506	001
PARKER 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
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
WEST WARD	0.025MG;2.5MG		A087765	001 Mar 15, 1982
LOGEN				
SUPERPHARM	0.025MG;2.5MG		A088962	001 May 10, 1985
LO-TROL				
VANGARD	0.025MG;2.5MG		A088009	001 Mar 25, 1983
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

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION				
ATROPINE AND DEMEROL				
ABBOTT	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 38 (of 346)

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION
 MEPERIDINE AND ATROPINE SULFATE
 WYETH AYERST 0.4MG/ML;100MG/ML A085121 003

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR
 ATNAA
 US ARMY 2.1MG/0.7ML;600MG/0.7ML N021175 001 Jan 17, 2002

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
 PROMETHEUS LABS 25MG N016324 002

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION
 IMURAN
 PROMETHEUS LABS EQ 100MG BASE/VIAL N017391 001

AZELASTINE HYDROCHLORIDE

SPRAY, METERED; NASAL
 ASTEPRO
 MEDA PHARMS EQ 0.125MG BASE/SPRAY N022203 001 Oct 15, 2008

AZITHROMYCIN

CAPSULE; ORAL
 ZITHROMAX
 PFIZER EQ 250MG BASE N050670 001 Nov 01, 1991

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 A062388 001 Sep 08, 1982
 EQ 2GM BASE/VIAL A062417 001 Oct 12, 1982
 EQ 2GM BASE/VIAL N050562 001 Sep 03, 1982
 EQ 3GM BASE/VIAL A062388 002 Sep 08, 1982
 EQ 3GM BASE/VIAL A062417 002 Oct 12, 1982
 EQ 3GM BASE/VIAL N050562 002 Sep 03, 1982
 EQ 4GM BASE/VIAL A062388 003 Sep 08, 1982
 EQ 4GM BASE/VIAL A062417 003 Oct 12, 1982
 EQ 4GM BASE/VIAL N050562 003 Sep 03, 1982

AZTREONAM

INJECTABLE; INJECTION
 AZACTAM
 BRISTOL MYERS SQUIBB 500MG/VIAL N050580 001 Dec 31, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 39 (of 346)

AZTREONAM

INJECTABLE; INJECTION
 AZACTAM IN PLASTIC CONTAINER
 BRISTOL MYERS SQUIBB 10MG/ML N050632 003 May 24, 1989

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION; ORAL
 SPECTROBID
 PFIZER 125MG/5ML N050556 001 Mar 23, 1982
 TABLET; ORAL
 SPECTROBID
 PFIZER 400MG N050520 001
 800MG 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
 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
 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
 MONARCH PHARMS 400 UNITS/GM;1%;EQ 3.5MG N050416 002
 BASE/GM;10,000 UNITS/GM
 ZINC BACITRACIN,NEOMYCIN SULFATE,POLYMYXIN B SULFATE & HYDROCORTISONE
 PHARMAFAIR 400 UNITS/GM;1%;EQ 3.5MG A062389 001 Jul 02, 1982
 BASE/GM;10,000 UNITS/GM
 OINTMENT; TOPICAL
 NEOMYCIN & POLYMYXIN B SULFATES & BACITRACIN ZINC & HYDROCORTISONE
 PHARMAFAIR 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 A062381 001 Sep 06, 1985
 UNITS/GM

BACITRACIN ZINC; LIDOCAINE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; TOPICAL
 LANABIOTIC
 COMBE 400 UNITS/GM;40MG/GM;EQ 5MG A062499 001 Jun 03, 1985
 BASE/GM;5,000 UNITS/GM

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 A062386 001 Sep 09, 1982
 UNITS/GM

DISCONTINUED DRUG PRODUCT LIST

6 - 40 (of 346)

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC BACITRACIN-NEOMYCIN-POLYMYXIN PHARMADEERM	400 UNITS/GM; EQ 3.5MG BASE/GM; 5,000 UNITS/GM	A062167	001
NEO-POLYCIN DOW PHARM	500 UNITS/GM; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	A060647	001
NEOSPORIN MONARCH PHARMS	400 UNITS/GM; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	N050417	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

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 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	A073092	001 Jan 28, 1994
	10MG	A074698	001 Aug 20, 1996
	20MG	A073093	001 Jan 28, 1994
	20MG	A074698	002 Aug 20, 1996
LIORESAL NOVARTIS	10MG	N017851	001
	20MG	N017851	003 Jan 20, 1982
TABLET, ORALLY DISINTEGRATING; ORAL KEMSTRO			
SCHWARZ PHARMA	10MG	N021589	001 Oct 30, 2003
	20MG	N021589	002 Oct 30, 2003

DISCONTINUED DRUG PRODUCT LIST

6 - 41 (of 346)

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, 1996

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL BENAZEPRIL HYDROCHLORIDE				
GENPHARM	5MG		A076476	001 Feb 11, 2004
	10MG		A076476	002 Feb 11, 2004
	20MG		A076476	003 Feb 11, 2004
	40MG		A076476	004 Feb 11, 2004

BENDROFLUMETHIAZIDE

TABLET; ORAL NATURETIN-10				
APOTHECON	10MG		N012164	003
NATURETIN-2.5				
APOTHECON	2.5MG		N012164	001
NATURETIN-5				
APOTHECON	5MG		N012164	002

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

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL DIDREX				
PHARMACIA AND UPJOHN	25MG		N012427	003

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION EMETE-CON				
PFIZER	EQ 50MG BASE/VIAL		N016820	001
SUPPOSITORIY; RECTAL EMETE-CON				
ROERIG	EQ 100MG BASE		N016818	006

DISCONTINUED DRUG PRODUCT LIST

6 - 42 (of 346)

BENZTHIAZIDE

TABLET; ORAL AQUATAG SOLVAY	25MG 50MG	N016001 001 N016001 002
BENZTHIAZIDE PVT FORM EXNA	50MG	A083206 001
AH ROBINS INC	50MG	N012489 001
FOVANE PFIZER	50MG	N012128 002
URESE PFIZER	25MG	N012128 003

BENZTROPINE MESYLATE

TABLET; ORAL BENZTROPINE MESYLATE ACTAVIS TOTOWA	0.5MG 1MG 2MG	A040699 001 Feb 14, 2008 A040705 001 Feb 14, 2008 A040706 001 Feb 14, 2008
MUTUAL PHARM	1MG 2MG	A081264 001 Jan 23, 1992 A081265 001 Jan 23, 1992
QUANTUM PHARMICS	0.5MG 1MG 2MG	A088514 001 Jan 31, 1984 A088510 001 Jan 31, 1984 A088511 001 Jan 31, 1984
USL PHARMA	0.5MG 1MG 2MG	A089211 001 Jun 14, 1988 A089212 001 Jun 14, 1988 A089213 001 Jun 14, 1988
COGENTIN MERCK	0.5MG 1MG 2MG	N009193 004 N009193 003 N009193 002

BENZYL BENZOATE

EMULSION; TOPICAL BENZYL BENZOATE LANNETT	50%	A084535 001
---	-----	-------------

BEPRIDIL HYDROCHLORIDE

TABLET; ORAL BEPADIN MEDPOINTE PHARM HLC	200MG 300MG 400MG	N019001 001 Dec 28, 1990 N019001 002 Dec 28, 1990 N019001 003 Dec 28, 1990
VASCOR JOHNSON AND JOHNSON	200MG 300MG 400MG	N019002 001 Dec 28, 1990 N019002 002 Dec 28, 1990 N019002 003 Dec 28, 1990

BETA CAROTENE

CAPSULE; ORAL SOLATENE ROCHE	30MG	N017589 001
------------------------------------	------	-------------

BETAMETHASONE

CREAM; TOPICAL CELESTONE SCHERING	0.2%	N014762 001
TABLET; ORAL CELESTONE SCHERING	0.6MG	N012657 003

DISCONTINUED DRUG PRODUCT LIST

6 - 43 (of 346)

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
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
TEVA PHARMS	EQ 0.05% BASE	A071882	001	Jun 06, 1988
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	

BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION BETAMETHASONE SODIUM PHOSPHATE			
WATSON LABS	EQ 3MG BASE/ML	A085738	001
CELESTONE			
SCHERING	EQ 3MG BASE/ML	N017561	001

DISCONTINUED DRUG PRODUCT LIST

6 - 44 (of 346)

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
BETA-VAL					
TEVA	EQ 0.1% BASE		A070069	001	Dec 19, 1985
VALISONE					
SCHERING	EQ 0.1% BASE		N016740	001	

BETAXOLOL HYDROCHLORIDE

TABLET; ORAL					
KERLONE					
SANOFI AVENTIS US	10MG		N019507	001	Oct 27, 1989
	20MG		N019507	002	Oct 27, 1989

BETAXOLOL HYDROCHLORIDE; CHLORTHALIDONE

TABLET; ORAL					
KERLEDEX					
SANOFI AVENTIS US	5MG;12.5MG		N019807	001	Oct 30, 1992
	10MG;12.5MG		N019807	002	Oct 30, 1992

BETAXOLOL 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	

DISCONTINUED DRUG PRODUCT LIST

6 - 45 (of 346)

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

ABLE	5MG	A040492	001	Jul 27, 2004
	10MG	A040483	001	Jul 27, 2004
	25MG	A040485	001	Jul 27, 2004
	50MG	A040509	001	Jul 27, 2004
ACTAVIS TOTOWA	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
IVAX SUB TEVA PHARMS	25MG	A084689	001	
LANNETT	5MG	A084702	001	
	10MG	A084712	001	
	25MG	A084074	001	
SANDOZ	5MG	A084353	001	
	10MG	A084378	001	
	10MG	A084379	001	
	25MG	A084383	001	
	25MG	A084384	001	
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

BIPERIDEN LACTATE

INJECTABLE; INJECTION

AKINETON

ABBOTT

5MG/ML

N012418 002

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

FOR SOLUTION, TABLET, DELAYED RELEASE; ORAL

HALFLYTLEY

BRAINTREE

5MG,N/A;N/A,210GM;N/A,0.74GM;N/A,2.86GM N021551 002 Sep 24, 2007
;N/A,5.6GM

DISCONTINUED DRUG PRODUCT LIST

6 - 46 (of 346)

BISOPROLOL FUMARATE

TABLET; ORAL

BISOPROLOL FUMARATE

MUTUAL PHARM	5MG	A075474	001	Oct 25, 2002
	10MG	A075474	002	Oct 25, 2002

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

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				
TEVA PARENTERAL	EQ 15 UNITS BASE/VIAL	A064084	001	Jun 01, 1996
	EQ 30 UNITS BASE/VIAL	A064084	002	Jun 01, 1996

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
BAXTER HLTHCARE	50MG/ML	A070545	001	May 14, 1986
	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
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				
BAXTER HLTHCARE	200MG/100ML	N019837	002	Apr 12, 1989
	400MG/100ML	N019837	001	Apr 12, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 47 (of 346)

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION				
BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER				
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				
XIBROM				
ISTA PHARMS INC	0.09%	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				
AMBDRYL				
PARKE DAVIS	25MG	N007984	001	

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL				
AMBENYL				
FOREST LABS	12.5MG/5ML;10MG/5ML	N009319	006	Jan 10, 1984
BROMANLYL				
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

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
SANDOZ	4MG	A083215	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 48 (of 346)

BROMPHENIRAMINE MALEATE

TABLET; ORAL					
BROMPHENIRAMINE MALEATE					
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; CODEINE PHOSPHATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL					
BROMANATE DC					
ALPHARMA US PHARMS	2MG/5ML;10MG/5ML;12.5MG/5ML		A088723	001	Feb 25, 1985
DIMETANE-DC					
ROBINS AH	2MG/5ML;10MG/5ML;12.5MG/5ML		N011694	006	Mar 29, 1984
MYPHETANE DC					
WOCKHARDT	2MG/5ML;10MG/5ML;12.5MG/5ML		A088904	001	Feb 21, 1985

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; PHENYLPROPANOLAMINE HYDROCHLORIDE

ELIXIR; ORAL					
BIPHETAP					
MORTON GROVE	4MG/5ML;25MG/5ML		A088687	001	Sep 26, 1984
BROMANATE					
ALPHARMA US PHARMS	4MG/5ML;25MG/5ML		A088688	001	Feb 06, 1985
DIMETAPP					
WYETH CONS	2MG/5ML;12.5MG/5ML		N013087	003	Mar 29, 1984
TABLET, EXTENDED RELEASE; ORAL					
BROMATAPP					
COPLEY PHARM	12MG;75MG		A071099	001	Jul 02, 1987
DIMETAPP					
WYETH CONS	12MG;75MG		N012436	003	May 14, 1985

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	0.32MG/INH		N020441	003	Jun 24, 1997

DISCONTINUED DRUG PRODUCT LIST

6 - 49 (of 346)

BUDESONIDE

SPRAY, METERED; NASAL RHINOCORT	ASTRAZENECA	0 . 0 6 4 M G / I N H	N020746	0 0 2	O c t 0 1 , 1 9 9 9
------------------------------------	-------------	-----------------------	---------	-------	---------------------

BUMETANIDE

INJECTABLE; INJECTION BUMETANIDE	HOSPIRA	0 . 2 5 M G / M L	A074160	0 0 1	O c t 3 0 , 1 9 9 7
	TEVA PARENTERAL	0 . 2 5 M G / M L	A074613	0 0 1	N o v 1 8 , 1 9 9 7
TABLET; ORAL BUMEX	VALIDUS PHARMS INC	0 . 2 5 M G / M L	N018226	0 0 1	F e b 2 8 , 1 9 8 3
	VALIDUS PHARMS INC	0 . 5 M G	N018225	0 0 2	F e b 2 8 , 1 9 8 3
		1 M G	N018225	0 0 1	F e b 2 8 , 1 9 8 3
		2 M G	N018225	0 0 3	J u n 1 4 , 1 9 8 5

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION BUPIVACAINE HYDROCHLORIDE KIT	HOSPIRA	0 . 0 7 5 %	N019978	0 0 1	S e p 0 3 , 1 9 9 2
		0 . 1 1 4 %	N019978	0 0 2	S e p 0 3 , 1 9 9 2
		0 . 2 3 %	N019978	0 0 3	S e p 0 3 , 1 9 9 2

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE	HOSPIRA	0 . 2 5 % ; 0 . 0 0 5 M G / M L	A071166	0 0 1	J u n 1 6 , 1 9 8 8
		0 . 5 % ; 0 . 0 0 5 M G / M L	A071169	0 0 1	J u n 1 6 , 1 9 8 8
		0 . 7 5 % ; 0 . 0 0 5 M G / M L	A071171	0 0 1	J u n 1 6 , 1 9 8 8

BUPIVACAINE HYDROCHLORIDE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION DUOCAINE	AMPHASTAR PHARMS INC	E Q 0 . 3 7 5 % (3 7 . 5 M G / 1 0 M L) ; E Q 1 % (1 0 0 M G / 1 0 M L)	N021496	0 0 1	M a y 2 3 , 2 0 0 3
-----------------------------------	----------------------	--	---------	-------	---------------------

BUPROPION HYDROCHLORIDE

TABLET; ORAL BUPROPION HYDROCHLORIDE	SANDOZ	75 M G	A075613	0 0 2	O c t 1 0 , 2 0 0 0
		100 M G	A075613	0 0 1	O c t 1 0 , 2 0 0 0
WELLBUTRIN	GLAXOSMITHKLINE	50 M G	N018644	0 0 1	D e c 3 0 , 1 9 8 5
TABLET, EXTENDED RELEASE; ORAL BUPROPION HYDROCHLORIDE	SANDOZ	100 M G	A076845	0 0 1	J u l 1 4 , 2 0 0 5
		150 M G	A076834	0 0 1	J u l 1 4 , 2 0 0 5
		150 M G	A076845	0 0 2	J u l 1 4 , 2 0 0 5
WELLBUTRIN SR	GLAXOSMITHKLINE	50 M G	N020358	0 0 1	O c t 0 4 , 1 9 9 6
ZYBAN	GLAXOSMITHKLINE	100 M G	N020711	0 0 2	M a y 1 4 , 1 9 9 7

BUSPIRONE HYDROCHLORIDE

CAPSULE; ORAL BUSPAR	BRISTOL MYERS SQUIBB	5 M G	N021190	0 0 1	D e c 2 0 , 2 0 0 0
		7 . 5 M G	N021190	0 0 2	D e c 2 0 , 2 0 0 0
		10 M G	N021190	0 0 3	D e c 2 0 , 2 0 0 0
		15 M G	N021190	0 0 4	D e c 2 0 , 2 0 0 0

DISCONTINUED DRUG PRODUCT LIST

6 - 50 (of 346)

BUSPIRONE HYDROCHLORIDETABLET; ORAL
BUPAR

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
BUSPIRONE HYDROCHLORIDE				
ACTAVIS TOTOWA	5MG	A075388	001	May 09, 2002
	10MG	A075388	002	May 09, 2002
	15MG	A075388	003	May 09, 2002
	30MG	A078302	001	Dec 17, 2007
IVAX SUB TEVA PHARMS	5MG	A075385	001	Mar 01, 2002
	10MG	A075385	002	Mar 01, 2002
	15MG	A075385	003	Mar 01, 2002
SANDOZ	5MG	A075413	001	Mar 19, 2002
	10MG	A075413	002	Mar 19, 2002
	15MG	A075413	003	Mar 19, 2002

BUTABARBITAL SODIUMCAPSULE; 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	30MG/5ML	A085383	001
---------------------	----------	---------	-----

WOCKHARDT	30MG/5ML	A085880	001
-----------	----------	---------	-----

BUTALAN	33 . 3MG/5ML	A084723	001
---------	--------------	---------	-----

LANNETT	33 . 3MG/5ML	A084723	001
---------	--------------	---------	-----

SARISOL	30MG/5ML	A084723	001
---------	----------	---------	-----

HALSEY	30MG/5ML	A084723	001
--------	----------	---------	-----

TABLET; ORAL

BUTABARBITAL	30MG	A085550	001
--------------	------	---------	-----

BUNDY	30MG	A085550	001
-------	------	---------	-----

BUTABARBITAL SODIUM	15MG	A084292	003
---------------------	------	---------	-----

SANDOZ	15MG	A085938	001
--------	------	---------	-----

	15MG	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
------	------	---------	-----

	30MG	A088631	001
--	------	---------	-----

WATSON LABS	15MG	A085764	001
-------------	------	---------	-----

	30MG	A085772	001
--	------	---------	-----

WHITEWORTH TOWN PLSN	15MG	A083325	002
----------------------	------	---------	-----

	30MG	A083337	001
--	------	---------	-----

BUTISOL SODIUM	15MG	N000793	002
----------------	------	---------	-----

	100MG	N000793	005
--	-------	---------	-----

SARISOL NO. 1	15MG	A084719	001
---------------	------	---------	-----

HALSEY	15MG	A084719	001
--------	------	---------	-----

SARISOL NO. 2	30MG	A084719	002
---------------	------	---------	-----

SODIUM BUTABARBITAL	30MG	A083484	001
---------------------	------	---------	-----

IVAX SUB TEVA PHARMS	15MG	A084040	001
----------------------	------	---------	-----

	30MG	A084040	001
--	------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 51 (of 346)

BUTABARBITAL SODIUM

TABLET; ORAL SODIUM BUTABARBITAL LANNETT	15MG 30MG 100MG	A085849 001 A085866 001 A085881 001
MARSHALL PHARMA	16.2MG 32.4MG	A083524 001 A083858 001
WEST WARD	15MG 30MG	A085418 001 A085432 001

BUTOCONAZOLE NITRATE

CREAM; VAGINAL FEMSTAT ROCHE PALO	2%	N019215 001 Nov 25, 1985
SUPPOSITORY; VAGINAL FEMSTAT ROCHE PALO	100MG	N019359 001 Nov 25, 1985

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION BUTORPHANOL TARTRATE HOSPIRA	1MG/ML 1MG/ML 2MG/ML 2MG/ML	A075342 001 Nov 04, 1999 A075559 001 Mar 20, 2000 A075342 002 Nov 04, 1999 A075559 002 Mar 20, 2000
BUTORPHANOL TARTRATE PRESERVATIVE FREE HOSPIRA	1MG/ML 1MG/ML 2MG/ML 2MG/ML	A074620 001 Jan 22, 1997 A075170 001 Sep 28, 1998 A074620 002 Jan 22, 1997 A075170 002 Sep 28, 1998
SPRAY, METERED; NASAL STADOL BRISTOL MYERS SQUIBB	1MG/SPRAY	N019890 001 Dec 12, 1991

CABERGOLINE

TABLET; ORAL DOSTINEX PHARMACIA AND UPJOHN	0.5MG	N020664 001 Dec 23, 1996
--	-------	--------------------------

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 0.05MG	N018312 001 N018312 002
--	------------------	----------------------------

CALCIPOTRIENE

OINTMENT; TOPICAL DOVONEX LEO PHARM	0.005%	N020273 001 Dec 29, 1993
---	--------	--------------------------

DISCONTINUED DRUG PRODUCT LIST

6 - 52 (of 346)

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
 ASTRAZENECA 200 IU/ML A073690 001 Apr 14, 1995
 MIACALCIN
 NOVARTIS 100 IU/ML N017808 001 Jul 03, 1986

CALCITRIOL

INJECTABLE; INJECTION
 CALCITRIOL
 HOSPIRA 0.001MG/ML A075816 001 Jan 16, 2004
 0.002MG/ML A075816 002 Jan 16, 2004
 TEVA PARENTERAL 0.001MG/ML A075823 001 Mar 31, 2003

CALCIUM ACETATE

CAPSULE; ORAL
 PHOSLO
 FRESENIUS MEDCL EQ 84.5MG CALCIUM N021160 001 Apr 02, 2001
 EQ 169MG CALCIUM N021160 002 Apr 02, 2001
 TABLET; ORAL
 CALCIUM ACETATE
 ROXANE EQ 169MG CALCIUM A077693 001 Jan 30, 2008
 PHOSLO
 FRESENIUS MEDCL EQ 169MG CALCIUM N019976 001 Dec 10, 1990

CALCIUM CARBONATE; RISEDRONATE SODIUM

TABLET, TABLET; ORAL
 ACTONEL WITH CALCIUM (COPACKAGED)
 WARNER CHILCOTT EQ 500MG BASE,N/A;N/A,35MG N021823 001 Aug 12, 2005

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE;
 SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION
 PRISMASOL B22GK 2/2.5 IN PLASTIC CONTAINER
 GAMBRO RENAL PRODS 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML; N021703 012 Oct 10, 2008
 3.05GM/1000ML;0.157GM/1000ML;2.21GM/100
 0ML;7.07GM/1000ML
 PRISMASOL BGK 4/0 IN PLASTIC CONTAINER
 GAMBRO RENAL PRODS N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.0 N021703 005 Oct 25, 2006
 5GM/1000ML;0.314GM/1000ML;3.09GM/1000ML
 ;6.46GM/1000ML
 PRISMASOL BGK 4/3.5 IN PLASTIC CONTAINER
 GAMBRO RENAL PRODS 5.15GM/1000ML;20GM/1000ML;5.4GM/1000ML; N021703 008 Oct 25, 2006
 2.03GM/1000ML;0.314GM/1000ML;3.09GM/100
 0ML;6.46GM/1000ML
 PRISMASOL BK 0/0 IN PLASTIC CONTAINER
 GAMBRO RENAL PRODS N/A/1000ML;N/A/1000ML;5.4GM/1000ML;3.05 N021703 007 Oct 25, 2006
 GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46
 GM/1000ML
 PRISMASOL BK 4/2.5 IN PLASTIC CONTAINER
 GAMBRO RENAL PRODS 3.68GM/1000ML;N/A/1000ML;5.4GM/1000ML;3 N021703 009 Oct 25, 2006
 .05GM/1000ML;0.314GM/1000ML;3.09GM/1000
 ML;6.46GM/1000ML

DISCONTINUED DRUG PRODUCT LIST

6 - 53 (of 346)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 37MG/100ML;5GM/100ML;31MG/100ML;120MG/100ML;330MG/100ML;88MG/100ML

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 35MG/100ML;5GM/100ML;30MG/100ML;74MG/100ML;640MG/100ML;500MG/100ML;74MG/100ML

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN 29MG/100ML;2.5GM/100ML;15MG/100ML;610MG/100ML;560MG/100ML

DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B BRAUN 29MG/100ML;1.5GM/100ML;15MG/100ML;610MG/100ML;560MG/100ML

DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B BRAUN 29MG/100ML;4.25GM/100ML;15MG/100ML;610MG/100ML;560MG/100ML

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B BRAUN 26MG/100ML;1.5GM/100ML;15MG/100ML;560MG/100ML;390MG/100ML

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN 26MG/100ML;5GM/100ML;5MG/100ML;530MG/100ML;450MG/100ML

DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B BRAUN 26MG/100ML;4.25GM/100ML;15MG/100ML;560MG/100ML;390MG/100ML

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/100ML;00ML

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/100ML;62MG/100ML

DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

B BRAUN 20MG/100ML;5GM/100ML;30MG/100ML;600MG/100ML;310MG/100ML

MILES 20MG/100ML;5GM/100ML;30MG/100ML;600MG/100ML;310MG/100ML

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA 20MG/100ML;5GM/100ML;104MG/100ML;600MG/100ML;310MG/100ML
20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA 20MG/100ML;5GM/100ML;254MG/100ML;600MG/100ML;310MG/100ML

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA 20MG/100ML;5GM/100ML;254MG/100ML;600MG/100ML;310MG/100ML

DISCONTINUED DRUG PRODUCT LIST

6 - 54 (of 346)

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER
 HOSPIRA 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

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 N018899 001 Oct 31, 1983

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

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	

DISCONTINUED DRUG PRODUCT LIST

6 - 55 (of 346)

CALCIUM METRIZOATE; MEGLUMINE METRIZOATE; METRIZOATE MAGNESIUM; METRIZOATE SODIUMINJECTABLE; INJECTION
ISOPAQUE 440

GE HEALTHCARE 0.78MG/ML;75.9MG/ML;0.15MG/ML;16.6MG/ML N016847 001

CALCIUM; MEGLUMINE; METRIZOIC ACIDINJECTABLE; INJECTION
ISOPAQUE 280

GE HEALTHCARE 0.35MG/ML;140.1MG/ML;461.8MG/ML N017506 001

CANDICIDINOINTMENT; VAGINAL
VANOBID

SANOFI AVENTIS US 0.6MG/GM A061596 001

TABLET; VAGINAL
VANOBID

SANOFI AVENTIS US 3MG A061613 001

CAPTOPRILTABLET; ORAL
CAPOTENPAR PHARM 37.5MG N018343 006 Sep 17, 1986
75MG N018343 007 Jun 13, 1995
150MG N018343 004 Jun 13, 1995

Captopril

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

CLONMEL HLTHCARE

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

ENDO LABS

12.5MG A074418 001 Feb 13, 1996
25MG A074418 002 Feb 13, 1996
50MG A074418 003 Feb 13, 1996
100MG A074418 004 Feb 13, 1996

IVAX SUB TEVA PHARMS

12.5MG A074590 004 Aug 30, 1996
25MG A074590 002 Aug 30, 1996
50MG A074590 001 Aug 30, 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

TEVA

12.5MG A074433 001 Feb 13, 1996
25MG A074433 002 Feb 13, 1996

DISCONTINUED DRUG PRODUCT LIST

6 - 56 (of 346)

CAPTOPRIL

TABLET; ORAL Captopril					
TEVA	50MG	A074433	003	Feb 13, 1996	
	100MG	A074433	004	Feb 13, 1996	
TEVA PHARMS	12.5MG	A074462	001	Feb 13, 1996	
	25MG	A074462	002	Feb 13, 1996	
	50MG	A074462	003	Feb 13, 1996	
	100MG	A074462	004	Feb 13, 1996	
WATSON LABS	12.5MG	A074576	001	Apr 23, 1996	
	25MG	A074576	002	Apr 23, 1996	
	50MG	A074576	003	Apr 23, 1996	
	100MG	A074576	004	Apr 23, 1996	

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL Captopril and Hydrochlorothiazide					
ENDO LABS	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	
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	
WATSON LABS	50MG;25MG	A074832	001	Dec 29, 1997	

CARBACHOL

SOLUTION; INTRAOCCULAR 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 GEOPEN					
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 - 57 (of 346)

CARBENICILLIN DISODIUM

INJECTABLE; INJECTION
PYOPEN

GLAXOSMITHKLINE	EQ 10GM BASE/VIAL	N050298 006
	EQ 20GM BASE/VIAL	N050298 007

CARBENICILLIN INDANYL SODIUM

TABLET; ORAL
GECILLIN
PFIZER

	EQ 382MG BASE	N050435 001
--	---------------	-------------

CARBIDOPA; LEVODOPA

TABLET; ORAL
CARBIDOPA AND LEVODOPA

SANDOZ	10MG;100MG	A073586 001	Jun 29, 1995
	25MG;100MG	A073587 001	Jun 29, 1995
	25MG;250MG	A073620 001	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

CARBINOXAMINE MALEATE

ELIXIR; ORAL
CLISTIN

MCNEIL	4MG / 5ML	N008955 001
--------	-----------	-------------

TABLET; ORAL

CLISTIN

ORTHO MCNEIL PHARM	4MG	N008915 001
--------------------	-----	-------------

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

HOSPIRA	50MG/VIAL	A076473 001	Oct 27, 2004
	150MG/VIAL	A076473 002	Oct 27, 2004
	450MG/VIAL	A076473 003	Oct 27, 2004

PARAPLATIN

BRISTOL MYERS SQUIBB	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

APP PHARMS	50MG/5ML (10MG/ML)	A077247 001	Oct 21, 2004
	150MG/15ML (10MG/ML)	A077247 002	Oct 21, 2004

SAGENT PHARMS	50MG/5ML (10MG/ML)	A077096 001	Jun 14, 2005
	150MG/15ML (10MG/ML)	A077096 002	Jun 14, 2005
	450MG/45ML (10MG/ML)	A077096 003	Jun 14, 2005

TEVA PARENTERAL	50MG/5ML (10MG/ML)	A077389 001	Mar 30, 2007
	150MG/15ML (10MG/ML)	A077389 002	Mar 30, 2007
	450MG/45ML (10MG/ML)	A077389 003	Mar 30, 2007

PARAPLATIN

BRISTOL MYERS SQUIBB	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

DISCONTINUED DRUG PRODUCT LIST

6 - 58 (of 346)

CARBOPLATIN

INJECTABLE; IV (INFUSION)

PARAPLATIN

BRISTOL MYERS SQUIBB 600MG/60ML (10MG/ML)

N020452 004 Jan 15, 2004

CARISOPRODOL

CAPSULE; ORAL

SOMA

MEDA PHARMS 250MG

N011792 003

TABLET; ORAL

CARISOPRODOL

ABLE 350MG

A040421 001 Jun 21, 2001

PIONEER PHARMS 350MG

A089390 001 Oct 13, 1988

SANDOZ 350MG

A081025 001 Apr 13, 1989

350MG

A089566 001 Aug 30, 1988

WATSON LABS 350MG

A040152 001 Dec 03, 1996

350MG

A085433 001

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

N018550 002 Dec 31, 1987

150MG

N018550 003 Dec 31, 1987

CARTEOLOL HYDROCHLORIDE

TABLET; ORAL

CARTROL

ABBOTT 2.5MG

N019204 001 Dec 28, 1988

5MG

N019204 002 Dec 28, 1988

10MG

N019204 003 Dec 28, 1988

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

CLONMEL HLTHCARE 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

MARSAM PHARMS LLC EQ 250MG BASE

A064148 001 May 23, 1996

EQ 500MG BASE

A064148 002 May 23, 1996

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

DISCONTINUED DRUG PRODUCT LIST

6 - 59 (of 346)

CEFACLORFOR SUSPENSION; ORAL
CECLOR

LILLY	EQ 125MG BASE/5ML	N050522	001	
	EQ 250MG BASE/5ML	N050522	002	
CEFACLOR				
CLONMEL HLTHCARE	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
MARSAM PHARMS LLC	EQ 125MG BASE/5ML	A064204	001	Feb 18, 1998
	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, 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 HEMIHYDRATECAPSULE; ORAL
CEFADROXIL

IVAX SUB TEVA PHARMS	EQ 500MG BASE	A062766	001	Mar 03, 1987
PUREPAC PHARM	EQ 500MG BASE	A063017	001	Jan 05, 1989
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				
APOTHECON	EQ 125MG BASE/5ML	A062334	001	
	EQ 250MG BASE/5ML	A062334	002	
	EQ 500MG BASE/5ML	A062334	003	
TEVA	EQ 125MG BASE/5ML	A062698	001	Mar 01, 1989
	EQ 250MG BASE/5ML	A062698	002	Mar 01, 1989
	EQ 500MG BASE/5ML	A062698	003	Mar 01, 1989
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				
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

DISCONTINUED DRUG PRODUCT LIST

6 - 60 (of 346)

CEFAMANDOLE NAFATEINJECTABLE; 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	
	EQ 10GM BASE/VIAL	N050504	004	

CEFAZOLIN SODIUMINJECTABLE; INJECTION
ANCEF

GLAXOSMITHKLINE	EQ 250MG BASE/VIAL	N050461	001	
	EQ 500MG BASE/VIAL	N050461	002	
	EQ 1GM BASE/VIAL	N050461	003	
	EQ 5GM BASE/VIAL	N050461	004	
	EQ 10GM BASE/VIAL	N050461	005	
ANCEF IN DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	EQ 10MG BASE/ML	N050566	003	Jun 08, 1983
	EQ 20MG BASE/ML	N050566	004	Jun 08, 1983
ANCEF IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	EQ 10MG BASE/ML	N050566	001	Jun 08, 1983
	EQ 20MG BASE/ML	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	A062688	002	Nov 17, 1986
	EQ 1GM BASE/VIAL	A062688	003	Nov 17, 1986
	EQ 10GM BASE/VIAL	A062688	004	Nov 17, 1986
	EQ 20GM BASE/VIAL	A062688	005	Aug 03, 1987
BEDFORD	EQ 250MG BASE/VIAL	A062894	001	Jul 21, 1988
	EQ 500MG BASE/VIAL	A062894	002	Jul 21, 1988
	EQ 1GM BASE/VIAL	A062894	003	Jul 21, 1988
	EQ 5GM BASE/VIAL	A062894	004	Jul 21, 1988
	EQ 10GM BASE/VIAL	A062894	005	Jul 21, 1988
ELKINS SINK	EQ 250MG BASE/VIAL	A062807	001	Jan 12, 1988
	EQ 500MG BASE/VIAL	A062807	002	Jan 12, 1988
	EQ 1GM BASE/VIAL	A062807	003	Jan 12, 1988
	EQ 5GM BASE/VIAL	A062807	004	Jan 12, 1988
	EQ 10GM BASE/VIAL	A062807	005	Jan 12, 1988
	EQ 20GM BASE/VIAL	A062807	006	Jan 12, 1988
GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	A064033	001	Oct 31, 1993
MARSAM PHARMS LLC	EQ 250MG BASE/VIAL	A062988	001	Dec 29, 1989
	EQ 500MG BASE/VIAL	A062988	002	Dec 29, 1989
	EQ 1GM BASE/VIAL	A062988	003	Dec 29, 1989
	EQ 5GM BASE/VIAL	A062989	001	Dec 29, 1989
	EQ 10GM BASE/VIAL	A062989	002	Dec 29, 1989
	EQ 20GM BASE/VIAL	A062989	003	Dec 29, 1989
TEVA	EQ 250MG BASE/VIAL	A063016	001	Mar 14, 1989
	EQ 500MG BASE/VIAL	A063016	002	Mar 14, 1989
	EQ 1GM BASE/VIAL	A063016	003	Mar 14, 1989
	EQ 5GM BASE/VIAL	A063018	001	Mar 05, 1990
	EQ 10GM BASE/VIAL	A063018	002	Mar 05, 1990
KEFZOL				
ACS DOBFAR	EQ 250MG BASE/VIAL	A061773	001	
	EQ 20GM BASE/VIAL	A061773	005	Sep 08, 1987
LILLY	EQ 500MG BASE/VIAL	A062557	001	Sep 10, 1985
	EQ 1GM BASE/VIAL	A062557	002	Sep 10, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 61 (of 346)

CEFDINIR

CAPSULE; ORAL OMNICEF ABBOTT	300MG	N050739	001	Dec 04, 1997
FOR SUSPENSION; ORAL OMNICEF ABBOTT	125MG/5ML 250MG/5ML	N050749	001	Dec 04, 1997
		N050749	002	Jul 29, 2004

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
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
CEFOBID IN PLASTIC CONTAINER PFIZER	EQ 20MG BASE/ML EQ 40MG BASE/ML	N050613	002	Jul 31, 1987
		N050613	001	Jul 23, 1986

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

DISCONTINUED DRUG PRODUCT LIST

6 - 62 (of 346)

CEFORANIDE

INJECTABLE; INJECTION PRECEF					
BRISTOL	500MG/VIAL	N050554	001	May 24, 1984	
	1GM/VIAL	N050554	002	May 24, 1984	
	2GM/VIAL	N050554	003	May 24, 1984	
	10GM/VIAL	N050554	004	May 24, 1984	
	20GM/VIAL	N050554	005	May 24, 1984	

CEFOTAXIME SODIUM

INJECTABLE; INJECTION 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	
CLAFORAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER					
SANOFI AVENTIS US	EQ 20MG BASE/ML	N050596	001	May 20, 1985	
	EQ 40MG BASE/ML	N050596	003	May 20, 1985	

CEFOTETAN DISODIUM

INJECTABLE; INJECTION CEFOTAN					
ASTRAZENECA	EQ 1GM BASE/VIAL	A063293	001	Apr 29, 1993	
	EQ 1GM BASE/VIAL	N050588	001	Dec 27, 1985	
	EQ 2GM BASE/VIAL	A063293	002	Apr 29, 1993	
	EQ 2GM BASE/VIAL	N050588	002	Dec 27, 1985	
	EQ 10GM BASE/VIAL	N050588	003	Apr 25, 1988	
CEFOTAN IN PLASTIC CONTAINER					
ASTRAZENECA	EQ 20MG BASE/ML	N050694	002	Jul 30, 1993	
	EQ 40MG BASE/ML	N050694	001	Jul 30, 1993	

CEFOTIAM HYDROCHLORIDE

INJECTABLE; INJECTION CERADON					
TAKEDA	EQ 1GM BASE/VIAL	N050601	001	Dec 30, 1988	

CEFOXITIN SODIUM

INJECTABLE; INJECTION MEFOXIN					
BIONICHE PHARMA	EQ 1GM BASE/VIAL	A062757	001	Jan 08, 1987	
	EQ 2GM BASE/VIAL	A062757	002	Jan 08, 1987	
MYLAN INSTITUTIONAL	EQ 1GM BASE/VIAL	N050517	001		
	EQ 2GM BASE/VIAL	N050517	002		
	EQ 10GM BASE/VIAL	N050517	003		
MEFOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER					
MERCK	EQ 20MG BASE/ML	N050581	003	Sep 20, 1984	
	EQ 40MG BASE/ML	N050581	004	Sep 20, 1984	
MEFOXIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER					
MERCK	EQ 20MG BASE/ML	N050581	002	Sep 20, 1984	
	EQ 40MG BASE/ML	N050581	001	Sep 20, 1984	

CEFPIRAMIDE SODIUM

INJECTABLE; INJECTION CEFPIRAMIDE SODIUM					
WYETH AYERST	EQ 1GM BASE/VIAL	N050633	002	Jan 31, 1989	
	EQ 2GM BASE/VIAL	N050633	003	Jan 31, 1989	
	EQ 10GM BASE/VIAL	N050633	005	Jan 31, 1989	

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL BANAN					
SANKYO	EQ 50MG BASE/5ML	N050688	002	Aug 07, 1992	
	EQ 100MG BASE/5ML	N050688	001	Aug 07, 1992	

DISCONTINUED DRUG PRODUCT LIST

6 - 63 (of 346)

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL					
VANTIN					
PHARMACIA AND UPJOHN	EQ 50MG BASE/5ML	N050675	001	Aug 07, 1992	
	EQ 100MG BASE/5ML	N050675	002	Aug 07, 1992	
TABLET; ORAL					
BANAN					
SANKYO	EQ 100MG BASE	N050687	001	Aug 07, 1992	
	EQ 200MG BASE	N050687	002	Aug 07, 1992	
VANTIN					
PHARMACIA AND UPJOHN	EQ 100MG BASE	N050674	001	Aug 07, 1992	
	EQ 200MG BASE	N050674	002	Aug 07, 1992	

CEFPROZIL

FOR SUSPENSION; ORAL					
CEFZIL					
BRISTOL MYERS SQUIBB	125MG/5ML	N050665	001	Dec 23, 1991	
	250MG/5ML	N050665	002	Dec 23, 1991	
TABLET; ORAL					
CEFZIL					
BRISTOL MYERS SQUIBB	250MG	N050664	001	Dec 23, 1991	
	500MG	N050664	002	Dec 23, 1991	

CEFTAZIDIME

INJECTABLE; INJECTION					
CEPTAZ					
GLAXOSMITHKLINE	500MG/VIAL	N050646	001	Sep 27, 1990	
	1GM/VIAL	N050646	002	Sep 27, 1990	
	2GM/VIAL	N050646	003	Sep 27, 1990	
	10GM/VIAL	N050646	004	Sep 27, 1990	
PENTACEF					
GLAXOSMITHKLINE	1GM/VIAL	A063322	001	Nov 07, 1995	
	1GM/VIAL	A064006	001	Mar 31, 1992	
	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					
GLAXOSMITHKLINE	EQ 10MG BASE/ML	N050634	001	Apr 28, 1989	

CEFTIBUTEN DIHYDRATE

FOR SUSPENSION; ORAL					
CEDAX					
PERNIX THERAP	EQ 180MG BASE/5ML	N050686	002	Dec 20, 1995	

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION					
CEFIZOX					
ASTELLAS	EQ 500MG BASE/VIAL	N050560	001	Sep 15, 1983	
	EQ 1GM BASE/VIAL	A063294	002	Mar 31, 1994	

DISCONTINUED DRUG PRODUCT LIST

6 - 64 (of 346)

CEFTIZOXIME SODIUMINJECTABLE; INJECTION
CEFIZOX

ASTELLAS	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

CEFTRIAZONE SODIUMINJECTABLE; *****
ROCEPHIN

HOFFMANN LA ROCHE	EQ 500MG BASE/VIAL	N050585	002	Dec 21, 1984
	EQ 1GM BASE/VIAL	N050585	003	Dec 21, 1984
	EQ 2GM BASE/VIAL	N050585	004	Dec 21, 1984

INJECTABLE; IM-IV

ROCEPHIN

HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL	N050585	001	Dec 21, 1984
-------------------	--------------------	---------	-----	--------------

INJECTABLE; INJECTION
ROCEPHIN

HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL	A063239	001	Aug 13, 1993
	EQ 500MG BASE/VIAL	A062654	001	Apr 30, 1987
	EQ 1GM BASE/VIAL	A062654	002	Apr 30, 1987
	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

ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER

HOFFMANN LA ROCHE	EQ 10MG BASE/ML	N050624	001	Feb 11, 1987
	EQ 20MG BASE/ML	N050624	002	Feb 11, 1987
	EQ 40MG BASE/ML	N050624	003	Feb 11, 1987

CEFTRIAZONE SODIUM; LIDOCAINEINJECTABLE; INJECTION
ROCEPHIN KIT

HOFFMANN LA ROCHE	EQ 500MG BASE/VIAL,N/A;N/A,1%	N050585	007	May 08, 1996
	EQ 1GM BASE/VIAL,N/A;N/A,1%	N050585	006	May 08, 1996

CEFUROXIME AXETILTABLET; ORAL
CEFUROXIME AXETIL

SANDOZ	EQ 250MG BASE	A065126	001	Oct 28, 2003
	EQ 500MG BASE	A065126	002	Oct 28, 2003

CEFUROXIME SODIUM

INJECTABLE; IM-IV

CEFUROXIME SODIUM MARSAM PHARMS LLC	EQ 750MG BASE/VIAL	A064035	001	Feb 26, 1993
KEFUROX ACS DOBFAR	EQ 750MG BASE/VIAL	A062591	001	Jan 10, 1986

INJECTABLE; INJECTION CEFUROXIME SODIUM MARSAM PHARMS LLC	EQ 1.5GM BASE/VIAL	A064035	002	Feb 26, 1993
	EQ 7.5GM BASE/VIAL	A064036	001	Feb 26, 1993

KEFUROX ACS DOBFAR	EQ 1.5GM BASE/VIAL	A062591	002	Jan 10, 1986
	EQ 7.5GM BASE/VIAL	A062591	003	Dec 17, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 65 (of 346)

CEFUROXIME SODIUM

INJECTABLE; INJECTION					
KEFUROX					
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
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
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 500MG BASE		A062869	001	Mar 17, 1988
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
	EQ 500MG BASE		A062823	001	Feb 05, 1988
YOSHITOMI	EQ 250MG BASE		A062872	001	Jun 20, 1988
	EQ 500MG BASE		A062871	001	Jul 05, 1988
KEFLEX					
SHIONOGI INC	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	
TEVA	EQ 125MG BASE/5ML		A062767	001	Jun 16, 1987
	EQ 125MG BASE/5ML		A062873	001	May 23, 1988
	EQ 250MG BASE/5ML		A062768	001	Jun 16, 1987
	EQ 250MG BASE/5ML		A062867	001	Apr 15, 1988
VITARINE	EQ 125MG BASE/5ML		A062779	001	Dec 22, 1987
	EQ 250MG BASE/5ML		A062781	001	Dec 22, 1987
KEFLEX					
LEX PHARMS	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		A062826	001	Aug 17, 1987
	EQ 500MG BASE		A062827	001	Aug 17, 1987
VITARINE	EQ 250MG BASE		A062863	001	Aug 11, 1988

DISCONTINUED DRUG PRODUCT LIST

6 - 66 (of 346)

CEPHALEXIN

TABLET; ORAL CEPHALEXIN VITARINE	EQ 500MG BASE EQ 1GM BASE	A062863 002 A062863 003	Aug 11, 1988 Aug 11, 1988
KEFLET LILLY	EQ 250MG BASE EQ 250MG BASE EQ 500MG BASE EQ 500MG BASE EQ 1GM BASE	A062745 001 N050440 003 A062745 002 N050440 001 N050440 002	Dec 01, 1986 Feb 26, 1987 Dec 01, 1986
TABLET, FOR SUSPENSION; ORAL PANIXINE DISPERDOSE	RANBAXY LABS LTD	EQ 125MG BASE EQ 250MG BASE	A065100 002 A065100 001
			Sep 11, 2003 Sep 11, 2003

CEPHALEXIN HYDROCHLORIDE

TABLET; ORAL KEFTAB LILLY	EQ 250MG BASE EQ 333MG BASE EQ 500MG BASE	N050614 001 N050614 003 N050614 002	Oct 29, 1987 May 16, 1988 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 A062426 002 A062426 003 A062426 004	May 03, 1985 May 03, 1985 May 03, 1985 May 03, 1985
CEPHALOTHIN SODIUM ABBOTT	EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL	A062547 001 A062548 001 A062547 002 A062548 002	Sep 11, 1985 Sep 11, 1985 Sep 11, 1985 Sep 11, 1985
ABRAXIS PHARM	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	A062666 002 A062666 001	Jun 10, 1987 Jun 10, 1987
BRISTOL	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 4GM BASE/VIAL	A062464 001 A062464 002 A062464 003	May 07, 1984 May 07, 1984 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 A062422 005 A062730 001 A062422 004 A062422 006 A062730 002	Jan 31, 1984 Jul 16, 1991 Mar 05, 1987 Jan 31, 1984 Jul 16, 1991 Mar 05, 1987
CEPHALOTHIN SODIUM W/ SODIUM CHLORIDE IN PLASTIC CONTAINER BAXTER HLTHCARE	EQ 20MG BASE/ML EQ 40MG BASE/ML	A062422 001 A062422 002	Jan 31, 1984 Jan 31, 1984
KEFLIN LILLY	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 4GM BASE/VIAL EQ 20GM BASE/VIAL	N050482 001 N050482 002 N050482 003 N050482 007	
KEFLIN IN PLASTIC CONTAINER LILLY	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	A062549 001 A062549 002	Sep 10, 1985 Sep 10, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 67 (of 346)

CEPHALOTHIN SODIUMINJECTABLE; INJECTION
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 SODIUMINJECTABLE; 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
BAXTER HLTHCARE	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

CEPHRADINECAPSULE; 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	
	250MG/5ML	A061866	002	

CEPHRADINE

BARR	125MG/5ML	A062858	001	May 19, 1988
	250MG/5ML	A062859	001	May 19, 1988

DISCONTINUED DRUG PRODUCT LIST

6 - 68 (of 346)

CEPHRADINE

FOR SUSPENSION; ORAL					
CEPHRADINE					
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 SODIUM

TABLET; 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 DIETHYLAMINE

INJECTABLE; 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
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY					
ACTAVIS MID ATLANTIC	5MG/5ML		A090378	002	May 09, 2008
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF					
ACTAVIS MID ATLANTIC	5MG/5ML		A090378	001	May 09, 2008

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					
SIGMA TAU	250MG		N018513	002	Jul 28, 1983

CHLOPHEDIANOL HYDROCHLORIDE

SYRUP; ORAL					
ULO					
3M	25MG/5ML		N012126	001	

CHLORAMPHENICOL

CAPSULE; ORAL					
AMPHICOL					
FERRANTE	100MG		A060058	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 69 (of 346)

CHLORAMPHENICOL

CAPSULE; ORAL AMPHICOL FERRANTE	250MG	A060058	002
CHLORAMPHENICOL IVAX SUB TEVA PHARMS	250MG	A062247	001
CHLOROMYCETIN PARKEDALE	50MG	A060591	001
	100MG	A060591	003
	250MG	A060591	002
MYCHEL ARMENPHARM	250MG	A060851	001
CREAM; TOPICAL CHLOROMYCETIN PARKE DAVIS	1%	N050183	001
FOR SOLUTION; OPHTHALMIC CHLOROMYCETIN PARKEDALE	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
CHLOROMYCETIN PARKEDALE	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
CHLOROFAIR PHARMAFAIR	0.5%	A062437	001
CHLOROPTIC ALLERGAN	0.5%	N050091	001
ECONOCHLOR ALCON	0.5%	A061645	001
OPHTHOCHLOR PARKEDALE	0.5%	A061220	001
OPTOMYCIN OPTOPICS	0.5%	A062171	001
SOLUTION/DROPS; OTIC CHLOROMYCETIN PARKEDALE	0.5%	N050205	001

CHLORAMPHENICOL PALMITATE

SUSPENSION; ORAL CHLOROMYCETIN PALMITATE PARKE DAVIS	EQ 150MG BASE/5ML	A062301	001
	EQ 150MG BASE/5ML	N050152	001

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	

DISCONTINUED DRUG PRODUCT LIST

6 - 70 (of 346)

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION
 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
 PARKADEALE 12.5MG/VIAL;25MG/VIAL N050202 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC
 OPHTHOCORT
 PARKADEALE 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	

DISCONTINUED DRUG PRODUCT LIST

6 - 71 (of 346)

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE				
IMPAK LABS	5MG	A086213	001	
	10MG	A085113	001	
	25MG	A086212	001	
IVAX SUB TEVA PHARMS	5MG	A083741	001	
	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	
SANDOZ	5MG	A084678	001	
	5MG	A084919	001	
	10MG	A084041	001	
	10MG	A084920	001	
	25MG	A084679	002	
	25MG	A084823	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
USL PHARMA	5MG	A084644	001	
	25MG	A084645	001	
VANGARD	5MG	A088129	001	Mar 28, 1983
	10MG	A088010	001	Mar 28, 1983
	25MG	A088130	001	Mar 28, 1983
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 PHARM INTL	100MG/AMP	N012301	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 72 (of 346)

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL				
MENRIUM 10-4				
ROCHE	10MG; 0.4MG		N014740	006
MENRIUM 5-2				
ROCHE	5MG; 0.2MG		N014740	002
MENRIUM 5-4				
ROCHE	5MG; 0.4MG		N014740	004

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL				
EXIDINE				
XTTRIUM	2.5%		N019421	001
MICRODERM				Dec 17, 1985
J AND J	4%		A072255	001
PREVACARE R				Apr 15, 1991
J AND J	0.5%		A072292	001
STERI-STAT				Jan 28, 1992
MATRIX MEDCL	4%		A070104	001
SPONGE; TOPICAL				Jul 24, 1986
CHLORHEXIDINE GLUCONATE				
KENDALL IL	4%		N019490	001
E-Z SCRUB				Mar 27, 1987
BECTON DICKINSON	4%		A073416	001
HIBICLENS				Mar 14, 2000
MOLNLYCKE HLTH	4%		N018423	001
MICRODERM				
J AND J	4%		A072295	001
				Feb 28, 1991

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-MPF				
APP PHARMS	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				
CHLOROQUINE PHOSPHATE				
MD PHARM	EQ 150MG BASE		A087228	001
PUREPAC PHARM	EQ 150MG BASE		A080886	001
TEVA	EQ 150MG BASE		A087504	001
WATSON LABS	EQ 150MG BASE		A087979	001
	EQ 300MG BASE		A088030	001
				Jan 13, 1982
				Dec 21, 1982
				Dec 21, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 73 (of 346)

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET; ORAL

ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE

SANOFI AVENTIS US EQ 300MG BASE;EQ 45MG BASE

N014860 002

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

ABC HOLDING	250MG	A085569	001
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			
LUNDBECK INC	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
250MG;250MGA070783 001 Nov 06, 1987
A070654 001 Nov 06, 1987CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE AND RESERPINE

WEST WARD 250MG;0.125MG
500MG;0.125MGA088557 001 Dec 22, 1983
A088365 001 Dec 22, 1983

CHLOROTHIAZIDE W/ RESERPINE

WATSON LABS 250MG;0.125MG
500MG;0.125MGA084853 001
A088151 001 Jun 09, 1983

CHLOROTHIAZIDE-RESERPINE

MYLAN 250MG;0.125MG
500MG;0.125MGA087744 001 May 06, 1982
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
25MG

N008102 004

72MG

N011444 001

N016235 001

CHLOROXINE

SHAMPOO; TOPICAL

CAPITROL

WESTWOOD SQUIBB 2%

N017594 001

DISCONTINUED DRUG PRODUCT LIST

6 - 74 (of 346)

CHLORPHENESIN CARBAMATE

TABLET; ORAL
MAOLATE
PHARMACIA AND UPJOHN 400MG N014217 002

CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL
CHLORPHENIRAMINE MALEATE
SANDOZ 12MG A070797 001 Aug 12, 1988
TELDRIN
GLAXOSMITHKLINE 8MG N017369 001
12MG N017369 002

INJECTABLE; INJECTION

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

CHLOR-TRIMETON

SCHERING PLOUGH 10MG/ML N008826 001
100MG/ML N008794 001

PYRIDAMAL 100

BEL MAR 100MG/ML A083733 001

SYRUP; ORAL

CHLORPHENIRAMINE MALEATE
PHARM ASSOC 2MG/5ML A087520 001 Feb 10, 1982
CHLOR-TRIMETON

SCHERING 2MG/5ML N006921 006

TABLET; ORAL

ANTAGONATE
BAYER PHARMS 4MG A083381 001

CHLORPHENIRAMINE MALEATE

ANABOLIC 4MG A083078 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
MUTUAL PHARM 4MG A080700 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
SANDOZ 4MG A080961 001
VITARINE 4MG A085837 001
WATSON LABS 4MG A080696 001
4MG A080791 001
4MG A085139 001
WEST WARD 4MG A083787 001

CHLOR-TRIMETON

SCHERING 4MG N006921 002

KLOROMIN

HALSEY 4MG A083629 001

PHENETRON

LANNETT 4MG A080846 001

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

SCHERING PLOUGH 8MG N007638 001

DISCONTINUED DRUG PRODUCT LIST

6 - 75 (of 346)

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL
 EFIDAC 24 CHLORPHENIRAMINE MALEATE
 ALZA 16MG N019746 002 Nov 18, 1994

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL CHLORPHENIRAMINE MALEATE AND PHENYLPROPANOLAMINE HYDROCHLORIDE WATSON LABS 12MG;75MG A088681 001 Sep 29, 1987
CHLORPHENIRAMINE MALEATE AND PHENYLPROPANOLAMINE HYDROCHLORIDE SANDOZ 12MG;75MG A088940 001 Jan 26, 1989
COLD CAPSULE IV GRAHAM DM 12MG;75MG N018793 001 Apr 25, 1985
COLD CAPSULE V GRAHAM DM 8MG;75MG N018794 001 Apr 23, 1985
CONTAC 12 HOUR GLAXOSMITHKLINE 8MG;75MG N018099 001
DRIZE ASCHER 12MG;75MG A088359 001 Feb 13, 1986
ORNDAE GLAXOSMITHKLINE 12MG;75MG N012152 004
PHENYLPROPANOLAMINE HYDROCHLORIDE W/ CHLORPHENIRAMINE MALEATE CENT PHARMS 8MG;75MG N018809 001 May 07, 1984
TABLET, EXTENDED RELEASE; ORAL CONTAC NOVARTIS 12MG;75MG N019613 001 Jun 13, 1986
DEMAZIN SCHERING PLOUGH 4MG;25MG N018556 001 May 14, 1984
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 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

CHLORPHENIRAMINE POLISTIREX; PHENYLPROPANOLAMINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL CORSYM UCB INC EQ 4MG MALEATE/5ML;EQ 37.5MG HCL/5ML N018050 001 Jan 04, 1984

CHLORPHENTERMINE HYDROCHLORIDE

TABLET; ORAL PRE-SATE PARKE DAVIS EQ 65MG BASE N014696 001
--

DISCONTINUED DRUG PRODUCT LIST

6 - 76 (of 346)

CHLORPROMAZINE

SUPPOSITORY; RECTAL		
THORAZINE		
GLAXOSMITHKLINE	25MG	N009149 024
	100MG	N009149 033

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL			
THORAZINE			
GLAXOSMITHKLINE	30MG	N011120 016	
	75MG	N011120 017	
	150MG	N011120 018	
	200MG	N011120 019	
	300MG	N011120 020	
CONCENTRATE; ORAL			
CHLORPROMAZINE HYDROCHLORIDE			
ACTAVIS MID ATLANTIC	100MG/ML	A086863 001	
PHARM ASSOC	30MG/ML	A040231 001	Dec 30, 1999
	100MG/ML	A040224 001	Jan 26, 1999
WOCKHARDT	30MG/ML	A087032 001	Jul 08, 1982
	100MG/ML	A087053 001	
CHLORPROMAZINE HYDROCHLORIDE INTENSOL			
ROXANE	30MG/ML	A088157 001	Apr 27, 1983
	100MG/ML	A088158 001	Apr 27, 1983
SONAZINE			
SANDOZ	30MG/ML	A080983 004	
	100MG/ML	A080983 005	
THORAZINE			
GLAXOSMITHKLINE	30MG/ML	N009149 032	
	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			
SANDOZ	10MG/5ML	A083040 001	
THORAZINE			
GLAXOSMITHKLINE	10MG/5ML	N009149 022	
TABLET; ORAL			
CHLORPROMAZINE HYDROCHLORIDE			
ABBOTT	10MG	A084414 001	
	25MG	A084415 001	
	50MG	A084411 001	
	100MG	A084412 001	
	200MG	A084413 001	
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 77 (of 346)

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

LEDERLE	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	
ROXANE	10MG	A085331	001	
	25MG	A085331	002	
	50MG	A085331	003	
	100MG	A085331	004	
	200MG	A085331	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	
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

BARR	100MG	A088812	001	Oct 19, 1984
	100MG	A089446	001	Nov 17, 1986
	250MG	A088813	001	Oct 19, 1984
	250MG	A089447	001	Nov 17, 1986
CLONMEL HLTHCARE	100MG	A089561	001	Sep 04, 1987
	250MG	A089562	001	Sep 04, 1987
DURAMED PHARMS BARR	100MG	A088918	001	Oct 16, 1984
	250MG	A088919	001	Oct 16, 1984
HALSEY	100MG	A089321	001	Jan 16, 1986
	250MG	A088662	001	Jan 09, 1986
IVAX PHARMS	100MG	A088840	001	Oct 25, 1984
	250MG	A087353	001	
PAR PHARM	100MG	A088175	001	Feb 27, 1984
	250MG	A088176	001	Feb 27, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 78 (of 346)

CHLORPROPAMIDE

TABLET; ORAL CHLORPROPAMIDE					
SANDOZ	100MG	A088725	001	Aug 31, 1984	
	250MG	A084669	001		
	250MG	A088726	001	Aug 31, 1984	
SUPERPHARM	100MG	A088694	001	Sep 17, 1984	
	250MG	A088695	001	Sep 17, 1984	
TEVA	100MG	A088768	001	Oct 11, 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	
GLUCAMIDE					
TEVA	250MG	A088641	001	Oct 11, 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		

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	
CLONMEL HLTHCARE	25MG	A087451	001		
	50MG	A087450	001		
IVAX PHARMS	25MG	A087555	001		
	25MG	A088164	001	Jan 09, 1984	
	50MG	A087176	001		
	50MG	A087947	001	Feb 27, 1984	
KV PHARM	25MG	A087311	001		
	50MG	A087312	001		
MUTUAL PHARM	25MG	A087292	001		
	25MG	A089285	001	Jul 21, 1986	
	25MG	A089738	001	Sep 19, 1988	
	50MG	A087293	001		
	50MG	A089286	001	Jul 21, 1986	
	50MG	A089739	001	Sep 19, 1988	
PIONEER PHARMS	50MG	A089591	001	Jul 21, 1988	
PUREPAC PHARM	25MG	A088139	001	Jul 16, 1986	
	50MG	A088140	001	Aug 11, 1983	
SANDOZ	25MG	A087380	001		
	50MG	A087118	001		

DISCONTINUED DRUG PRODUCT LIST

6 - 79 (of 346)

CHLORTHALIDONE

TABLET; ORAL					
CHLORTHALIDONE					
SANDOZ	50MG		A087381	001	
SUPERPHARM	25MG		A087473	001	Feb 09, 1983
	50MG		A087247	001	Feb 09, 1983
TEVA	50MG		A088651	001	May 30, 1985
USL PHARMA	25MG		A089051	001	Jun 01, 1987
	50MG		A089052	001	Jun 01, 1987
VANGARD	25MG		A088012	001	Jul 14, 1982
	50MG		A088073	001	Mar 25, 1983
WARNER CHILCOTT	25MG		A087515	001	Jan 24, 1983
	50MG		A087516	001	Feb 09, 1983
WATSON LABS	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					
MONARCH PHARMS	25MG		A088051	001	Nov 12, 1982
	25MG		N019574	002	Feb 12, 1992

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

CHLORTHALIDONE; METOPROLOL TARTRATE

CAPSULE; ORAL					
LOPRESSIDONE					
NOVARTIS	25MG;100MG		N019451	001	Dec 31, 1987
	25MG;200MG		N019451	002	Dec 31, 1987

CHLORTHALIDONE; RESERPINE

TABLET; ORAL					
DEMI-REGROTON					
SANOFI AVENTIS US	25MG;0.125MG		N015103	002	
REGROTON					
SANOFI AVENTIS US	50MG;0.25MG		N015103	001	

CHLORZOXAZONE

TABLET; ORAL					
CHLORZOXAZONE					
ACTAVIS TOTOWA	250MG		A088928	001	May 08, 1987
	500MG		A040113	001	Sep 29, 1995
MUTUAL PHARM	500MG		A089970	001	Sep 27, 1990
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

DISCONTINUED DRUG PRODUCT LIST

6 - 80 (of 346)

CHLORZOXAZONE

TABLET; ORAL CHLORZOXAZONE					
SANDOZ	250MG		A089852	001	May 04, 1988
	500MG		A089853	001	May 04, 1988
WATSON LABS	250MG		A086901	001	
	250MG		A086948	001	Aug 09, 1982
	500MG		A081019	001	Jul 29, 1991
PARAFLEX					
ORTHO MCNEIL PHARM	250MG		N011300	003	
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					
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
TEVA PHARMS	EQ 4GM RESIN/PACKET		A074554	001	Oct 02, 1996
	EQ 4GM RESIN/SCOOPFUL		A074554	002	Oct 02, 1996
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
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
TABLET; ORAL QUESTRAN					
APOTHECON	EQ 800MG RESIN		A073403	002	Dec 27, 1999
	EQ 1GM RESIN		A073403	001	Apr 28, 1994

CHORIOGONADOTROPIN ALFA

INJECTABLE; INJECTION OVIDREL					
EMD SERONO	0.25MG/VIAL		N021149	001	Sep 20, 2000

CHROMIC CHLORIDE

INJECTABLE; INJECTION CHROMIC CHLORIDE					
ABRAXIS PHARM	EQ 0.004MG CHROMIUM/ML		N019271	001	May 05, 1987

CHROMIC PHOSPHATE P-32

INJECTABLE; INJECTION PHOSPHOCOL P32					
MALLINCKRODT	5mCi/ML		N017084	001	

CHYMOPAPAIN

INJECTABLE; INJECTION CHYMODIACTIN					
CHART MEDCL	4,000 UNITS/VIAL		N018663	002	Aug 21, 1984
	10,000 UNITS/VIAL		N018663	001	Nov 10, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 81 (of 346)

CHYMOPAPAIN

INJECTABLE; INJECTION
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

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION
PRIMAXIN
MERCK EQ 750MG BASE/VIAL; 750MG/VIAL N050630 002 Dec 14, 1990

CILOSTAZOL

TABLET; ORAL
CILOSTAZOL
ACTAVIS TOTOWA 100MG A077028 002 Nov 26, 2004
IVAX SUB TEVA PHARMS 100MG A077020 002 Mar 01, 2005
MUTUAL PHARM 50MG A077208 002 Mar 29, 2006
100MG A077208 001 Mar 29, 2006

CIMETIDINE

SUSPENSION; ORAL
TAGAMET HB 200
GLAXOSMITHKLINE 200MG/20ML N020951 001 Jul 09, 1999

TABLET; ORAL CIMETIDINE ENDO PHARMS	200MG 300MG 400MG 800MG	A074281 001 A074281 002 A074281 003 A074329 001	May 17, 1994 May 17, 1994 May 17, 1994 May 17, 1994
IVAX SUB TEVA PHARMS	200MG 300MG 400MG 800MG	A074401 001 A074401 002 A074401 003 A074402 001	May 30, 1995 May 30, 1995 May 30, 1995 May 30, 1995
LEK PHARMS	100MG 200MG 200MG 300MG 400MG 800MG	A075122 001 A074250 001 A075122 002 A074250 002 A074250 003 A074250 004	Jun 19, 1998 Jun 29, 1995 Jun 19, 1998 Jun 29, 1995 Jun 29, 1995 Jun 29, 1995
PERRIGO	100MG	A074972 001	Jun 19, 1998
ROXANE	300MG 400MG 800MG	A074361 001 A074361 002 A074371 001	Dec 23, 1994 Dec 23, 1994 Dec 23, 1994
SANDOZ	200MG 300MG 400MG 800MG	A074100 001 A074100 002 A074100 003 A074100 004	Jan 31, 1995 Jan 31, 1995 Jan 31, 1995 Jan 31, 1995
TEVA	200MG 300MG 400MG 800MG	A074365 001 A074365 002 A074365 003 A074365 004	Feb 28, 1995 Feb 28, 1995 Feb 28, 1995 Feb 28, 1995
TAGAMET HB GLAXOSMITHKLINE	100MG	N020238 001	Jun 19, 1995

DISCONTINUED DRUG PRODUCT LIST

6 - 82 (of 346)

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

ENDO PHARMS	EQ 300MG BASE/2ML	A074005	001	Aug 31, 1994
HOSPIRA	EQ 300MG BASE/2ML	A074296	001	Mar 28, 1997
	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
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				
ACTAVIS MID ATLANTIC	EQ 300MG BASE/5ML	A074176	001	Jun 01, 1994
DURAMED PHARMS BARR	EQ 300MG BASE/5ML	A075110	001	Jun 18, 1998
ENDO PHARMS	EQ 300MG BASE/5ML	A074251	001	Dec 22, 1994
ROXANE	EQ 300MG BASE/5ML	A074541	001	Aug 05, 1997
TEVA PHARMS	EQ 300MG BASE/5ML	A074859	001	Jul 09, 1998
TAGAMET				
GLAXOSMITHKLINE	EQ 300MG BASE/5ML	N017924	001	

CINOXACIN

CAPSULE; ORAL

CINOBAC

LILLY

250MG

N018067 001

500MG

N018067 002

CINOXACIN

TEVA

250MG

A073005 001

Feb 28, 1992

500MG

A073006 001

Feb 28, 1992

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

BAYER HLTHCARE

1200MG/120ML (10MG/ML)

N019847 003

Dec 26, 1990

CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

BAYER PHARMS

200MG/100ML

N019858 001

Dec 26, 1990

CIPROFLOXACIN

APP PHARMS

200MG/20ML (10MG/ML)

A076484 001

Aug 28, 2006

400MG/40ML (10MG/ML)

A076484 002

Aug 28, 2006

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

CIPROFLOXACIN IN DEXTROSE 5%

HIKMA FARMACEUTICA

200MG/100ML

A076757 001

Apr 21, 2008

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

BEDFORD

200MG/100ML

A078114 001

Mar 18, 2008

400MG/200ML

A078114 002

Mar 18, 2008

TEVA PHARMS

200MG/100ML

A077138 001

Mar 18, 2008

400MG/200ML

A077138 002

Mar 18, 2008

DISCONTINUED DRUG PRODUCT LIST

6 - 83 (of 346)

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

BARR	EQ 250MG BASE	A074124	001	Jun 09, 2004
	EQ 500MG BASE	A074124	002	Jun 09, 2004
	EQ 750MG BASE	A074124	003	Jun 09, 2004
NOSTRUM LABS	EQ 250MG BASE	A076138	001	Jun 09, 2004
	EQ 500MG BASE	A076138	002	Jun 09, 2004
	EQ 750MG BASE	A076138	003	Jun 09, 2004
SANDOZ	EQ 100MG BASE	A075939	001	Mar 03, 2005
	EQ 250MG BASE	A075939	002	Jun 09, 2004
	EQ 250MG BASE	A076593	002	Jun 09, 2004
	EQ 500MG BASE	A075939	003	Jun 09, 2004
	EQ 500MG BASE	A076593	003	Jun 09, 2004
	EQ 750MG BASE	A075939	004	Jun 09, 2004
	EQ 750MG BASE	A076593	004	Jun 09, 2004
TEVA	EQ 250MG BASE	A076136	001	Jun 09, 2004
	EQ 500MG BASE	A076136	002	Jun 09, 2004
	EQ 750MG BASE	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

CIPROFLOXACIN EXTENDED RELEASE

SANDOZ	212.6MG;EQ 287.5MG BASE	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	N020210	001	Jul 29, 1993
	EQ 20MG BASE	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	A074713	001	Nov 14, 2000
	50MG/VIAL	A074713	002	Nov 14, 2000
PLATINOL				
CORDEN PHARMA	10MG/VIAL	N018057	001	
	50MG/VIAL	N018057	002	
PLATINOL-AQ				
CORDEN PHARMA	0.5MG/ML	N018057	003	Jul 18, 1984
	1MG/ML	N018057	004	Nov 08, 1988

CITALOPRAM HYDROBROMIDE

SOLUTION; ORAL

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
MUTUAL PHARM	EQ 10MG BASE	A077052	001	Jul 03, 2006

DISCONTINUED DRUG PRODUCT LIST

6 - 84 (of 346)

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

MUTUAL PHARM	EQ 20MG BASE	A077052	002	Jul 03, 2006
	EQ 40MG BASE	A077052	003	Jul 03, 2006
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	A077040	001	Aug 17, 2005
	EQ 20MG BASE	A077040	002	Aug 17, 2005
	EQ 40MG BASE	A077040	003	Aug 17, 2005
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
TABLET, ORALLY DISINTEGRATING; ORAL				
CITALOPRAM HYDROBROMIDE				
BIOVAI 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
---------	--------------------------------------	---------	-----	--------------

CLARITHROMYCIN

FOR SUSPENSION; ORAL

BIAxin

ABBOTT	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

TABLET, EXTENDED RELEASE; ORAL

CLARITHROMYCIN

RANBAXY	1GM	A065210	001	Jan 26, 2005
---------	-----	---------	-----	--------------

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

ACTAVIS MID ATLANTIC	EQ 0.5MG BASE/5ML	A074075	001	Oct 31, 1993
----------------------	-------------------	---------	-----	--------------

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

TEVA	1.34MG	A073282	001	Jan 31, 1992
------	--------	---------	-----	--------------

	1.34MG	A073282	002	Dec 03, 1992
--	--------	---------	-----	--------------

TAVIST

NOVARTIS	2.68MG	N017661	001	
----------	--------	---------	-----	--

TAVIST-1

NOVARTIS	1.34MG	N017661	002	
----------	--------	---------	-----	--

	1.34MG	N017661	003	Aug 21, 1992
--	--------	---------	-----	--------------

CLEMASTINE FUMARATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TAVIST-D

NOVARTIS	EQ 1MG BASE;75MG	N018298	001	Dec 15, 1982
----------	------------------	---------	-----	--------------

	1.34MG;75MG	N018298	002	Aug 21, 1992
--	-------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 85 (of 346)

CLEMASTINE FUMARATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
 TAVIST-D
 NOVARTIS 1.34MG;75MG N020640 001 Aug 09, 1996

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
 TEVA EQ 75MG BASE A063027 001 Sep 20, 1989
 WATSON LABS EQ 75MG BASE A063082 001 Jul 31, 1991

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
 ASTRAZENECA EQ 150MG BASE/ML A062928 001 Feb 13, 1989
 BAXTER HLTHCARE 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
 BEDFORD EQ 150MG BASE/ML A063163 001 Jun 30, 1994
 BRISTOL MYERS SQUIBB EQ 150MG BASE/ML A062908 001 Feb 01, 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
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%
 ABRAXIS PHARM EQ 12MG BASE/ML N050636 001 Dec 22, 1989
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER
 ABBOTT 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
 ACTAVIS MID ATLANTIC EQ 1% BASE A062811 001 Sep 01, 1988
 BOCA PHARMA EQ 1% BASE A062930 001 Jun 28, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 86 (of 346)

CLINDAMYCIN PHOSPHATE

SOLUTION; TOPICAL CLINDAMYCIN PHOSPHATE	COPLEY PHARM COREPHARMA	EQ 1% BASE EQ 1% BASE	A062944 001	Jan 11, 1989
			A064108 001	Sep 27, 1996

CLIOQUINOL; NYSTATIN

OINTMENT; TOPICAL NYSTAFORM	BAYER PHARMS	10MG/GM;100,000 UNITS/GM	N050235 001
--------------------------------	--------------	--------------------------	-------------

CLOBETASOL PROPIONATE

CREAM; TOPICAL CLOBETASOL PROPIONATE	ACTAVIS MID ATLANTIC COREPHARMA	0.05% 0.05%	A074139 001	Aug 03, 1994
CLOBETASOL PROPIONATE (EMOLlient)	COREPHARMA	0.05%	A075733 001	Aug 22, 2001
OINTMENT; TOPICAL CLOBETASOL PROPIONATE	ACTAVIS MID ATLANTIC COREPHARMA	0.05% 0.05%	A074128 001 A075057 001	Aug 03, 1994 Aug 12, 1998
SOLUTION; TOPICAL CLOBETASOL PROPIONATE	ACTAVIS MID ATLANTIC	0.05%	A074331 001	Dec 15, 1995

CLOFAZIMINE

CAPSULE; ORAL LAMPRENE	NOVARTIS	100MG	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
---------------------------	-------	------	-------------	--------------

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL CLOMIPRAMINE HYDROCHLORIDE	SANDOZ	25MG	A074953 001	Jun 25, 1997
		50MG	A074953 002	Jun 25, 1997
		75MG	A074953 003	Jun 25, 1997
	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

DISCONTINUED DRUG PRODUCT LIST

6 - 87 (of 346)

CLONAZEPAM

TABLET; ORAL CLONAZEPAM SANDOZ	0.5MG 1MG 2MG	A074925 001 A074925 002 A074925 003	Sep 30, 1997 Sep 30, 1997 Sep 30, 1997
TEVA	0.5MG 1MG 2MG	A074920 001 A074920 002 A074920 003	Aug 04, 1998 Aug 04, 1998 Aug 04, 1998
KLONOPIN ROCHE	0.125MG 0.25MG	N017533 005 N017533 006	Apr 09, 1997 Apr 09, 1997
TABLET, ORALLY DISINTEGRATING; ORAL KLONOPIN RAPIDLY DISINTEGRATING ROCHE	0.125MG 0.25MG 0.5MG 1MG 2MG	N020813 001 N020813 002 N020813 003 N020813 004 N020813 005	Dec 23, 1997 Dec 23, 1997 Dec 23, 1997 Dec 23, 1997 Dec 23, 1997

CLONIDINE

SUSPENSION, 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 EQ 0.26MG BASE	N022500 001 N022500 002	Dec 03, 2009 Dec 03, 2009

CLONIDINE HYDROCHLORIDE

TABLET; ORAL CLONIDINE HYDROCHLORIDE AM THERAP	0.1MG 0.2MG 0.3MG	A070881 001 A070882 001 A070883 001	Jul 08, 1986 Jul 08, 1986 Jul 08, 1986
DURAMED PHARMS BARR	0.1MG 0.2MG 0.3MG	A071103 001 A071102 001 A071101 001	Aug 14, 1986 Aug 14, 1986 Aug 14, 1986
INTERPHARM	0.1MG 0.2MG 0.3MG	A071252 001 A071253 001 A071254 001	Oct 01, 1986 Oct 01, 1986 Oct 01, 1986
PAR PHARM	0.1MG 0.2MG 0.3MG	A070461 001 A070460 001 A070459 001	Jul 08, 1986 Jul 08, 1986 Jul 08, 1986
SANDOZ	0.1MG 0.2MG 0.3MG	A070887 001 A070886 001 A071294 001	Aug 31, 1988 Aug 31, 1988 Aug 31, 1988
TEVA	0.1MG 0.2MG 0.3MG	A070747 001 A070702 001 A070659 001	Jul 08, 1986 Jul 08, 1986 Jul 08, 1986
WARNER CHILCOTT	0.1MG 0.2MG 0.3MG	A072138 001 A072139 001 A072140 001	Jun 13, 1988 Jun 13, 1988 Jun 13, 1988
WATSON LABS	0.1MG 0.1MG 0.2MG 0.2MG 0.3MG	A070395 001 A070965 001 A070396 001 A070964 001 A070397 001	Mar 23, 1987 Jul 08, 1986 Mar 23, 1987 Jul 08, 1986 Mar 23, 1987
TABLET, EXTENDED RELEASE; ORAL JENLOGA SHIONOGI INC	0.1MG 0.2MG	N022331 001 N022331 002	Sep 30, 2009 May 25, 2010
KAPVAY SHIONOGI INC	0.2MG	N022331 004	Sep 28, 2010

DISCONTINUED DRUG PRODUCT LIST

6 - 88 (of 346)

CLOPIDOGREL BISULFATE

TABLET; ORAL
 CLOPIDOGREL BISULFATE
 DR REDDYS LABS INC EQ 75MG BASE A076273 001 Jan 14, 2008

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
CLONMEL HLTHCARE	3.75MG	A071742	001	Dec 14, 1987
	7.5MG	A071743	001	Dec 14, 1987
	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
SANDOZ	3.75MG	A072219	001	Aug 26, 1988
	7.5MG	A072220	001	Aug 26, 1988
	15MG	A072112	001	Aug 26, 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				
LUNDBECK INC	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
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
SANDOZ	3.75MG	A072512	001	May 11, 1990

DISCONTINUED DRUG PRODUCT LIST

6 - 89 (of 346)

CLORAZEPATE DIPOTASSIUM

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

SANDOZ	7.5MG	A072513	001	May 11, 1990
	15MG	A072514	001	May 11, 1990
WARNER CHILCOTT	3.75MG	A071828	001	Mar 03, 1988
	7.5MG	A071829	001	Mar 03, 1988
	15MG	A071830	001	Mar 03, 1988
TRANXENE SD				
LUNDBECK INC	11.25MG	N017105	005	
	22.5MG	N017105	004	

CLOTTRIMAZOLE

CREAM; TOPICAL

LOTTRIMIN

SCHERING PLOUGH	1%	N017619	001	
MYCELEX				
BAYER PHARMS	1%	N018183	001	
LOTION; TOPICAL				
LOTTRIMIN				
SCHERING	1%	N018813	001	Feb 17, 1984
SOLUTION; TOPICAL				
LOTTRIMIN				
SCHERING PLOUGH	1%	N017613	001	
TABLET; VAGINAL				
GYNIX				
TEVA PHARMS	100MG	A073249	001	Feb 13, 1998
MYCELEX-G				
BAYER PHARMS	500MG	N019069	001	Apr 19, 1985

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				
AZUR PHARMA INTL	50MG	N021590	003	Jun 03, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 90 (of 346)

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 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

COLCHICINE; PROBENECID

TABLET; ORAL COLBENEMID MERCK	0.5MG;500MG	N012383 001	
PROBEN-C WATSON LABS	0.5MG;500MG	A085552 001	
PROBENECID AND COLCHICINE BEECHAM	0.5MG;500MG	A084321 001	
IMPAX LABS	0.5MG;500MG	A083720 002	
IVAX SUB TEVA PHARMS	0.5MG;500MG	A083734 001	
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 - 91 (of 346)

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; IV (INFUSION)

VAPRISOL

ASTELLAS

20MG/4ML (5MG/ML)

N021697 001 Dec 29, 2005

COPPERINTRAUTERINE DEVICE; INTRAUTERINE
CU-7

GD SEARLE LLC 89MG

N017408 001

TATUM-T

GD SEARLE LLC 120MG

N018205 001

CORTICOTROPININJECTABLE; INJECTION
ACTHPARKEDALE 25 UNITS/VIAL
40 UNITS/VIAL

N008317 002

N008317 004

ACTHAR

SANOFI AVENTIS US 25 UNITS/VIAL
40 UNITS/VIAL

N007504 002

N007504 003

CORTICOTROPIN

ORGANICS LAGRANGE 40 UNITS/ML
80 UNITS/ML

N010831 001

N010831 002

WATSON LABS 40 UNITS/VIAL

A088772 001 Nov 21, 1984

H.P. ACTHAR GEL

QUESTCOR PHARMS 40 UNITS/ML

N008372 006

PURIFIED CORTROPHIN GEL

ORGANON USA INC 40 UNITS/ML
80 UNITS/ML

N008975 001

N008975 002

CORTICOTROPIN-ZINC HYDROXIDEINJECTABLE; INJECTION
CORTROPHIN-ZINC

ORGANON USA INC 40 UNITS/ML

N009854 001

CORTISONE ACETATEINJECTABLE; INJECTION
CORTISONE ACETATEPHARMACIA AND UPJOHN 25MG/ML
WATSON LABS 25MG/ML
25MG/ML
50MG/ML
50MG/ML

N008126 002

A083147 003

A085677 001

A083147 004

A085677 002

CORTONE

MERCK 25MG/ML
50MG/ML

N007110 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

A083471 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

DISCONTINUED DRUG PRODUCT LIST

6 - 92 (of 346)

CORTISONE ACETATE

TABLET; ORAL				
CORTISONE ACETATE				
VITARINE	25MG		A080333	001
WATSON LABS	25MG		A085884	001
WHITEWORTH TOWN PLSN	25MG		A080341	001
CORTONE				
MERCK	25MG		N007750	003

CROMOLYN SODIUM

CAPSULE; INHALATION				
INTAL				
SANOFI AVENTIS US	20MG		N016990	001
CAPSULE; ORAL				
GASTROCROM				
UCB INC	100MG		N019188	001
CONCENTRATE; ORAL				Dec 22, 1989
CROMOLYN SODIUM				
GENERA PHARMS	100MG/5ML		A090954	001
SOLUTION; INHALATION				Dec 18, 2009
CROMOLYN SODIUM				
ACTAVIS MID ATLANTIC	10MG/ML		A075067	001
PHARMASCIENCE INC	10MG/ML		A075437	001
ROXANE	10MG/ML		A075175	001
INTAL				Sep 30, 1999
KING PHARMS	10MG/ML		N018596	001
SOLUTION/DROPS; OPHTHALMIC				May 28, 1982
CROMOPTIC				
KING PHARMS	4%		A075088	001
SPRAY, METERED; NASAL				Apr 27, 1999
CROMOLYN SODIUM				
ACTAVIS MID ATLANTIC	5.2MG/SPRAY		A074800	001
NASALCROM				Jul 26, 2001
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
INJECTABLE; INJECTION				Nov 05, 1996
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 - 93 (of 346)

CYANOCOBALAMININJECTABLE; INJECTION
CYANOCOBALAMIN

ABRAXIS PHARM	0.03MG/ML	A080510	003	
	0.1MG/ML	A080510	001	
	1MG/ML	A080510	002	
AKORN	1MG/ML	A087969	001	Nov 10, 1983
APP PHARMS	0.1MG/ML	A080557	002	
BAXTER HLTHCARE	1MG/ML	A080515	002	
BIONICHE PHARMA	1MG/ML	A040451	001	Sep 23, 2003
DELL LABS	0.03MG/ML	A080689	001	
	0.1MG/ML	A080689	002	
	1MG/ML	A080689	003	
LUITPOLD	0.03MG/ML	A080668	001	
LYPHOMED	1MG/ML	A083075	001	
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	A080573	002	
	0.1MG/ML	A083120	001	
	1MG/ML	A080573	001	
	1MG/ML	A083120	002	
WYETH AYERST	0.1MG/ML	A080554	001	
	1MG/ML	A080554	002	
REDISOL				
MERCK	1MG/ML	N006668	010	
RUBIVITE				
BEL MAR	0.03MG/ML	N010791	004	
	0.05MG/ML	N010791	001	
	0.1MG/ML	N010791	002	
	0.12MG/ML	N010791	005	
	1MG/ML	N010791	003	
RUBRAMIN PC				
BRISTOL MYERS SQUIBB	0.1MG/ML	N006799	002	
	1MG/ML	N006799	004	
	1MG/ML	N006799	010	Apr 28, 1988
RUVITE				
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

DISCONTINUED DRUG PRODUCT LIST

6 - 94 (of 346)

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A; N/A
 CYANOCOBALAMIN CO 57 SCHILLING TEST KIT
 MALLINCKRODT 0.1MG;0.5uCi;60MG N016635 001

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

TABLET; ORAL
 CYCLOBENZAPRINE HYDROCHLORIDE
 SANDOZ 10MG A073683 001 Feb 26, 1993
 WATSON LABS 10MG A073143 001 Nov 27, 1991
 10MG A074436 001 Nov 30, 1994

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
 AK-PENTOLATE
 AKORN 1% A085555 001
 CYCLOPENTOLATE HYDROCHLORIDE
 ALCON UNIVERSAL 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
 LYOPHILIZED CYTOXAN
 BAXTER HLTHCARE 100MG/VIAL N012142 006 Dec 05, 1985
 200MG/VIAL N012142 007 Dec 10, 1985
 500MG/VIAL N012142 008 Jan 04, 1984
 1GM/VIAL N012142 010 Sep 24, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 95 (of 346)

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION LYOPHILIZED CYTOXAN BAXTER HLTHCARE NEOSAR TEVA PARENTERAL	2GM/VIAL 100MG/VIAL 100MG/VIAL 200MG/VIAL 200MG/VIAL 500MG/VIAL 500MG/VIAL 1GM/VIAL 1GM/VIAL 2GM/VIAL 2GM/VIAL	N012142 009	Dec 10, 1985
	A040015 001	Apr 29, 1993	
	A087442 001	Feb 16, 1982	
	A040015 002	Apr 29, 1993	
	A087442 002	Feb 16, 1982	
	A040015 003	Apr 29, 1993	
	A087442 003	Feb 16, 1982	
	A040015 004	Apr 29, 1993	
	A087442 004	Jul 08, 1983	
	A040015 005	Apr 29, 1993	
	A087442 005	Mar 30, 1989	
TABLET; ORAL CYTOXAN BAXTER HLTHCARE	25MG 50MG	N012141 002 N012141 001	

CYCLOSPORINE

CAPSULE; ORAL NEORAL NOVARTIS	50MG	N050715 003	Jul 14, 1995
-------------------------------------	------	-------------	--------------

CYCLOTHIAZIDE

TABLET; ORAL ANHYDRON LILLY FLUIDIL PHARMACIA AND UPJOHN	2MG 2MG 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 HALSEY MORTON GROVE NASKA PERIACTIN MERCK	2MG/5ML 2MG/5ML 2MG/5ML 2MG/5ML 2MG/5ML	A086833 001 A089199 001 A087001 001 A089021 001 N013220 002	Jul 03, 1986 Nov 04, 1982 Dec 21, 1987
TABLET; ORAL CYPROHEPTADINE HYDROCHLORIDE AM THERAP ASCOT DURAMED PHARMS BARR HALSEY KV PHARM MD PHARM MYLAN PIONEER PHARMS PLIVA SANDOZ SUPERPHARM TG UNITED LABS VITARINE WATSON LABS	4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG 4MG	A088798 001 A087685 001 A088232 001 A089057 001 A086737 001 A087566 001 A086678 001 A087839 001 A088205 001 A086808 001 A087405 001 A088212 001 A087284 001 A085245 001	Feb 15, 1985 Oct 25, 1982 Oct 25, 1983 Jul 03, 1986 Nov 10, 1982 Feb 08, 1984 Jul 26, 1983 May 26, 1983

DISCONTINUED DRUG PRODUCT LIST

6 - 96 (of 346)

CYPROHEPTADINE HYDROCHLORIDE

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

WATSON LABS	4MG	A086165	001
	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

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

ABRAXIS PHARM	100MG/VIAL	A070962	001	Aug 28, 1986
	200MG/VIAL	A070990	001	Aug 28, 1986

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; IV (INFUSION)

SYNERCID

KING PHARMS	420MG/VIAL;180MG/VIAL	N050748	002	Aug 24, 2000
-------------	-----------------------	---------	-----	--------------

DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

EISAI INC	7,500 IU/0.75ML	N020287	008	Apr 04, 2002
-----------	-----------------	---------	-----	--------------

DANAPAROID SODIUM

INJECTABLE; INJECTION

ORGARAN

ORGANON USA 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

200MG

N017557 004

N017557 002

DANTROLENE SODIUM

CAPSULE; ORAL

DANTROLENE SODIUM

ACTAVIS TOTOWA	25MG	A076686	001	Oct 24, 2005
	50MG	A076686	002	Oct 24, 2005
	100MG	A076686	003	Oct 24, 2005

DAPIPRAZOLE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

REV-EYES

FERA PHARMS	0.5%	N019849	001	Dec 31, 1990
-------------	------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 97 (of 346)

DAPTOMYCIN

INJECTABLE; IV (INFUSION)
 CUBICIN
 CUBIST 250MG/VIAL N021572 001 Sep 12, 2003

DARUNAVIR ETHANOLATE

TABLET; ORAL
 PREZISTA
 TIBOTEC EQ 300MG BASE N021976 001 Jun 23, 2006

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

DEMECARIUM BROMIDE

SOLUTION/DROPS; OPHTHALMIC
 HUMORSOL
 MERCK 0.125% N011860 002
 0.25% N011860 001

DEMECLOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL
 DECLEOMYCIN
 LEDERLE 150MG N050262 001
 SYRUP; ORAL
 DECLEOMYCIN
 LEDERLE 75MG/5ML N050257 001
 TABLET; ORAL
 DECLEOMYCIN
 COREPHARMA 75MG N050261 001

DESERPIDIENE

TABLET; ORAL
 HARMONYL
 ABBOTT 0.1MG N010796 001
 0.25MG N010796 002

DESERPIDIENE; HYDROCHLORTIAZIDE

TABLET; ORAL
 ORETICYL 25
 ABBOTT 0.125MG;25MG N012148 001
 ORETICYL 50
 ABBOTT 0.125MG;50MG N012148 003
 ORETICYL FORTE
 ABBOTT 0.25MG;25MG N012148 002

DESERPIDIENE; METHYCLOTHIAZIDE

TABLET; ORAL
 ENDURONYL
 ABBOTT 0.25MG;5MG N012775 001
 ENDURONYL FORTE
 ABBOTT 0.5MG;5MG N012775 002

DISCONTINUED DRUG PRODUCT LIST

6 - 98 (of 346)

DESERPIDINE; METHYCLOTHIAZIDE

TABLET; ORAL

METHYCLOTHIAZIDE AND DESERPIDINE

WATSON LABS	0.25MG;5MG 0.5MG;5MG	A088486 001	Aug 10, 1984
		A088452 001	Aug 10, 1984

DESIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

PERTOFRANE

SANOFI AVENTIS US	25MG 50MG	N013621 001
		N013621 002

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

PLIVA	25MG 50MG 75MG 100MG 150MG	A071800 001 A071801 001 A071802 001 A071803 001 A071804 001	Dec 08, 1987 Dec 08, 1987 Dec 08, 1987 May 29, 1997 May 29, 1997
USL PHARMA	25MG 50MG 75MG 100MG	A071864 001 A071865 001 A071866 001 A071867 001	Sep 09, 1987 Sep 09, 1987 Sep 09, 1987 Sep 09, 1987

DESLANOSIDE

INJECTABLE; INJECTION

CEDILANID-D

NOVARTIS	0.2MG/ML	N009282 002
----------	----------	-------------

DESMOPRESSIN ACETATEINJECTABLE; INJECTION
DDAVP

SANOFI AVENTIS US	0.015MG/ML	N018938 002	Apr 25, 1995
DESMOPRESSIN ACETATE			
BEDFORD	0.004MG/ML	A074575 001	Feb 18, 2000
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			
SANOFI AVENTIS US	0.01MG/SPRAY	N017922 002	Feb 06, 1989
STIMATE			
CSL BEHRING	0.15MG/SPRAY	N020355 001	Mar 07, 1994

DESOGESTREL; ETHINYLL ESTRADIOL

TABLET; ORAL-21

DESOGEN

ORGANON USA INC	0.15MG;0.03MG	N020071 001	Dec 10, 1992
DESOGESTREL AND ETHINYLL ESTRADIOL			
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

DESONIDE

CREAM; TOPICAL

DESONIDE

TEVA PHARMS	0.05%	A074027 001	Sep 28, 1992
-------------	-------	-------------	--------------

DESOXIMETASONE

OINTMENT; TOPICAL

DESOXIMETASONE

ALTANA	0.25%	A073440 001	Apr 01, 1998
--------	-------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 99 (of 346)

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

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		
IMPAK LABS	0.75MG	A085376 001
MUTUAL PHARM	0.25MG	A084013 001
	0.25MG	A084764 001
	0.5MG	A084084 001
	0.5MG	A084766 001
	0.75MG	A084081 001
	0.75MG	A084765 001
	1.5MG	A084086 001
	1.5MG	A084763 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
SANDOZ	0.75MG	A080399 001
UPSHER SMITH	0.75MG	A087534 001
	1.5MG	A087533 001
WATSON LABS	0.25MG	A085455 001
	0.5MG	A085458 001
	0.75MG	A080968 001
	0.75MG	A084457 001
	0.75MG	A085818 001
	1.5MG	A085456 001
	1.5MG	A085840 001

DISCONTINUED DRUG PRODUCT LIST

6 - 100 (of 346)

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

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			
ORGANON USA INC	0.5MG	N012675	004
	0.75MG	N012675	007
	1.5MG	N012675	009
	4MG	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

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

APP PHARMS

EQ 10MG PHOSPHATE/ML

A088469 001 Jan 25, 1984

DEXAMETHASONE SODIUM PHOSPHATE

AKORN EQ 4MG PHOSPHATE/ML

A084493 001

BAXTER HLTHCARE EQ 4MG PHOSPHATE/ML

A084282 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

WATSON LABS EQ 4MG 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

WYETH AYERST

HEXDROL

ORGANON USA INC EQ 4MG PHOSPHATE/ML

N014694 002

EQ 10MG PHOSPHATE/ML

N014694 003

EQ 20MG PHOSPHATE/ML

N014694 004

DISCONTINUED DRUG PRODUCT LIST

6 - 101 (of 346)

DEXAMETHASONE SODIUM PHOSPHATE

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 Oct 30, 1995
NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE				
ALCON UNIVERSAL	EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML		A062714	001 Jul 21, 1986
PHARMAFAIR	EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML		A062539	001 Jan 10, 1985

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC				
DEXACIDIN				
NOVARTIS	0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM		A062566	001 Feb 22, 1985
DEXASPORIN				
PHARMAFAIR	0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM		A062411	001 May 16, 1983
SUSPENSION/DROPS; OPHTHALMIC				
DEXACIDIN				
NOVARTIS	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML		A062544	001 Oct 29, 1984
DEXASPORIN				
PHARMAFAIR	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML		A062428	001 May 18, 1983

DEXBROMPHENIRAMINE MALEATE

SYRUP; ORAL				
DISOMER				
SCHERING	2MG/5ML		N011814	002
TABLET; ORAL				
DISOPHROL				
SCHERING	2MG		N011814	001

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHENDRINE SULFATE

TABLET; ORAL				
DISOPHROL				
SCHERING	2MG;60MG		N012394	002

DISCONTINUED DRUG PRODUCT LIST

6 - 102 (of 346)

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
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				
PLIVA	2MG	A088682	001	Jan 17, 1986
POLARAMINE				
SCHERING	2MG	A086835	001	

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				
ENDO PHARMS	5MG	A040299	001	May 13, 1999
HALSEY	10MG	A083930	001	
LANNETT	5MG	A083903	001	
	10MG	A083903	003	
	15MG	A085652	001	
MAST MM	5MG	A086521	001	
PUREPAC PHARM	5MG	A084125	001	
SANDOZ	5MG	A085370	001	
	10MG	A085371	001	
VITARINE	5MG	A084986	001	
	10MG	A085892	001	
DEXTROSTAT				
SHIRE	5MG	A084051	001	
	10MG	A084051	002	
FERNDEX				
FERNDALE LABS	5MG	A084001	001	

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL PHERAZINE DM				
HALSEY	15MG/5ML;6.25MG/5ML	A088913	001	Mar 02, 1987
PROMETHAZINE HYDROCHLORIDE AND DESTROMETHORPHAN HYDROBROMIDE				
ANI PHARMS	15MG/5ML;6.25MG/5ML	N011265	002	Apr 02, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 103 (of 346)

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 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; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION					
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 W/ DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	5GM/100ML;30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML	N018274	001		

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	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 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	5GM/100ML;220MG/100ML	N019699	005	Sep 29, 1989	

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 W/ DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG/100ML;91MG/100ML	N018270	001		

DISCONTINUED DRUG PRODUCT LIST

6 - 104 (of 346)

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 15MEQ IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5GM/100ML;224MG/100ML;450MG/100ML	N018008	003	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5GM/100ML;300MG/100ML;450MG/100ML	N018008	001	
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5GM/100ML;75MG/100ML;450MG/100ML	N018008	002	

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	
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 105 (of 346)

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
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

ABBOTT	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	85%	N011620	002

BRACCO

DIATRIZOATE MEGLUMINE

BRACCO

HYPAQUE

GE HEALTHCARE

RENO-60

RENO-DIP

UROVIST MEGLUMINE DIU/CT

BAYER HLTHCARE

SOLUTION; URETERAL

RENO-30

BRACCO

UROVIST CYSTO

BAYER HLTHCARE

UROVIST CYSTO PEDIATRIC

BAYER HLTHCARE

SOLUTION; URETHRAL

HYPAQUE-CYSTO

GE HEALTHCARE

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

ANGIOVIST 292

BAYER HLTHCARE

ANGIOVIST 370

BAYER HLTHCARE

DIATRIZOATE-60

INT'L MEDICATION

HYPAQUE-76

GE HEALTHCARE

DISCONTINUED DRUG PRODUCT LIST

6 - 106 (of 346)

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION				
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	
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	5MG/ML (5MG/ML)	N020648	002	Jul 29, 1997
	10MG/2ML (5MG/ML)	N020648	003	Jul 29, 1997
	15MG/3ML (5MG/ML)	N020648	004	Jul 29, 1997
	20MG/4ML (5MG/ML)	N020648	005	Jul 29, 1997
INJECTABLE; INJECTION				
DIAZEPAM				
ABRAXIS PHARM	5MG/ML	A070662	001	Jun 25, 1986
BAXTER HLTHCARE	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
MARSAM PHARMS LLC	5MG/ML	A072370	001	Jan 29, 1993
	5MG/ML	A072371	001	Jan 29, 1993
	5MG/ML	A072397	001	Jan 29, 1993
PARENTA PHARMS	5MG/ML	A076815	001	Apr 15, 2004

DISCONTINUED DRUG PRODUCT LIST

6 - 107 (of 346)

DIAZEPAM

INJECTABLE; INJECTION

DIAZEPAM

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	A070911	001	Aug 28, 1986
	5MG/ML	A070912	001	Aug 28, 1986
	5MG/ML	A070930	001	Dec 01, 1986

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	5MG	A070227	001	Sep 26, 1985
	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
PAR PHARM	2MG	A070462	001	Feb 25, 1986
	5MG	A070463	001	Feb 25, 1986
	10MG	A070464	001	Feb 25, 1986
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
SANDOZ	2MG	A070302	001	Dec 20, 1985
	5MG	A070303	001	Dec 20, 1985
	10MG	A070304	001	Dec 20, 1985
TEVA PHARMS	5MG	A070153	001	Nov 01, 1985
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
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

DISCONTINUED DRUG PRODUCT LIST

6 - 108 (of 346)

DIAZOXIDE

CAPSULE; ORAL PROGLYCEM				
TEVA BRANDED PHARM	50MG 100MG		N017425 001 N017425 002	
INJECTABLE; INJECTION DIAZOXIDE				
ABRAXIS PHARM	15MG/ML		A071519 001	Aug 26, 1987
HYPERSTAT SCHERING	15MG/ML		N016996 001	

DIBUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION HEAVY SOLUTION NUPERCAINE				
NOVARTIS	2.5MG/ML		N006203 001	

DICHLORPHENAMIDE

TABLET; ORAL DARANIDE				
TARO	50MG		N011366 001	

DICLOFENAC POTASSIUM

TABLET; ORAL CATAFLAM				
NOVARTIS	25MG		N020142 001	Nov 24, 1993
DICLOFENAC POTASSIUM				
MUTUAL PHARM	50MG		A075470 001	Feb 21, 2002
SANDOZ	50MG		A075582 001	Feb 23, 2001
WATSON LABS	50MG		A075152 001	Nov 27, 1998

DICLOFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC DICLOFENAC SODIUM				
FALCON PHARMS	0.1%		N020809 001	May 04, 1998
TABLET, DELAYED RELEASE; ORAL DICLOFENAC SODIUM				
NOSTRUM LABS	50MG		A074986 001	Feb 26, 1999
	75MG		A074986 002	Feb 26, 1999
PLIVA	50MG		A074432 002	Jul 29, 1999
	75MG		A074432 003	Jul 29, 1999
ROXANE	25MG		A074391 001	Jun 29, 1995
	50MG		A074391 002	Jun 29, 1995
	75MG		A074391 003	Jun 29, 1995
TEVA	50MG		A074723 001	Mar 30, 1999
	75MG		A074390 001	Aug 15, 1996
TEVA PHARMS	25MG		A074459 001	Jun 25, 1997
	50MG		A074459 002	Jun 25, 1997
	75MG		A074459 003	Jun 25, 1997
VOLTAREN				
NOVARTIS	25MG		N019201 001	Jul 28, 1988
	50MG		N019201 002	Jul 28, 1988
	75MG		N019201 003	Jul 28, 1988

DICLOXA CILLIN 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

DISCONTINUED DRUG PRODUCT LIST

6 - 109 (of 346)

DICLOxacillin Sodium

FOR SUSPENSION; ORAL DICLOACILLIN 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			
ABBOTT	25MG	N005545	003
	50MG	N005545	004
	100MG	N005545	005

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL DICYCLOMINE HYDROCHLORIDE			
MUTUAL PHARM	10MG	A084505	001 Oct 21, 1986
PIONEER PHARMS	10MG	A089361	001 Jan 10, 1989
WATSON LABS	10MG	A083179	001 Feb 12, 1986
INJECTABLE; INJECTION DICYCLOMINE HYDROCHLORIDE			
WATSON LABS	10MG/ML	A080614	001 Feb 11, 1986
SYRUP; ORAL DICYCLOMINE HYDROCHLORIDE			
ALPHARMA US PHARMS	10MG/5ML	A084479	001
TABLET; ORAL DICYCLOMINE HYDROCHLORIDE			
MUTUAL PHARM	20MG	A084600	001 Jul 29, 1985
PIONEER PHARMS	20MG	A088585	001 Aug 20, 1986
WATSON LABS	20MG	A084361	001 Feb 06, 1986

DIDANOSINE

FOR SOLUTION; ORAL 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
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

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

DISCONTINUED DRUG PRODUCT LIST

6 - 110 (of 346)

DIETHYLCARBAMAZINE CITRATE

TABLET; ORAL HETRAZAN LEDERLE	50MG	N006459 001
-------------------------------------	------	-------------

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL DIETHYLPROPION HYDROCHLORIDE		
SANDOZ	25MG	A085916 001
TEVA	25MG	A088642 001
TG UNITED LABS	25MG	A088267 001
	25MG	A088268 001
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

STILBESTROL

TABLICAPS	0 .5MG	A083004 001
	1MG	A083002 001
	5MG	A083006 001

STILBETIN

BRISTOL MYERS SQUIBB	0 .1MG	N004056 007
	0 .25MG	N004056 017
	0 .5MG	N004056 008
	1MG	N004056 009
	5MG	N004056 010

TABLET, DELAYED RELEASE; ORAL
DIETHYLSTILBESTROL

LILLY	0 .1MG	N004039 002
	0 .25MG	N004039 005
	0 .5MG	N004039 003
	1MG	N004039 004
	5MG	N004039 006

STILBESTROL

TABLICAPS	0 .5MG	A083003 001
	1MG	A083005 001

DISCONTINUED DRUG PRODUCT LIST

6 - 111 (of 346)

DIETHYLSILBESTROL

TABLET, DELAYED RELEASE; ORAL STILBESTROL TABLICAPS	5MG	A083007	001
STILBETIN BRISTOL MYERS SQUIBB	0.1MG	N004056	011
	0.5MG	N004056	012
	1MG	N004056	013
	5MG	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 FLORONE PHARMACIA AND UPJOHN	0.05%	N017741	001
FLORONE E PHARMACIA AND UPJOHN	0.05%	N019259	001 Aug 28, 1985
PSORCON SANOFI AVENTIS US	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 PUREPAC PHARM	250MG	A074285	001 May 07, 1996
	500MG	A074285	002 May 07, 1996
ROXANE	250MG	A073562	001 Nov 27, 1992
	500MG	A073563	001 Nov 27, 1992
SANDOZ	500MG	A074604	001 Jun 10, 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

DIGITOXIN

INJECTABLE; INJECTION CRYSTODIGIN LILLY	0.2MG/ML	A084100	005
---	----------	---------	-----

DIGOXIN

CAPSULE; ORAL LANOXICAPS GLAXOSMITHKLINE LLC	0.05MG	N018118	002 Jul 26, 1982
	0.1MG	N018118	003 Jul 26, 1982
	0.15MG	N018118	004 Sep 24, 1984
	0.2MG	N018118	001 Jul 26, 1982
INJECTABLE; INJECTION DIGOXIN ABRAXIS PHARM	0.25MG/ML	A083217	001
HOSPIRA	0.25MG/ML	A040206	001 Aug 28, 1998

DISCONTINUED DRUG PRODUCT LIST

6 - 112 (of 346)

DIGOXIN

INJECTABLE; INJECTION DIGOXIN					
WYETH AYERST	0.25MG/ML		A084386	001	
DIGOXIN PEDIATRIC HOSPIRA	0.1MG/ML		A040092	001	Apr 25, 1996
TABLET; ORAL DIGOXIN					
ACTAVIS TOTOWA	0.125MG		A040282	001	Dec 23, 1999
	0.25MG		A040282	002	Dec 23, 1999
LANOXIN					
COVIS PHARMA	0.0625MG		N020405	001	Sep 30, 1997
	0.1875MG		N020405	003	Sep 30, 1997
	0.375MG		N020405	005	Sep 30, 1997
	0.5MG		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		N018885	001	Nov 30, 1984
	0.5MG/0.7ML; 5,000 UNITS/0.7ML; 7.46MG/0.7ML		N018885	002	Nov 30, 1984

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL CARDIZEM SR					
BIOVAIL	60MG		N019471	001	Jan 23, 1989
	90MG		N019471	002	Jan 23, 1989
	120MG		N019471	003	Jan 23, 1989
	180MG		N019471	004	Jan 23, 1989
DILTIAZEM HYDROCHLORIDE					
BIOVAIL	60MG		A074845	001	Sep 15, 1999
	90MG		A074845	002	Sep 15, 1999
	120MG		A074845	003	Sep 15, 1999
	120MG		N020939	001	Jan 28, 2000
	180MG		N020939	002	Jan 28, 2000
	240MG		N020939	003	Jan 28, 2000
	300MG		N020939	004	Jan 28, 2000
	360MG		N020939	005	Sep 14, 2001
	420MG		N020939	006	Sep 14, 2001
TEVA	60MG		A074079	001	Nov 30, 1993
	90MG		A074079	002	Nov 30, 1993
	120MG		A074079	003	Nov 30, 1993
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
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
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

DISCONTINUED DRUG PRODUCT LIST

6 - 113 (of 346)

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

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 MALATETABLET, 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 MALEATETABLET, EXTENDED RELEASE; ORAL
TECZEM
BIOVAIL

BIOVAIL	EQ 180MG HCL;5MG	N020507	001	Oct 04, 1996
---------	------------------	---------	-----	--------------

DIMENHYDRINATEINJECTABLE; INJECTION
DIMENHYDRINATE

BAXTER HLTHCARE	50MG/ML	A084767	001
WATSON LABS	50MG/ML	A083531	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; PERFLEXANE

INJECTABLE; INTRAVENOUS IMAGENT IMCOR PHARMS CO	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
---	----------------	---------	-----

DIPHEMANIL METHYLSULFATE

TABLET; ORAL PRANTAL SCHERING	100MG	N008114	004
-------------------------------------	-------	---------	-----

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; 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
HEATHER	25MG	A084524	001
			Jun 04, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 114 (of 346)

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

HEATHER	50MG	A083953	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	
	25MG	A089488	001	Jan 02, 1987
	50MG	A089489	001	Jan 02, 1987
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	
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	
WEST WARD	50MG	A083567	001	
WHITEWORTH TOWN PLSN	25MG	A083441	001	
	50MG	A080800	001	
ELIXIR; ORAL				
BELIX				
HALSEY	12.5MG/5ML	A086586	001	Oct 03, 1983
BENADRYL				
MCNEIL CONS	12.5MG/5ML	N005845	004	
DIBENIL				
CENCI	12.5MG/5ML	A088304	001	Dec 16, 1983
DIPHEN				
USL PHARMA	12.5MG/5ML	A084640	001	
DIPHENHYDRAMINE HYDROCHLORIDE				
BUNDY	12.5MG/5ML	A083674	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 115 (of 346)

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

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	
DIPHENHYDRAMINE HYDROCHLORIDE				
BAXTER HLTHCARE	50MG/ML	A083183	001	
BEL MAR	10MG/ML	A080822	001	
LYPHOMED	10MG/ML	A087066	001	
WATSON LABS	10MG/ML	A083533	001	
WYETH AYERST	50MG/ML	A080577	001	
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE				
ABRAXIS PHARM	50MG/ML	A080586	002	
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

DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

BENYLIN

PARKE DAVIS	12.5MG/5ML; 30MG/5ML	N019014	001	Jun 11, 1985
-------------	----------------------	---------	-----	--------------

DIPHENIDOL HYDROCHLORIDE

TABLET; ORAL

VONTROL

GLAXOSMITHKLINE	EQ 25MG BASE	N016033	001
-----------------	--------------	---------	-----

DIPHENYL PYRALINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

HISPRIL

GLAXOSMITHKLINE	5MG	N011945	001
-----------------	-----	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 116 (of 346)

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC DIPIVEFRIN HYDROCHLORIDE BAUSCH AND LOMB	0.1%	A074188	001	May 19, 1995
---	------	---------	-----	--------------

DIPYRIDAMOLE

INJECTABLE; INJECTION DIPYRIDAMOLE HOSPIRA	5MG/ML	A074601	001	Dec 19, 1997
IV PERSANTINE BOEHRINGER INGELHEIM	5MG/ML	N019817	001	Dec 13, 1990
TABLET; ORAL DIPYRIDAMOLE PUREPAC PHARM	50MG	A089426	001	Jul 12, 1990
	75MG	A089427	001	Jul 12, 1990
SANDOZ	25MG	A086944	002	Apr 16, 1991
	50MG	A087562	001	Feb 25, 1992
	75MG	A087561	001	Feb 25, 1992

DIRITHROMYCIN

TABLET, DELAYED RELEASE; ORAL DYNABAC LILLY RES LABS	250MG	N050678	001	Jun 19, 1995
--	-------	---------	-----	--------------

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL DISOPYRAMIDE PHOSPHATE 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
MUTUAL PHARM	EQ 100MG BASE	A070351	001	Dec 17, 1985
	EQ 150MG BASE	A070352	001	Dec 17, 1985
MYLAN	EQ 100MG BASE	A070138	001	Jun 14, 1985
	EQ 150MG BASE	A070139	001	Jun 14, 1985
SANDOZ	EQ 100MG BASE	A070470	001	Dec 10, 1985
	EQ 150MG BASE	A070471	001	Dec 10, 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

DISULFIRAM

TABLET; ORAL ANTABUSE TEVA WOMENS	250MG	N007883	003	
	500MG	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	A086890	001	
	500MG	A087974	001	Aug 05, 1983

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE; ORAL DEPAKOTE CP ABBOTT	EQ 250MG BASE	N019794	001	Jul 11, 1990
	EQ 500MG BASE	N019794	002	Jul 11, 1990

DISCONTINUED DRUG PRODUCT LIST

6 - 117 (of 346)

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

ASTRAZENECA	EQ 12.5MG BASE/ML	A074098	001	Feb 21, 1995
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
MARSAM PHARMS LLC	EQ 12.5MG BASE/ML	A074279	001	Feb 18, 1998
TEVA PARENTERAL	EQ 12.5MG BASE/ML	A074995	001	Mar 31, 1998
DOBUTREX	EQ 12.5MG BASE/ML	A074206	001	Oct 19, 1993
LILLY	EQ 12.5MG BASE/ML	N017820	002	

DOCETAXEL

INJECTABLE; INJECTION

TAXOTERE

SANOFI AVENTIS US 40MG/ML

N020449 001 May 14, 1996

DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

EISAI INC 5MG/5ML
TABLET, ORALLY DISINTEGRATING; ORAL
DONEPEZIL HYDROCHLORIDE
MUTUAL PHARM 5MG
10MGN021719 001 Oct 18, 2004
A077975 002 Dec 11, 2009
A077975 001 Dec 11, 2009DOPAMINE 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
ASTRAZENECA	40MG/ML	A070087	001	Oct 23, 1985
	40MG/ML	N018656	001	Jun 28, 1983
	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
BAXTER HLTHCARE	40MG/ML	N018398	001	
	80MG/ML	N018398	002	Mar 22, 1982
HOSPIRA	40MG/ML	A074403	001	May 23, 1996
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
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 118 (of 346)

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION
INTROPIN
HOSPIRA 160MG/ML N017395 003

DOXACURIUM CHLORIDE

INJECTABLE; INJECTION
NUROMAX
ABBOTT EO 1MG BASE/ML N019946 001 Mar 07, 1991

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
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
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
SANDOZ	EQ 1MG BASE	A075432	001	Oct 18,	2000
	EQ 1MG BASE	A075646	001	Oct 18,	2000
	EQ 2MG BASE	A075432	002	Oct 18,	2000
	EQ 2MG BASE	A075646	002	Oct 18,	2000
	EQ 4MG BASE	A075432	003	Oct 18,	2000
	EQ 4MG BASE	A075646	003	Oct 18,	2000
	EQ 8MG BASE	A075432	004	Oct 18,	2000
	EQ 8MG BASE	A075646	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

DOXEPI N HYDROCHLORIDE

CAPSULE; ORAL					
DOXEPI N HYDROCHLORIDE					
CLONMEL HLTHCARE	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
MUTUAL PHARM	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
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		
	EQ 150MG BASE	N016987	007	Apr 13,	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

DISCONTINUED DRUG PRODUCT LIST

6 - 119 (of 346)

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

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
WATSON LABS	EQ 10MG BASE	A070952	001	Mar 04, 1987
	EQ 10MG BASE	A072985	001	Mar 29, 1991
	EQ 25MG BASE	A070953	001	May 15, 1986
	EQ 25MG BASE	A072986	001	Mar 29, 1991
	EQ 50MG BASE	A070954	001	May 15, 1986
	EQ 50MG BASE	A072987	001	Mar 29, 1991
	EQ 75MG BASE	A071763	001	Feb 09, 1988
	EQ 100MG BASE	A070955	001	May 15, 1986
	EQ 150MG BASE	A071764	001	Feb 09, 1988
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				
SINEQUAN				
PFIZER	EQ 10MG BASE/ML	N017516	001	

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

PHARMACIA AND UPJOHN	2MG/ML	A063165	001	Jan 30, 1991
	2MG/ML	N050629	001	Dec 23, 1987
	200MG/100ML	A063165	002	Jan 30, 1991
	200MG/100ML	N050629	002	May 03, 1988
ADRIAMYCIN RDF				
PHARMACIA AND UPJOHN	10MG/VIAL	N050467	001	
	20MG/VIAL	N050467	003	May 20, 1985
	50MG/VIAL	N050467	002	
	150MG/VIAL	N050467	004	Jul 22, 1987
RUBEX				
BRISTOL MYERS SQUIBB	10MG/VIAL	A062926	001	Apr 13, 1989
	50MG/VIAL	A062926	002	Apr 13, 1989
	100MG/VIAL	A062926	003	Apr 13, 1989

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

PAR PHARM	EQ 75MG BASE	A065055	004	Apr 18, 2005
FOR SUSPENSION; ORAL				
DOXYCHEL				
RACHELLE	EQ 25MG BASE/5ML	A061720	001	
TABLET; ORAL				
DOXYCYCLINE				
MUTUAL PHARM	EQ 50MG BASE	A065471	001	Apr 17, 2009
	EQ 75MG BASE	A065471	002	Apr 17, 2009
	EQ 100MG BASE	A065471	003	Apr 17, 2009

DISCONTINUED DRUG PRODUCT LIST

6 - 120 (of 346)

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

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
MYLAN	EQ 50MG BASE	A062337	001	Mar 29, 1982
	EQ 100MG BASE	A062337	002	Mar 29, 1982
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	
DOXY-LEMMON				
TEVA	EQ 50MG BASE	A062497	001	Aug 23, 1984
	EQ 100MG BASE	A062497	002	Jun 15, 1984
PERIOSTAT				
COLLAGENEX	EQ 20MG BASE	N050744	001	Sep 30, 1998
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
	EQ 100MG BASE	N050582	001	Jul 22, 1985
WARNER CHILCOTT	EQ 100MG BASE	A062653	001	Oct 30, 1985
INJECTABLE; INJECTION				
DOXYCHEL HYCLATE				
RACHELLE	EQ 100MG BASE/VIAL	A061953	001	
DOXYCYCLINE				
BAXTER HLTHCARE	EQ 100MG BASE/VIAL	A062450	001	Oct 27, 1983
	EQ 200MG BASE/VIAL	A062450	002	Oct 27, 1983
BEDFORD	EQ 200MG BASE/VIAL	A062569	002	Mar 09, 1988
DOXYCYCLINE HYCLATE				
BAXTER HLTHCARE	EQ 100MG BASE/VIAL	A062992	001	Feb 16, 1989
	EQ 200MG BASE/VIAL	A062992	002	Feb 16, 1989
VIBRAMYCIN				
PFIZER	EQ 100MG BASE/VIAL	N050442	002	
	EQ 200MG BASE/VIAL	N050442	001	
TABLET; ORAL				
DOXYCYCLINE HYCLATE				
HEATHER	EQ 100MG BASE	A062462	001	May 11, 1983
INTERPHARM	EQ 100MG BASE	A062764	001	Sep 02, 1988
LARKEN LABS	EQ 20MG BASE	A065287	001	Feb 28, 2006
MUTUAL PHARM	EQ 100MG BASE	A062391	001	Sep 30, 1982
MYLAN	EQ 100MG BASE	A062432	001	Feb 15, 1983
SUPERPHARM	EQ 100MG BASE	A062494	001	Feb 20, 1985
TRUXTON INC	EQ 50MG BASE	A062269	003	
	EQ 100MG BASE	A062269	002	Nov 08, 1982
WARNER CHILCOTT	EQ 100MG BASE	A062593	001	Aug 28, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 121 (of 346)

DOXYCYCLINE HYCLATE

TABLET; ORAL DOXYCYCLINE HYCLATE					
WATSON LABS	EQ 50MG BASE EQ 100MG BASE		A062392	001	Mar 31, 1983
DOXY-LEMMON			A062392	002	Mar 31, 1983
TEVA	EQ 100MG BASE		A062581	001	Mar 15, 1985
VIBRA-TABS					
PFIZER	EQ 100MG BASE		N050533	001	

DOXYLAMINE SUCCINATE

CAPSULE; ORAL UNISOM					
PFIZER	25MG		N019440	001	Feb 05, 1986
TABLET; ORAL DECAPRYN					
SANOFI AVENTIS US	12.5MG 25MG		N006412	015 014	
DOXYLAMINE SUCCINATE					
COPLEY PHARM	25MG		A088900	002	Feb 12, 1988
QUANTUM PHARMICS	25MG		A088603	001	Aug 07, 1984
DOXY-SLEEP-AID					
PAR PHARM	25MG		A070156	001	Jul 02, 1987

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL BENDECTIN					
SANOFI AVENTIS US	10MG;10MG		N010598	002	

DROMOSTANOLONE PROPIONATE

INJECTABLE; INJECTION DROLBAN					
LILLY	50MG/ML		N012936	001	

DROPERIDOL

INJECTABLE; INJECTION DROPERIDOL					
ABRAXIS PHARM	2.5MG/ML 2.5MG/ML		A070992	001	Nov 17, 1986
ASTRAZENECA	2.5MG/ML 2.5MG/ML 2.5MG/ML 2.5MG/ML		A072018	001	Oct 20, 1988
HOSPIRA	2.5MG/ML 2.5MG/ML		A072019	001	Oct 19, 1988
LUITPOLD	2.5MG/ML		A072020	001	Oct 19, 1988
SMITH AND NEPHEW	2.5MG/ML		A072021	001	Oct 19, 1988
SOLOPAK	2.5MG/ML 2.5MG/ML		A071645	001	Apr 07, 1988
WATSON LABS	2.5MG/ML 2.5MG/ML 2.5MG/ML		A072272	001	Aug 31, 1995
			A072335	001	Oct 24, 1988
			A071750	001	Sep 06, 1988
			A071754	001	Sep 06, 1988
			A071755	001	Sep 06, 1988
			A073520	001	Nov 27, 1991
			A073521	001	Nov 27, 1991
			A073523	001	Nov 27, 1991

DROPERIDOL; FENTANYL CITRATE

INJECTABLE; INJECTION FENTANYL CITRATE AND DROPERIDOL					
ASTRAZENECA	2.5MG/ML;EQ 0.05MG BASE/ML 2.5MG/ML;EQ 0.05MG BASE/ML 2.5MG/ML;EQ 0.05MG BASE/ML		A072026	001	Apr 13, 1989
HOSPIRA	2.5MG/ML;EQ 0.05MG BASE/ML		A072027	001	Apr 13, 1989
INNOVAR			A072028	001	Apr 13, 1989
AKORN MFG	2.5MG/ML;EQ 0.05MG BASE/ML		A071982	001	May 04, 1988
			N016049	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 122 (of 346)

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
NEOTHYLLINE			
TEVA	200MG	N007794	001
	400MG	N007794	002

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC PHOSPHOLINE IODIDE			
WYETH PHARMS INC	0.03%	N011963	002
	0.06%	N011963	004
	0.25%	N011963	003

ECONAZOLE NITRATE

CREAM; TOPICAL SPECTAZOLE			
ORTHO JANSSEN	1%	N018751	001 Dec 23, 1982

EDETADE CALCIUM DISODIUM

TABLET; ORAL CALCIUM DISODIUM VERSENATE			
MEDICIS	500MG	N008922	002

EDETADE DISODIUM

INJECTABLE; INJECTION DISODIUM EDETADE			
WATSON LABS	150MG/ML	A084356	001
EDETADE DISODIUM			
WATSON LABS	150MG/ML	A080391	001
SODIUM VERSENATE			
3M	200MG/ML	N010573	001

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

DISCONTINUED DRUG PRODUCT LIST

6 - 123 (of 346)

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
--	---------	-----	--------------

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
RANBAXY 2.5MG	A075556	001	Aug 22, 2000
5MG	A075556	002	Aug 22, 2000
10MG	A075556	003	Aug 22, 2000
20MG	A075556	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

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE; ORAL LEXCEL ASTRAZENECA 5MG; 2.5MG	N020668	002	Oct 28, 1998
5MG; 5MG	N020668	001	Dec 27, 1996

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE IVAX SUB TEVA PHARMS 5MG; 12.5MG	A075736	001	Mar 25, 2003
10MG; 25MG	A075736	002	Mar 25, 2003

ENALAPRILAT

INJECTABLE; INJECTION ENALAPRILAT HOSPIRA 1.25MG/ML	A075456	001	Aug 22, 2000
1.25MG/ML	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
---	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 124 (of 346)

ENOXACIN

TABLET; ORAL PENETREX	SANOFI AVENTIS US	200MG 400MG	N019616 004 N019616 005	Dec 31, 1991 Dec 31, 1991
--------------------------	-------------------	----------------	----------------------------	------------------------------

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS LOVENOX (PRESERVATIVE FREE)	SANOFI AVENTIS US	90MG/0.6ML (150MG/ML)	N020164 006	Jun 02, 2000
---	-------------------	-----------------------	-------------	--------------

EPINEPHRINE

AEROSOL, METERED; INHALATION BRONKAID MIST	STERLING	0.25MG/INH	N016803 001	
PRIMATENE MIST	WYETH CONS	0.2MG/INH	N016126 001	
INJECTABLE; INJECTION SUS-PHRINE SULFITE-FREE	FOREST LABS	1.5MG/AMP 5MG/ML	N007942 003 N007942 001	Feb 05, 1999
EPI E Z PEN JR	MERIDIAN MEDCL TECHN	0.15MG/DELIVERY	N019430 004	Aug 03, 1995
EPIPEN E Z PEN	MERIDIAN MEDCL TECHN	0.3MG/DELIVERY	N019430 003	Aug 03, 1995

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%	N017751 006	
		0.005MG/ML;1.5%	N017751 007	
	DENTSPLY PHARM	0.005MG/ML;1.5%	N021384 001	

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION CITANEST 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 001	
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE	ELKINS SINK	0.01MG/ML;1% 0.01MG/ML;2%	A080406 001 A080406 002	
	GRAHAM CHEM	0.01MG/ML;2% 0.02MG/ML;2%	A080504 004 A080504 005	Oct 19, 1983 Oct 19, 1983
	HOSPIRA	0.005MG/ML;1% 0.005MG/ML;1.5%	A089649 001 A089650 001	Jun 21, 1988 Jun 21, 1988

DISCONTINUED DRUG PRODUCT LIST

6 - 125 (of 346)

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION					
LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE					
ABBOTT	0.01MG/ML;1%		A083154	001	
BEL MAR	0.01MG/ML;1%		A080820	001	
	0.01MG/ML;2%		A080757	001	
DELL LABS	0.01MG/ML;1%		A083389	001	
	0.01MG/ML;2%		A083390	001	
INTL MEDICATION	0.01MG/ML;1%		A086402	001	
WATSON LABS	0.01MG/ML;1%		A080377	003	
	0.01MG/ML;1%		A085463	001	
	0.01MG/ML;2%		A080377	004	
LIDOCATON					
PHARMATON	0.01MG/ML;2%		A084729	001	Aug 17, 1983
	0.02MG/ML;2%		A084728	001	Aug 17, 1983
XYLOCAINE W/ EPINEPHRINE					
APP PHARMS	0.01MG/ML;2%		N006488	003	
ASTRAZENECA	0.005MG/ML;1%		N010418	006	
	0.005MG/ML;1.5%		N010418	010	
	0.005MG/ML;2%		N010418	008	
PATCH; IONTOPHORESIS, TOPICAL					
LIDOSITE TOPICAL SYSTEM KIT					
VYTERIS	1.05MG/PATCH;100MG/PATCH		N021504	001	May 06, 2004
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; IV (INFUSION)					
EPIRUBICIN HYDROCHLORIDE					
HOSPIRA	200MG/VIAL		N050807	002	Sep 15, 2006

EPLERENONE

TABLET; ORAL					
INSPRA					
GD SEARLE LLC	100MG		N021437	003	Sep 27, 2002

EPROSARTAN MESYLATE

TABLET; ORAL					
TEVETEN					
ABBOTT	EQ 300MG BASE		N020738	004	Dec 22, 1997

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	
IMPAX LABS	50,000 IU		A080951	001	
LANNETT	50,000 IU		A080825	001	
VITARINE	50,000 IU		A084053	001	
WESTWARD	50,000 IU		A083102	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 126 (of 346)

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
WATSON LABS	1MG	A086433	001	May 27, 1982
	1MG	A087244	001	Aug 16, 1982
GERIMAL WATSON LABS	1MG	A088207	001	Mar 22, 1984
HYDERGINE NOVARTIS	0.5MG	N017993	003	
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	
MUTUAL PHARM	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	
	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 PARKE DAVIS	2MG	A088337	001	Jun 08, 1984
WIGRETTES ORGANON USA INC	2MG	A086750	001	Jul 29, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 127 (of 346)

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL ERYC					
PARKE DAVIS	250MG	A062546	001	Jul 25, 1985	
	250MG	A062618	001	Sep 25, 1985	
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 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		
POWDER; FOR RX COMPOUNDING ERYTHROMYCIN					
PADDOCK LLC	100%	N050610	001	Nov 07, 1986	
SOLUTION; TOPICAL A/T/S					
TARO PHARMS NORTH	2%	A062405	001	Nov 18, 1982	
C-SOLVE-2					
BIOGLAN PHARMA	2%	A062468	001	Jul 03, 1985	
ERYDERM					
ARBOR PHARMS INC	2%	A062290	001		
ERYMAX					
MERZ PHARMS	2%	A062508	002	Jul 11, 1985	
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	
COREPHARMA	2%	A064127	001	Feb 14, 1997	
LILLY	2%	N050532	001		
PHARMAFAIR	1.5%	A062485	001	Jul 11, 1984	
	2%	A062616	001	Jul 25, 1985	
SANSAC					
DOW PHARM SCIENCES	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					
ORTHO JANSSEN	2%	N050594	001	Feb 15, 1985	
ERYTHROMYCIN COREPHARMA	2%	A064128	001	Jul 03, 1996	
T-STAT					
WESTWOOD SQUIBB	2%	A062748	001	Jul 23, 1987	
TABLET, DELAYED RELEASE; ORAL E-BASE					
BARR	333MG	A063028	001	May 15, 1990	

DISCONTINUED DRUG PRODUCT LIST

6 - 128 (of 346)

ERYTHROMYCIN

TABLET, DELAYED RELEASE; ORAL

E-BASE

BARR	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
-------	-------	---------	-----

ROBIMYCIN

ROBINS AH	250MG	A061633	001
-----------	-------	---------	-----

R-P MYCIN

SOLVAY	250MG	A061659	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
	EQ 250MG BASE/5ML	A062409	001

BARR	EQ 125MG BASE/5ML	A062169	001
	EQ 250MG BASE/5ML	A062169	002

LIFE LABS	EQ 250MG BASE/5ML	A062362	001
-----------	-------------------	---------	-----

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

BARR	EQ 200MG BASE/5ML	A062055	001
------	-------------------	---------	-----

PEDIAMYCIN			
------------	--	--	--

ROSS LABS	EQ 200MG BASE/5ML	A062305	001
-----------	-------------------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 129 (of 346)

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION; ORAL				
E-MYCIN E				
PHARMACIA AND UPJOHN	EQ 200MG BASE/5ML EQ 400MG BASE/5ML	A062198 A062198	001 002	
ERYTHROMYCIN ETHYLSUCCINATE				
ALPHARMA US PHARMS	EQ 200MG BASE/5ML EQ 400MG BASE/5ML	A062200 A062200	001 002	
DISTA	EQ 200MG BASE/5ML EQ 400MG BASE/5ML	A062177 A062177	001 002	
NASKA	EQ 400MG BASE/5ML	A062674	001	Mar 10, 1987
PARKE DAVIS	EQ 200MG BASE/5ML EQ 400MG BASE/5ML	A062231 A062231	001 002	
PHARMAFAIR	EQ 200MG BASE/5ML EQ 400MG BASE/5ML	A062559 A062558	001 001	Mar 15, 1985 Mar 15, 1985
WYAMYCIN E				
WYETH AYERST	EQ 200MG BASE/5ML EQ 400MG BASE/5ML	A062123 A062123	002 001	
SUSPENSION/DROPS; ORAL				
PEDIAMYCIN				
ROSS LABS	EQ 100MG BASE/2.5ML	A062305	002	
TABLET; ORAL				
E.E.S. 400				
ARBOR PHARMS INC	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				
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 EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	N050370 N050370 N050370	001 002 003	

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION				
ERYTHROCIN				
ABBOTT	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A062586 A062586	001 002	Jan 04, 1988 Jan 04, 1988
HOSPIRA	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	N050182 N050182	002 003	
ERYTHROMYCIN	EQ 1GM BASE/VIAL	N050609	002	Sep 24, 1986
ELKINS SINK	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A062563 A062563	001 002	Mar 28, 1985 Mar 28, 1985
ERYTHROMYCIN LACTOBIONATE				
ABRAXIS PHARM	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A062604 A062604	001 002	Nov 24, 1986 Nov 24, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 130 (of 346)

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION					
<u>ERYTHROMYCIN LACTOBIONATE</u>					
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
	EQ 1GM BASE/VIAL		A063253	002	Jul 30, 1993

ERYTHROMYCIN STEARATE

TABLET; ORAL					
<u>BRISTAMYCIN</u>					
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 INC	EQ 125MG BASE		A060359	002	
	EQ 500MG BASE		A060359	003	
ERYTHROMYCIN STEARATE					
BARR	EQ 250MG BASE		A061591	001	
	EQ 500MG BASE		A063179	001	May 15, 1990
IVAX SUB TEVA PHARMS	EQ 250MG BASE		A061461	001	
	EQ 500MG BASE		A061461	002	
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					
<u>ESCITALOPRAM OXALATE</u>					
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

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION					
<u>BREVIBLOC</u>					
BAXTER HLTHCARE CORP	10MG/ML		N019386	003	Aug 15, 1988
	20MG/ML		N019386	007	May 28, 2003

ESTAZOLAM

TABLET; ORAL					
<u>PROSOM</u>					
ABBOTT	1MG		N019080	001	Dec 26, 1990
	2MG		N019080	002	Dec 26, 1990

DISCONTINUED DRUG PRODUCT LIST

6 - 131 (of 346)

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL
ESCLIM

WOMEN FIRST HLTHCARE	0.025MG/24HR 0.0375MG/24HR 0.05MG/24HR 0.075MG/24HR 0.1MG/24HR	N020847 001 N020847 002 N020847 003 N020847 004 N020847 005	Aug 04, 1998 Aug 04, 1998 Aug 04, 1998 Aug 04, 1998 Aug 04, 1998
----------------------	--	---	--

ESTRADIOL

ORTHO MCNEIL PHARM	0.05MG/24HR 0.075MG/24HR 0.1MG/24HR	N021048 001 N021048 002 N021048 003	Sep 20, 1999 Sep 20, 1999 Sep 20, 1999
--------------------	---	---	--

FEMPATCH

PARKE DAVIS	0.025MG/24HR
-------------	--------------

N020417 001 Dec 03, 1996

VIVELLE

NOVARTIS	0.025MG/24HR 0.0375MG/24HR 0.075MG/24HR	N020323 005 N020323 001 N020323 003	Aug 16, 2000 Oct 28, 1994 Oct 28, 1994
----------	---	---	--

GEL; TOPICAL

ESTROGEL

ASCEND	0.06%	N021166 001	Feb 09, 2004
--------	-------	-------------	--------------

TABLET; ORAL

ESTRACE

BRISTOL MYERS SQUIBB	0.5MG	A081295 001	Jun 30, 1993
----------------------	-------	-------------	--------------

ESTRADIOL

AAI PHARMA INC	0.5MG 1MG 2MG	A040138 001 A040138 002 A040138 003	Jan 30, 1998 Jan 30, 1998 Jan 30, 1998
HERITAGE PHARMS INC	0.5MG 1MG 2MG	A040275 001 A040275 002 A040275 003	Dec 29, 1998 Dec 29, 1998 Dec 29, 1998

GYNODIOL

DURAMED PHARMS BARR	0.5MG 1MG 1.5MG 2MG	A040212 001 A040212 002 A040212 003 A040212 004	Dec 29, 1997 Dec 29, 1997 Dec 29, 1997 Dec 29, 1997
---------------------	------------------------------	--	--

INNOFEM

NOVO NORDISK INC	0.5MG 1MG 2MG	A040312 001 A040312 002 A040312 003	Nov 19, 1999 Nov 19, 1999 Nov 19, 1999
------------------	---------------------	---	--

TABLET; VAGINAL

VAGIFEM

NOVO NORDISK INC	25MCG	N020908 001	Mar 26, 1999
------------------	-------	-------------	--------------

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION
DEPO-ESTRADIOL

PHARMACIA AND UPJOHN	1MG/ML 3MG/ML	A085470 001 A085470 002
----------------------	------------------	----------------------------

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	N017968 001
----------------------	----------------	-------------

TESTOSTERONE CYPIONATE-ESTRADIOL CYPIONATE

WATSON LABS	2MG/ML;50MG/ML	A085603 001	Mar 13, 1986
-------------	----------------	-------------	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 132 (of 346)

ESTRADIOL VALERATE

INJECTABLE; INJECTION
 ESTRADIOL VALERATE
 WATSON LABS 10MG/ML A083546 001

ESTRADIOL VALERATE; TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION
 DITATE-DS
 SAVAGE LABS 8MG/ML;180MG/ML A086423 001
 TESTOSTERONE ENANTHATE AND ESTRADIOL VALERATE
 WATSON LABS 4MG/ML;90MG/ML A085865 001
 8MG/ML;180MG/ML A085860 001

ESTROGENS, CONJUGATED

TABLET; ORAL
 PREMARIN
 WYETH PHARMS INC 2.5MG N004782 002

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 A083266 001
 0.625MG A083266 002
 1.25MG A083266 003
 2.5MG A083266 004
 ESTERIFIED ESTROGENS
 PVT FORM 0.625MG A083414 001
 1.25MG A083765 001
 2.5MG A085907 001
 SANDOZ 1.25MG A085302 001
 ESTRATAB
 SOLVAY 0.3MG A086715 001
 0.625MG A083209 001
 1.25MG A083856 001
 2.5MG A083857 001
 EVEX
 ROCHE PALO 0.625MG A084215 001
 1.25MG A083376 002
 FEMOGEN
 PVT FORM 0.625MG A085076 001
 1.25MG A085008 001
 2.5MG A085007 001

DISCONTINUED DRUG PRODUCT LIST

6 - 133 (of 346)

ESTRONE

INJECTABLE; INJECTION ESTROGENIC SUBSTANCE				
WYETH AYERST	2MG/ML	A083488	001	
ESTRONE				
WATSON LABS	2MG/ML	A083397	001	
NATURAL ESTROGENIC SUBSTANCE-ESTRONE				
WATSON LABS	2MG/ML	A085237	001	Nov 23, 1982
THEELIN				
PARKEDALE	1MG/ML	N003977	001	
	2MG/ML	N003977	002	
	5MG/ML	N003977	003	

ESTROPIPATE

CREAM; VAGINAL OGEN				
PHARMACIA AND UPJOHN	1.5MG/GM	A084710	001	
TABLET; ORAL ESTROPIPATE				
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

ESZOPICLONE

TABLET; ORAL ESZOPICLONE				
LUPIN LTD	1MG	A091124	001	Sep 13, 2011
	2MG	A091124	002	Sep 13, 2011
	3MG	A091124	003	Sep 13, 2011
TEVA	1MG	A091169	001	May 23, 2011
	2MG	A091169	002	May 23, 2011
	3MG	A091169	003	May 23, 2011
WOCKHARDT LTD	1MG	A091165	001	Jul 14, 2011
	2MG	A091165	002	Jul 14, 2011
	3MG	A091165	003	Jul 14, 2011

ETHACRYNIC ACID

TABLET; ORAL EDECRIN				
ATON	50MG	N016092	002	

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL MYAMBUTOL				
STI PHARMA LLC	200MG	N016320	002	
	500MG	N016320	004	

ETHCHLORVYNOL

CAPSULE; ORAL ETHCHLORVYNOL				
BANNER PHARMACAPS	100MG	A084463	001	
	200MG	A084463	002	
	500MG	A084463	003	
	750MG	A084463	004	
PLACIDYL				
ABBOTT	100MG	N010021	004	
	200MG	N010021	007	
	500MG	N010021	002	
	750MG	N010021	010	

DISCONTINUED DRUG PRODUCT LIST

6 - 134 (of 346)

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 LABS	0.035MG;1MG	A072720 001 Dec 30, 1991
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 PREVEN EMERGENCY CONTRACEPTIVE KIT DURAMED	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
LEVILITE 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

DISCONTINUED DRUG PRODUCT LIST

6 - 135 (of 346)

ETHINYLMESTRADIOL; LEVONORGESTREL

TABLET; ORAL-21						
LEVORA 0.15/30-21						
WATSON LABS	0.03MG;0.15MG		A073592	001	Dec 13,	1993
NORDETTE-21						
DURAMED RES	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						
WATSON LABS	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 ETHINYLMESTRADIOL						
BARR	0.02MG;0.1MG		A075862	002	Apr 29,	2003
TRIPHASICL-28						
WYETH PHARMS INC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG		N019190	001	Nov 01,	1984

ETHINYLMESTRADIOL; NORETHINDRONE

TABLET; ORAL-21						
BALZIVA-21						
BARR	0.035MG;0.4MG		A076198	001	Apr 22,	2004
BREVICON 21-DAY						
WATSON LABS	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
NORETHINDRONE AND ETHINYLMESTRADIOL						
WATSON LABS	0.035MG;0.5MG		A070684	001	Jan 29,	1987
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 JANSEN	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,1M G		N018985	001	Apr 04,	1984
OVCON-35						
WARNER CHILCOTT	0.035MG;0.4MG		N018127	001		
OVCON-50						
WARNER CHILCOTT	0.05MG;1MG		N018128	001		
TRI-NORINYL 21-DAY						
WATSON LABS	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

DISCONTINUED DRUG PRODUCT LIST

6 - 136 (of 346)

ETHINYLMESTRADIOL; NORETHINDRONE

TABLET; ORAL-28					
NORCEPT-E 1/35 28					
ORTHO MCNEIL PHARM	0.035MG;1MG		A071546	001	Feb 09, 1989
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

ETHINYLMESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-21					
ESTROSTEP 21					
WARNER CHILCOTT	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	

ETHINYLMESTRADIOL; 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

ETHINYLMESTRADIOL; NORGESTREL

TABLET; ORAL-21					
LO/OVRAL					
WYETH PHARMS INC	0.03MG;0.3MG		N017612	001	
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					
OVRAL-28					
WYETH PHARMS	0.05MG;0.5MG		N016806	001	

ETHIODIZED OIL

OIL; INTRALYMPHATIC					
ETHIODOL					
GUERBET	99%		N009190	001	

ETHOPROPRAZINE HYDROCHLORIDE

TABLET; ORAL					
PARSIDOL					
PARKE DAVIS	10MG		N009078	003	
	50MG		N009078	006	
	100MG		N009078	008	

ETHOTOIN

TABLET; ORAL					
PEGANONE					
LUNDBECK INC	500MG		N010841	003	

ETHOXZOLAMIDE

TABLET; ORAL					
CARDRASE					
PHARMACIA AND UPJOHN	62.5MG		N011047	002	
	125MG		N011047	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 137 (of 346)

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
---	---------	--------------------------

ETODOLAC

CAPSULE; ORAL ETODOLAC AAIPHARMA LLC	300MG	A074929 001 Jan 30, 1998
ENDO PHARMS	200MG	A074842 001 Jul 17, 1997
	300MG	A074842 002 Jul 17, 1997
IVAX SUB TEVA PHARMS	200MG	A074899 001 Jul 08, 1997
	300MG	A074899 002 Jul 08, 1997
MYLAN	200MG	A074932 001 May 16, 1997
	200MG	A075071 001 Sep 30, 1998
	300MG	A074932 002 May 16, 1997
	300MG	A075071 002 Sep 30, 1998
SANDOZ	200MG	A074840 001 Aug 29, 1997
	200MG	A074942 001 Sep 30, 1997
	300MG	A074840 002 Aug 29, 1997
	300MG	A074942 002 Sep 30, 1997
TEVA	200MG	A075126 001 Sep 16, 1999
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		
AAIPHARMA LLC	400MG	A074927 001 Oct 30, 1997
ACTAVIS ELIZABETH	400MG	A074819 001 Feb 28, 1997
	500MG	A074819 002 Apr 28, 1998
ENDO PHARMS	400MG	A074841 001 Jun 27, 1997

DISCONTINUED DRUG PRODUCT LIST

6 - 138 (of 346)

ETODOLACTABLET; ORAL
ETODOLAC

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
RANBAXY	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				
POINT HOLDINGS	400MG	A075696	001	Jul 31, 2000
SANDOZ	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

ETOPOSIDECAPSULE; ORAL
VEPESID

BRISTOL MYERS SQUIBB	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
MARSAM PHARMS LLC	20MG/ML	A074968	001	Jan 09, 1998
PIERRE FABRE	20MG/ML	A074813	001	Jul 09, 1997
TEVA PARENTERAL	20MG/ML	A074510	001	Jun 29, 1995
WATSON LABS	20MG/ML	A074228	001	Oct 15, 1996
TOPOSAR				
TEVA PARENTERAL	20MG/ML	A074166	001	Feb 27, 1995
VEPESID				
BRISTOL MYERS SQUIBB	20MG/ML	N018768	001	Nov 10, 1983

ETOPOSIDE PHOSPHATEINJECTABLE; 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

ETRETINATECAPSULE; ORAL
TEGISON
ROCHE

ROCHE	10MG	N019369	001	Sep 30, 1986
	25MG	N019369	002	Sep 30, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 139 (of 346)

EVANS BLUE

INJECTABLE; INJECTION
 EVANS BLUE
 PARKE DAVIS 0.5% N008041 001

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
 FAMOTIDINE PRESERVATIVE FREE
 APOTHECON 10MG/ML A075708 001 Apr 16, 2001
 HOSPIRA 10MG/ML A075669 001 Apr 16, 2001
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER
 ABBOTT 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 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
 MUTUAL PHARM 20MG A075639 002 Dec 12, 2001
 40MG A075639 001 Dec 12, 2001
 SANDOZ 20MG A075302 001 Apr 16, 2001
 20MG A075793 001 Apr 16, 2001
 40MG A075302 002 Apr 16, 2001
 40MG A075793 002 Apr 16, 2001
 TABLET, CHEWABLE; ORAL
 PEPCID AC
 MERCK SHARP DOHME 10MG N020801 001 Sep 24, 1998
 TABLET, ORALLY DISINTEGRATING; ORAL
 FLUXID
 SCHWARZ PHARMA 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

FENOFIBRATE

CAPSULE; ORAL
 ANTARA (MICRONIZED)
 LUPIN ATLANTIS 87MG N021695 002 Nov 30, 2004
 LIPIDIL
 ABBOTT 100MG N019304 001 Dec 31, 1993
 LIPOFEN
 CIPHER PHARMS INC 100MG N021612 002 Jan 11, 2006
 TRICOR (MICRONIZED)
 ABBOTT 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
 ABBOTT 54MG N021203 001 Sep 04, 2001
 160MG N021203 003 Sep 04, 2001

DISCONTINUED DRUG PRODUCT LIST

6 - 140 (of 346)

FENOLDOPAM MESYLATEINJECTABLE; 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 CALCIUMCAPSULE; ORAL
FENOPROFEN CALCIUM

AM THERAP	EQ 200MG BASE	A072307	001	Aug 22, 1988
	EQ 300MG BASE	A072308	001	Aug 22, 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
SANDOZ	EQ 200MG BASE	A072394	001	Oct 17, 1988
	EQ 300MG BASE	A072395	001	Oct 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 300MG BASE	A072981	001	Aug 19, 1991
	EQ 300MG BASE	A072293	001	Aug 17, 1988
	EQ 300MG BASE	A072982	001	Aug 19, 1991

NALFON

PEDINOL 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
CLONMEL HLTHCARE	EQ 600MG BASE	A072326	001	Aug 17, 1988
HALSEY	EQ 600MG BASE	A072357	001	Aug 17, 1988
MUTUAL PHARM	EQ 600MG BASE	A072902	001	Dec 21, 1990
PAR PHARM	EQ 600MG BASE	A072429	001	Aug 17, 1988
QUANTUM PHARMICS	EQ 600MG BASE	A072194	001	Aug 17, 1988
SANDOZ	EQ 600MG BASE	A072396	001	Oct 17, 1988
USL PHARMA	EQ 600MG BASE	A072362	001	Aug 17, 1988
WATSON LABS	EQ 600MG BASE	A072165	001	Aug 17, 1988
	EQ 600MG BASE	A072407	001	Aug 17, 1988
	EQ 600MG BASE	A072602	001	Oct 11, 1988

NALFON

DISTA EQ 600MG BASE

N017710 001

FENTANYL CITRATEINJECTABLE; 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

MARSAM PHARMS LLC EQ 0.05MG BASE/ML

A074917 001 Feb 03, 1998

TABLET; Buccal

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

DISCONTINUED DRUG PRODUCT LIST

6 - 141 (of 346)

FENTANYL HYDROCHLORIDE

SYSTEM; IONTOPHORESIS, TRANSDERMAL
 IONSYS
 INCLINE THERAP 10.8MCG N021338 001 May 22, 2006

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

FEXOFENADINE HYDROCHLORIDE

CAPSULE; ORAL
 ALLEGRA
 SANOFI AVENTIS US 60MG N020625 001 Jul 25, 1996
 FEXOFENADINE HYDROCHLORIDE
 BARR 60MG A076169 001 Jul 13, 2005

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
 URISPAS
 ORTHO MCNEIL JANSSEN 100MG N016769 001

FLECAINIDE ACETATE

TABLET; ORAL
 FLECAINIDE ACETATE
 SANDOZ 50MG A076030 001 Oct 28, 2002
 100MG A076030 002 Oct 28, 2002
 150MG A076030 003 Oct 28, 2002
 TAMBOCOR
 MEDICIS 200MG N018830 002 Oct 31, 1985

FLOXURIDINE

INJECTABLE; INJECTION
 FUDR
 HOSPIRA 500MG/VIAL N016929 001

FLUCONAZOLE

INJECTABLE; INJECTION
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
 APOTEX INC 200MG/100ML (2MG/ML) A076889 001 Mar 25, 2005
 400MG/200ML (2MG/ML) A076889 002 Mar 25, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 142 (of 346)

FLUCONAZOLE

INJECTABLE; INJECTION

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

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
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
SANDOZ	50MG	A076086	001	Jul 29, 2004
	100MG	A076086	002	Jul 29, 2004
	150MG	A076086	003	Jul 29, 2004
	200MG	A076086	004	Jul 29, 2004

FLUDEOXYGLUCOSE F-18

INJECTABLE; INJECTION

FLUDEOXYGLUCOSE F 18

DOWNSTATE CLINCL	4-40mCi/ML	N020306	001	Aug 19, 1994
	4-90mCi/ML	N020306	002	Sep 25, 2001

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

KING PHARMS	0.1MG	N010060	001
-------------	-------	---------	-----

FLUMETHASONE PIVALATE

CREAM; TOPICAL

LOCORTEN

NOVARTIS	0.03%	N016379	001
----------	-------	---------	-----

FLUNISOLIDE

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
------------------	--------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 143 (of 346)

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL					
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
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

FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM; TOPICAL					
NEO-SYNALAR					
MEDIMETRIKS PHARMS	0.025%;EQ 3.5MG BASE/GM		A060700	001	

FLUOCINONIDE

CREAM; TOPICAL					
FLUOCINONIDE					
PERRIGO NEW YORK	0.05%		A071790	001	Jul 13, 1988
TARO	0.05%		A071500	001	Jun 10, 1987
FLUOCINONIDE EMULSIFIED BASE					
ACTAVIS MID ATLANTIC	0.05%		A074204	001	Jun 13, 1995
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	
INJECTABLE; INTRAVENOUS					
AK-FLUOR 25%					
AKORN	EQ 500MG BASE/2ML (EQ 250MG BASE/ML)		N022186	002	Aug 08, 2008

FLUOROMETHOLONE

CREAM; TOPICAL					
OXYLONE					
PHARMACIA AND UPJOHN	0.025%		N011748	001	
SUSPENSION/DROPS; OPHTHALMIC					
FLUOR-OP					
NOVARTIS	0.1%		A070185	001	Feb 27, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 144 (of 346)

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
 50MG/ML A089519 001 Mar 12, 1987
 APP PHARMS 50MG/ML A040291 001 Mar 24, 1999
 50MG/ML A040379 001 Nov 15, 2000
 BEDFORD 50MG/ML A089508 001 Jan 26, 1988
 MARCHAR 50MG/ML A087791 001 Jan 18, 1983
 SMITH AND NEPHEW 50MG/ML A088766 001 Dec 28, 1984
 50MG/ML A088767 001 Dec 28, 1984
 50MG/ML A089434 001 Mar 26, 1987
 VALEANT 500MG/10ML (50MG/ML) N012209 001
 WATSON LABS 50MG/ML A087792 001 Oct 13, 1982
 SOLUTION; TOPICAL
 FLUOROPLEX
 ELORAC 1% N016765 001

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL
 FLUOXETINE
 MUTUAL PHARMA 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
 BARR EQ 40MG BASE A076251 001 May 18, 2005
 PAR PHARM EQ 10MG BASE A076922 001 Dec 16, 2004
 EQ 20MG BASE A076922 002 Dec 16, 2004
 EQ 40MG BASE A076922 003 Dec 16, 2004
 SANDOZ EQ 10MG BASE A075807 001 Jan 29, 2002
 EQ 20MG BASE A075807 002 Jan 29, 2002
 PROZAC
 LILLY EQ 60MG BASE N018936 004 Jun 15, 1999
 SOLUTION; ORAL
 FLUOXETINE HYDROCHLORIDE
 ACTAVIS MID ATLANTIC EQ 20MG BASE/5ML A075690 001 Jan 31, 2002
 HI TECH PHARMA EQ 20MG BASE/5ML A075525 001 Jun 27, 2002
 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 A076024 001 Jan 29, 2002

DISCONTINUED DRUG PRODUCT LIST

6 - 145 (of 346)

FLUOXETINE HYDROCHLORIDE

TABLET; ORAL PROZAC LILLY	EQ 10MG BASE EQ 20MG BASE	N020974 001 Mar 09, 1999 N020974 002 Mar 09, 1999
---------------------------------	------------------------------	--

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
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 TEVA PHARMS	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 TEVA PHARMS	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 2.5MG 5MG 10MG	A088555 001 Dec 18, 1987 A088544 001 Dec 18, 1987 A088527 001 Dec 18, 1987 A088550 001 Dec 18, 1987
PERMITIL SCHERING	0.25MG 2.5MG 5MG 10MG	N012034 001 N012034 004 N012034 005 N012034 006

DISCONTINUED DRUG PRODUCT LIST

6 - 146 (of 346)

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL PROLIXIN APOTHECON	1MG 2.5MG 5MG 10MG	N011751 004 N011751 001 N011751 003 N011751 002
TABLET, EXTENDED RELEASE; ORAL PERMITIL SCHERING	1MG	N012419 004

FLUPREDNISOLONE

TABLET; ORAL ALPHADROL PHARMACIA AND UPJOHN	1.5MG	N012259 002
---	-------	-------------

FLURANDRENOLIDE

CREAM; TOPICAL CORDRAN SP WATSON PHARMS	0.05%	N012806 002
LOTION; TOPICAL FLURANDRENOLIDE ALPHARMA US PHARMS	0.05%	A087203 001 Apr 29, 1982
OINTMENT; TOPICAL CORDRAN WATSON PHARMS	0.025% 0.05%	N012806 004 N012806 001

FLURANDRENOLIDE; NEOMYCIN SULFATE

CREAM; 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 HYDROCHLORIDE

CAPSULE; ORAL DALMANE VALEANT PHARM INTL	15MG 30MG	N016721 001 N016721 002
FLURAZEPAM HYDROCHLORIDE HALSEY	15MG 30MG	A071808 001 Jan 07, 1988 A071809 001 Jan 07, 1988
MUTUAL PHARM	15MG 30MG	A070454 001 Aug 04, 1986 A070455 001 Aug 04, 1986
PAR PHARM	15MG 30MG	A070444 001 Mar 20, 1986 A070445 001 Mar 20, 1986
PUREPAC PHARM	15MG 30MG	A071927 001 Sep 09, 1987 A071551 001 Sep 09, 1987
SANDOZ	15MG 30MG	A071716 001 Jul 31, 1991 A071717 001 Jul 31, 1991
SUPERPHARM	15MG 30MG	A071659 001 Aug 04, 1988 A071660 001 Aug 04, 1988
USL PHARMA	15MG 30MG	A070562 001 Jul 09, 1987 A070563 001 Jul 09, 1987
WARNER CHILCOTT	15MG 30MG	A071767 001 Dec 04, 1987 A071768 001 Dec 04, 1987
WATSON LABS	15MG	A072368 001 Mar 30, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 147 (of 346)

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

IVAX SUB TEVA PHARMS	50MG 100MG	A074411 001 A074411 002	May 31, 1995 May 31, 1995
PLIVA	50MG 100MG	A074647 001 A074647 002	Apr 01, 1997 Apr 01, 1997
SANDOZ	50MG 100MG	A074448 001 A074448 002	Jul 28, 1995 Jul 28, 1995
TEVA	50MG 100MG	A074405 002 A074405 001	May 24, 1995 May 24, 1995
THERAGEN	100MG	A074560 002	May 16, 1997

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

SCHERING

125MG

N018554 001 Jan 27, 1989

FLUTICASONE PROPIONATE

AEROSOL, METERED; INHALATION

FLOVENT

GLAXOSMITHKLINE	0.044MG/INH 0.11MG/INH 0.22MG/INH	N020548 001 N020548 002 N020548 003	Mar 27, 1996 Mar 27, 1996 Mar 27, 1996
-----------------	---	---	--

OINTMENT; TOPICAL

FLUTICASONE PROPIONATE

TARO PHARM IND'S 0.005%

A077145 001 Jun 14, 2005

POWDER; INHALATION

FLOVENT

GLAXOSMITHKLINE	0.044MG/INH 0.088MG/INH 0.22MG/INH	N020549 001 N020549 002 N020549 003	Nov 07, 1997 Nov 07, 1997 Nov 07, 1997
-----------------	--	---	--

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

ACTAVIS ELIZABETH	25MG 50MG 100MG	A075901 001 A075901 002 A075901 003	Dec 28, 2000 Dec 28, 2000 Dec 28, 2000
-------------------	-----------------------	---	--

IVAX SUB TEVA PHARMS	25MG 50MG 100MG	A075898 001 A075898 002 A075898 003	Mar 12, 2001 Mar 12, 2001 Mar 12, 2001
----------------------	-----------------------	---	--

MUTUAL PHARM	25MG 50MG 100MG	A076125 001 A076125 002 A076125 003	Apr 29, 2002 Apr 29, 2002 Apr 29, 2002
--------------	-----------------------	---	--

MYLAN	50MG 100MG	A075950 001 A075950 002	Oct 15, 2001 Oct 15, 2001
-------	---------------	----------------------------	------------------------------

SANDOZ	25MG 50MG 100MG	A075887 001 A075887 002 A075887 003	Jan 05, 2001 Jan 05, 2001 Jan 05, 2001
--------	-----------------------	---	--

SYNTTHON PHARMS	25MG 50MG 100MG	A075899 001 A075899 002 A075899 003	Jan 17, 2001 Jan 17, 2001 Jan 17, 2001
-----------------	-----------------------	---	--

WATSON LABS	25MG 50MG 100MG	A075894 001 A075894 002 A075894 003	Apr 18, 2001 Apr 18, 2001 Apr 18, 2001
-------------	-----------------------	---	--

LUVOX	25MG 50MG 100MG 150MG	N020243 001 N020243 002 N020243 003 N020243 004	Dec 05, 1994 Dec 05, 1994 Dec 05, 1994 Dec 05, 1994
-------	--------------------------------	--	--

DISCONTINUED DRUG PRODUCT LIST

6 - 148 (of 346)

FOLIC ACID

INJECTABLE; INJECTION					
FOLIC ACID					
BEN VENUE	5MG/ML	A081066	001	Dec 29,	1993
FOLVITE		N005897	008		
HIKMA (MAPLE)	5MG/ML				
TABLET; ORAL					
FOLIC ACID					
BARR	1MG	A089177	001	Jan 08,	1986
EVERYLIFE	1MG	A080755	001		
HALSEY	1MG	A083598	001		
IMPAK LABS	1MG	A080686	001		
IVAX SUB TEVA PHARMS	1MG	A083000	001		
LANNETT	1MG	A080816	001		
LILLY	1MG	N006135	003		
MK LABS	1MG	A083526	001		
MUTUAL PHARM	1MG	A040582	001	Jul 18,	2005
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		
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					
HIKMA (MAPLE)	1MG	N005897	004		

FOLLITROPIN ALFA/BETA

INJECTABLE; IM-SC					
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	150 IU/0.18ML	N021211	003	Feb 11,	2004
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

FOMIVIRSEN SODIUM

INJECTABLE; INJECTION					
VITRAVENE PRESERVATIVE FREE					
NOVARTIS	6.6MG/ML	N020961	001	Aug 26,	1998

FORMOTEROL FUMARATE

POWDER; INHALATION					
FORADIL CERTIHALER					
NOVARTIS	0.0085MG/INH	N021592	001	Dec 15,	2006

DISCONTINUED DRUG PRODUCT LIST

6 - 149 (of 346)

FOSINOPRIL SODIUM

TABLET; ORAL					
	FOSINOPRIL SODIUM				
SANDOZ	10MG		A076188	001	Oct 08, 2004
	20MG		A076188	002	Oct 08, 2004
	40MG		A076188	003	Oct 08, 2004

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL					
	FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE				
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					
CEREBYX					
PARKE DAVIS	EQ 50MG PHENYTOIN NA/ML		N020450	001	Aug 05, 1996

FURAZOLIDONE

SUSPENSION; ORAL					
FUROXONE					
SHIRE	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
ASTRAZENECA	10MG/ML		A070014	001	Sep 09, 1985
	10MG/ML		A070095	001	Sep 09, 1985
	10MG/ML		A070096	001	Sep 09, 1985
BAXTER HLTHCARE	10MG/ML		A071439	001	Sep 14, 1990
	10MG/ML		N018267	001	
HOSPIRA	10MG/ML		A072080	001	Aug 13, 1991
	10MG/ML		A074337	001	Oct 31, 1994
MARSAM PHARMS LLC	10MG/ML		A074017	001	Jun 30, 1994
ORGANON USA INC	10MG/ML		A070017	001	Dec 15, 1986
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
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					
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
MUTUAL PHARM	20MG		A070043	001	Sep 26, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 150 (of 346)

FUROSEMIDE

TABLET; ORAL

FUROSEMIDE

MUTUAL PHARM	40MG	N018790	001	Nov 29, 1983
	80MG	A070100	001	Jan 26, 1988
SANDOZ	40MG	N018750	002	Jul 30, 1984
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	N018369	001	May 14, 1982
	40MG	A070413	001	Feb 26, 1986
	40MG	N018369	002	May 14, 1982

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

IVAX SUB TEVA PHARMS	100MG	A075477	001	Mar 23, 2005
	300MG	A075477	002	Mar 23, 2005
	400MG	A075477	003	Mar 23, 2005
MUTUAL PHARM	100MG	A076537	001	Jun 30, 2005
	300MG	A076537	002	Jun 30, 2005
	400MG	A076537	003	Jun 30, 2005
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

TABLET; ORAL

GABAPENTIN

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
---------------	-------------------------	---------	-----	--------------

GALANTAMINE HYDROBROMIDE

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
YABAO BIOPHARMS	EQ 4MG BASE	A077604	001	Feb 06, 2009
	EQ 8MG BASE	A077604	002	Feb 06, 2009
	EQ 12MG BASE	A077604	003	Feb 06, 2009

GALLAMINE TRIETHIODIDE

INJECTABLE; INJECTION

FLAXEDIL

DAVIS AND GECK	20MG/ML	N007842	001
	100MG/ML	N007842	002

DISCONTINUED DRUG PRODUCT LIST

6 - 151 (of 346)

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION			
GALLIUM CITRATE GA 67			
GE HEALTHCARE	1mCi/ML	N017700	001
NEOSCAN			
GE HEALTHCARE	2mCi/ML	N017655	001

GANCICLOVIR

CAPSULE; ORAL			
CYTOVENE			
ROCHE PALO	250MG	N020460	001 Dec 22, 1994
	500MG	N020460	002 Dec 12, 1997

GANCICLOVIR SODIUM

INJECTABLE; INJECTION			
GANCICLOVIR SODIUM			
BEDFORD	EQ 500MG BASE/VIAL	A076222	001 Jul 16, 2003

GEFITINIB

TABLET; ORAL			
IRESSA			
ASTRAZENECA	250MG	N021399	001 May 05, 2003

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			
MYLAN	600MG	A074452	001 Feb 16, 1995
PUREPAC PHARM	600MG	A074360	001 Aug 31, 1994
SANDOZ	600MG	A074615	001 Sep 29, 1995
WATSON LABS	600MG	A074156	001 Oct 24, 1994

GEMTUZUMAB OZOGAMICIN

INJECTABLE; INJECTION			
MYLOTARG			
WYETH PHARMS INC	5MG/VIAL	N021174	001 May 17, 2000

GENTAMICIN SULFATE

CREAM; TOPICAL			
GARAMYCIN			
SCHERING	EQ 0.1% BASE	A060462	001
GENTAFAIR			
PARMAFAIR	EQ 0.1% BASE	A062458	001 Sep 01, 1983
GENTAMICIN SULFATE			
ALPHARMA US PHARMS	EQ 0.1% BASE	A062471	001 Sep 27, 1983
BAUSCH AND LOMB	EQ 0.1% BASE	A064056	001 Apr 29, 1994
PHARMADERM	EQ 1MG BASE/GM	A062530	001 Jul 05, 1984
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

DISCONTINUED DRUG PRODUCT LIST

6 - 152 (of 346)

GENTAMICIN SULFATE

INJECTABLE; INJECTION

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

BAXTER HLTHCARE

EQ 10MG BASE/ML

A062251 002

EQ 40MG BASE/ML

A062251 001

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

WYETH AYERST

EQ 10MG BASE/ML

A062264 001

EQ 40MG BASE/ML

A062264 002

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA

EQ 1.2MG BASE/ML

A062588 001 Jan 06, 1986

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

A062588 004 Jan 06, 1986

EQ 2MG BASE/ML

A062588 005 Jan 06, 1986

EQ 60MG BASE/100ML

A062588 006 Jan 06, 1986

EQ 70MG BASE/100ML

A062588 007 Jan 06, 1986

EQ 80MG BASE/100ML

A062588 008 Jan 06, 1986

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

BAUSCH AND LOMB

EQ 0.1% BASE

A064054 001 Apr 29, 1994

PHARMADERM

EQ 0.1% BASE

A062534 001 Oct 10, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 153 (of 346)

GENTAMICIN SULFATE

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 UNIVERSAL	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	20MG/VIAL	N020622	001	Dec 20, 1996

GLIMEPIRIDE

TABLET; ORAL GLIMEPIRIDE				
RANBAXY	3MG	A077366	001	Oct 06, 2005
	6MG	A077366	002	Oct 06, 2005

GLIPIZIDE

TABLET; ORAL GLIPIZIDE				
ENDO PHARMS	5MG	A074378	001	Nov 28, 1994
	10MG	A074378	002	Nov 28, 1994
PLIVA	5MG	A074619	001	Apr 04, 1997
	10MG	A074619	002	Apr 04, 1997
SANDOZ	5MG	A074542	001	Jun 20, 1995
	10MG	A074542	002	Jun 20, 1995
TEVA	5MG	A074387	001	Mar 04, 1996
	10MG	A074387	002	Mar 04, 1996
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 - 154 (of 346)

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		
SANDOZ	500MG	A083234	002		
UCB INC	500MG	A085171	001		
VITARINE	500MG	A087297	001		
WATSON LABS	500MG	A084362	001		
	500MG	A085763	001		

GLYBURIDE

TABLET; ORAL GLYBURIDE					
ACTAVIS TOTOWA	1.5MG	A075947	001	Nov 14, 2002	
	3MG	A075947	002	Nov 14, 2002	
	6MG	A075947	003	Nov 14, 2002	
GLYBURIDE (MICRONIZED)					
SANDOZ	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 GLYBURIDE AND METFORMIN HYDROCHLORIDE					
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		

GLCOPYRROLATE

INJECTABLE; INJECTION GLCOPYRROLATE					
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		
TABLET; ORAL GLCOPYRROLATE					
WATSON LABS	1MG	A085562	001		
	1MG	A086902	001		
	2MG	A085563	001		
	2MG	A086178	001		

DISCONTINUED DRUG PRODUCT LIST

6 - 155 (of 346)

GLYCOPYRROLATE

TABLET; ORAL
 GLYCOPYRROLATE
 WATSON LABS

2MG

A086900 001

GONADORELIN 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

HIKMA (MAPLE)

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

APP PHARMS

5,000 UNITS/VIAL
15,000 UNITS/VIAL
20,000 UNITS/VIALN017067 001
N017067 004
N017067 003

BEL MAR

5,000 UNITS/VIAL
10,000 UNITS/VIALN017054 001
N017054 002

FERRING

2,000 UNITS/VIAL
2,000 UNITS/VIAL
5,000 UNITS/VIAL
15,000 UNITS/VIAL
20,000 UNITS/VIALN017016 009 Dec 27, 1984
N017016 011 Feb 16, 1990
N017016 006
N017016 010 Feb 15, 1985
N017016 004

FOLLUTEIN

BRISTOL MYERS SQUIBB

10,000 UNITS/VIAL

N017056 001

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

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

NEO-POLYCIN

DOW PHARM

0.025MG/ML; EQ 1.75MG BASE/ML; 10,000 UNITS/ML

A060427 001

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

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

KYTRIL

ROCHE

EQ 2MG BASE/10ML

N021238 001 Jun 27, 2001

DISCONTINUED DRUG PRODUCT LIST

6 - 156 (of 346)

GRANISETRON HYDROCHLORIDE

TABLET; ORAL					
<u>GRANISETRON HYDROCHLORIDE</u>					
BARR	EQ 1MG BASE		A078221	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					
<u>RAXAR</u>					
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					
<u>GRISACTIN</u>					
WYETH AYERST	125MG		N050051	002	
	250MG		N050051	001	
<u>SUSPENSION; ORAL</u>					
<u>GRIFULVIN V</u>					
JOHNSON AND JOHNSON	125MG/5ML		N050448	001	
TABLET; ORAL					
<u>FULVICIN-U/F</u>					
ELORAC	250MG		A060569	002	
	500MG		A060569	001	
<u>GRIFULVIN V</u>					
J AND J	125MG		A060618	001	
	250MG		A060618	002	
	500MG		A060618	003	
ORTHONEUTROGENA	125MG		A062279	001	
	250MG		A062279	002	
GRISACTIN					
WYETH AYERST	500MG		A060212	001	

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL					
<u>FULVICIN P/G</u>					
ELORAC	125MG		A061996	001	
	250MG		A061996	002	
<u>FULVICIN P/G 165</u>					
ELORAC	165MG		A061996	003	Apr 06, 1982
FULVICIN P/G 330					
ELORAC	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

GUANABENZ ACETATE

TABLET; ORAL					
<u>GUANABENZ ACETATE</u>					
SANDOZ	EQ 4MG BASE		A074517	001	Sep 30, 1998
	EQ 8MG BASE		A074517	002	Sep 30, 1998
TEVA PHARMS	EQ 4MG BASE		A074267	001	Jun 01, 1994
	EQ 8MG BASE		A074267	002	Jun 01, 1994
WATSON LABS	EQ 4MG BASE		A074025	001	Feb 28, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 157 (of 346)

GUANABENZ ACETATE

TABLET; ORAL GUANABENZ ACETATE				
WATSON LABS	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	N018104	001	Dec 29, 1982
	25MG	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

HALAZEPAM

TABLET; ORAL PAXIPAM				
SCHERING	20MG	N017736	003	
	40MG	N017736	004	

HALCINONIDE

CREAM; TOPICAL HALOG				
WESTWOOD SQUIBB	0.025%	N017818	001	
HALOG-E				
RANBAXY	0.1%	N018234	001	
OINTMENT; TOPICAL HALOG				
BRISTOL MYERS SQUIBB	0.025%	N018125	001	
SOLUTION; TOPICAL HALOG				
RANBAXY	0.1%	N017823	001	

HALOBETASOL PROPIONATE

OINTMENT; TOPICAL HALOBETASOL PROPIONATE				
ACTAVIS MID ATLANTIC	0.05%	A077109	001	Jun 14, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 158 (of 346)

HALOFANTRINE HYDROCHLORIDE

TABLET; ORAL

HALFAN

GLAXOSMITHKLINE 250MG

N020250 001 Jul 24, 1992

HALOPERIDOL

TABLET; ORAL

HALDOL

ORTHO MCNEIL 0.5MG
1MG
2MG
5MG
10MG
20MGN015921 001
N015921 002
N015921 003
N015921 004
N015921 005
N015921 006 Feb 02, 1982

HALDOL SOLUTAB

ORTHO MCNEIL PHARM 1MG

N017079 001

HALOPERIDOL

DURAMED PHARMS BARR 0.5MG
1MG
2MG
5MG
10MG
20MGA071216 001 Dec 04, 1986
A071217 001 Dec 04, 1986
A071218 001 Dec 04, 1986
A071219 001 Dec 04, 1986
A071220 001 Jul 07, 1987
A071221 001 Jul 07, 1987LEDERLE 0.5MG
1MG
2MG
5MG
10MG
20MGA072727 001 Sep 19, 1989
A072728 001 Sep 19, 1989
A072729 001 Sep 19, 1989
A072730 001 Sep 19, 1989
A072731 001 Sep 19, 1989
A072732 001 Sep 19, 1989MUTUAL PHARM 0.5MG
1MG
2MG
5MG
10MG
20MGA071156 001 Jan 02, 1987
A071157 001 Jan 02, 1987
A071172 001 Jan 02, 1987
A071212 001 Jan 07, 1988
A071173 001 Jan 07, 1988
A071177 001 Jan 07, 1988PAR PHARM 20MG
PUREPAC PHARM 0.5MG
1MG
2MG
5MG
10MG
20MGA071328 001 Jul 20, 1987
A071071 001 Nov 03, 1986
A071072 001 Nov 03, 1986
A071073 001 Nov 03, 1986
A071074 001 Nov 03, 1986
A071075 001 Aug 04, 1987
A071076 001 Aug 04, 1987QUANTUM PHARMICS 0.5MG
1MG
2MG
5MGA071255 001 Feb 17, 1987
A071269 001 Feb 17, 1987
A071256 001 Feb 17, 1987
A071257 001 Feb 17, 1987ROXANE 0.5MG
1MG
2MG
5MG
10MG
20MGA071128 001 Feb 17, 1987
A071129 001 Feb 17, 1987
A071130 001 Feb 17, 1987
A071131 001 Feb 17, 1987
A071132 001 May 12, 1987
A071133 001 May 12, 1987ROYCE LABS 0.5MG
1MG
2MG
5MG
10MG
20MGA071722 001 Dec 24, 1987
A071723 001 Dec 24, 1987
A071724 001 Dec 24, 1987
A071725 001 Dec 24, 1987
A072121 001 Dec 24, 1987
A072122 001 Dec 24, 1987SCS 0.5MG
1MG
2MG
5MG
10MGA070720 001 Jun 10, 1986
A070721 001 Jun 10, 1986
A070722 001 Jun 10, 1986
A070723 001 Jun 10, 1986
A070724 001 Jun 10, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 159 (of 346)

HALOPERIDOLTABLET; ORAL
HALOPERIDOL

SCS	20MG	A070725	001	Sep 24, 1986
VINTAGE	0.5MG	A071233	001	Nov 03, 1986
	1MG	A071234	001	Nov 03, 1986
	2MG	A071235	001	Nov 03, 1986
	5MG	A071236	001	Nov 03, 1986
	10MG	A071237	001	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 DECANOATEINJECTABLE; INJECTION
HALOPERIDOL DECANOATE

HOSPIRA	EQ 50MG BASE/ML	A075176	001	Feb 09, 2000
	EQ 100MG BASE/ML	A075176	002	Feb 09, 2000
SANDOZ	EQ 50MG BASE/ML	A076463	001	Jun 24, 2005
	EQ 100MG BASE/ML	A076463	002	Jun 24, 2005

HALOPERIDOL LACTATECONCENTRATE; 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				
ROXANE	EQ 2MG BASE/ML	A072045	001	Apr 12, 1988
INJECTABLE; INJECTION				
HALOPERIDOL				
ABRAXIS PHARM	EQ 5MG BASE/ML	A071187	001	Jan 20, 1987
MARSAM PHARMS LLC	EQ 5MG BASE/ML	A072516	001	Feb 25, 1993
	EQ 5MG BASE/ML	A072517	001	Feb 25, 1993
SANDOZ	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	
-----------------	----	---------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 160 (of 346)

HALOPROGIN

SOLUTION; TOPICAL HALOTEX WESTWOOD SQUIBB	1%	N016943 001
---	----	-------------

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
INTL 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
APP PHARMS	1,000 UNITS/ML	N017651 005
	5,000 UNITS/ML	N017029 002
	10,000 UNITS/ML	N017651 003
	20,000 UNITS/ML	N017651 008
BAXTER HLTHCARE	5,000 UNITS/0.5ML	N017037 013 Apr 07, 1986
BAXTER HLTHCARE CORP	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
	7,500 UNITS/ML	N017007 003
	10,000 UNITS/ML	N017007 004
	15,000 UNITS/ML	N017007 005
	20,000 UNITS/ML	N017007 006

DISCONTINUED DRUG PRODUCT LIST

6 - 161 (of 346)

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

CHAMBERLIN PARENTERL	1,000 UNITS/ML 5,000 UNITS/ML 10,000 UNITS/ML 20,000 UNITS/ML	N017130 001 N017130 002 N017130 003 N017130 004	
DELL LABS	1,000 UNITS/ML 5,000 UNITS/ML 10,000 UNITS/ML 20,000 UNITS/ML 40,000 UNITS/ML	N017540 001 N017540 002 N017540 003 N017540 004 N017540 005	
HOSPIRA	2,500 UNITS/ML 10,000 UNITS/ML	A088099 001 A040095 001	Apr 28, 1983 Jul 26, 1996
LILLY	1,000 UNITS/ML 10,000 UNITS/ML 20,000 UNITS/ML	N005521 001 N005521 002 N005521 004	
LUITPOLD	1,000 UNITS/ML	A087452 001	Oct 31, 1983
MARSAM PHARMS LLC	1,000 UNITS/ML 1,000 UNITS/ML	A040007 001 A040008 001	Jun 07, 1996 Oct 10, 1995
ORGANON USA INC	1,000 UNITS/ML 5,000 UNITS/ML 10,000 UNITS/ML	N000552 008 N000552 009 N000552 010	
PARKE DAVIS	1,000 UNITS/ML 5,000 UNITS/ML 7,500 UNITS/ML 10,000 UNITS/ML 20,000 UNITS/ML	N017346 001 N017346 002 N017346 003 N017346 004 N017346 005	
PHARM SPEC	1,000 UNITS/ML 5,000 UNITS/ML 10,000 UNITS/ML 20,000 UNITS/ML 40,000 UNITS/ML	N017780 001 N017780 002 N017780 003 N017780 004 N017780 005	
PHARMACIA AND UPJOHN	1,000 UNITS/ML 5,000 UNITS/ML 10,000 UNITS/ML	N004570 001 N004570 002 N004570 003	
SMITH AND NEPHEW	1,000 UNITS/ML	A088239 001	Jul 26, 1984
SOLOPAK	1,000 UNITS/ML 5,000 UNITS/ML 5,000 UNITS/0.5ML 10,000 UNITS/ML 10,000 UNITS/0.5ML	A087043 001 A087077 001 A087395 001 A087107 001 A087363 001	
WATSON LABS	1,000 UNITS/ML 2,500 UNITS/ML 3,000 UNITS/ML 4,000 UNITS/ML 5,000 UNITS/ML 6,000 UNITS/ML 7,500 UNITS/ML 10,000 UNITS/ML 20,000 UNITS/ML 40,000 UNITS/ML	N017064 002 N017064 015 N017064 016 N017064 017 N017064 003 N017064 018 N017064 019 N017064 004 N017064 005 N017064 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 IN DEXTROSE 5%			
HOSPIRA	10,000 UNITS/100ML	N018911 006	Jan 30, 1985
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	2,000 UNITS/100ML	N018814 002	Jul 09, 1985
HEPARIN SODIUM 10,000 UNITS IN SODIUM CHLORIDE 0.45%			
HOSPIRA	10,000 UNITS/100ML 10,000 UNITS/100ML	N018911 001 N018916 005	Jan 30, 1985 Jan 31, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 162 (of 346)

HEPARIN SODIUM

INJECTABLE; INJECTION

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 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
MARSAM PHARMS LLC	1,000 UNITS/ML	A089464	001	Jun 03, 1986
PHARMA SERVE NY	1,000 UNITS/ML	A086129	001	
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 163 (of 346)

HEPARIN SODIUM

INJECTABLE; INJECTION					
LIQUAEMIN SODIUM					
ORGANON USA INC	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

HETACILLIN

FOR 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 POTASSIUM

CAPSULE; ORAL					
VERSAPEN-K					
BRISTOL	EQ 225MG AMPICIL		A061396	001	
	EQ 450MG AMPICIL		A061396	002	

HEXACHLOROPHENE

AEROSOL; TOPICAL					
SEPTISOL					
VESTAL LABS	0.23%		N017424	001	
TURGEX					
XTTRIUM	3%		N018375	001	
EMULSION; TOPICAL					
HEXA-GERM					
HUNTINGTON LABS	3%		N017411	001	
PHISOHEX					
SANOFI AVENTIS US	3%		N008402	001	
SOY-DOME					
BAYER PHARMS	3%		N017405	001	
TURGEX					
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 164 (of 346)

HEXACHLOROPHEN

SOLUTION; TOPICAL SEPTISOL	0 . 25 %	N017423	001
VESTAL LABS			
SPONGE; TOPICAL E-Z SCRUB	450MG	N017452	001
BECTON DICKINSON			
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

HEXYOCYCLIUM METHYLSULFATE

TABLET; ORAL TRAL			
ABBOTT	25MG	N010599	001

HEXYLCAINE HYDROCHLORIDE

SOLUTION; TOPICAL CYCLAINE			
MERCK	5 %	N008472	001

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

DISCONTINUED DRUG PRODUCT LIST

6 - 165 (of 346)

HOMATROPRINE METHYLBROMIDE; HYDROCODONE BITARTRATE

TABLET; ORAL

HOMATROPRINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

ACTAVIS TOTOWA	1.5MG;5MG	A040295	001	Dec 01, 2000
HYCODAN		N005213	001	Jul 26, 1988

HYALURONIDASE

INJECTABLE; INJECTION

VITRASE

ISTA PHARMS	6,200 UNITS/VIAL	N021640	001	May 05, 2004
WYDASE		N006343	002	
BAXTER HLTHCARE	150 UNITS/ML	N006343	006	
	150 UNITS/VIAL	N006343	005	
	1,500 UNITS/VIAL			

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 TOTOWA	25MG	A088560	001	Oct 04, 1984
	50MG	A088649	001	Oct 18, 1984
ASCOT	25MG	A088310	001	Dec 19, 1984
	50MG	A088311	001	Dec 19, 1984
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
IMPAK 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
	10MG	A089359	001	Jul 25, 1986
	25MG	A084106	002	
	25MG	A089258	001	May 05, 1986
	50MG	A084107	002	
	50MG	A089259	001	May 05, 1986
	100MG	A088729	001	Apr 11, 1985
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
SANDOZ	10MG	A083241	001	
	25MG	A083560	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 166 (of 346)

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

SANDOZ	50MG	A083561	001	
	50MG	A085088	001	
SUPERPHARM	10MG	A088787	001	Aug 28, 1984
	25MG	A088788	001	Aug 28, 1984
	50MG	A088789	001	Aug 28, 1984
TG UNITED LABS	10MG	A088846	001	Feb 26, 1985
	25MG	A088847	001	Feb 26, 1985
	50MG	A088848	001	Feb 26, 1985
	100MG	A088849	001	Feb 26, 1985
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	A085532	002	May 24, 1982
	50MG	A085533	002	May 25, 1982
WEST WARD	25MG	A088240	001	May 27, 1983
	50MG	A088241	001	May 27, 1983

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

NOVARTIS	25MG;25MG	A084735	001	
	50MG;50MG	A084810	001	
	100MG;50MG	A084811	001	

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
-------------	-----------	---------	-----	--------------

HYDRA-ZIDE

PAR PHARM	100MG;50MG	A088961	001	Oct 21, 1985
-----------	------------	---------	-----	--------------

TABLET; ORAL

APRESOLINE-ESIDRIX

NOVARTIS	25MG;15MG	N012026	002	
----------	-----------	---------	-----	--

HYDRALAZINE AND HYDROCHLORTIAZIDE

WATSON LABS	25MG;15MG	A085827	001	
-------------	-----------	---------	-----	--

HYDROCHLORTIAZIDE W/ HYDRALAZINE

WATSON LABS	25MG;15MG	A085373	001	
-------------	-----------	---------	-----	--

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

TG UNITED LABS	25MG;15MG;0.1MG	A084897	001	
----------------	-----------------	---------	-----	--

HYDRALAZINE HYDROCHLORIDE, HYDROCHLORTIAZIDE AND RESERPINE

IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A084291	001	
----------------------	-----------------	---------	-----	--

HYDRALAZINE HYDROCHLORIDE-HYDROCHLORTIAZIDE-RESERPINE

MYLAN	25MG;15MG;0.1MG	A087085	001	
-------	-----------------	---------	-----	--

HYDRALAZINE, HYDROCHLORTIAZIDE W/ RESERPINE

WATSON LABS	25MG;15MG;0.1MG	A085771	001	
-------------	-----------------	---------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 167 (of 346)

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL				
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				
MUTUAL PHARM	25MG;15MG;0.1MG		A088570	001 Apr 10, 1984
SOLVAY	25MG;15MG;0.1MG		A088376	001 Oct 28, 1983
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

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				
ESISDIRIX				
NOVARTIS	25MG		N011793	005
	50MG		N011793	008
	100MG		N011793	009
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
BARR	50MG		A084771	001
DAVA PHARMS INC	100MG		A087060	001
ELKINS SINK	50MG		A085152	002
HEATHER	50MG		A084135	001
IMPAK LABS	25MG		A084029	001
	50MG		A083607	002
	100MG		A085098	001
INWOOD LABS	25MG		A084776	001
	25MG		A085067	001
	50MG		A084776	002
IVAX SUB TEVA PHARMS	50MG		A084658	001
	100MG		A085022	001

DISCONTINUED DRUG PRODUCT LIST

6 - 168 (of 346)

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

MAST MM	25MG	A086192	001	
	50MG	A086192	002	
MUTUAL PHARM	25MG	A083972	001	
	50MG	A083972	002	
	100MG	A083972	003	
MYLAN	25MG	A084880	001	
	50MG	A085112	001	
PVT FORM	50MG	A086597	001	
ROXANE	25MG	A085004	001	
	50MG	A084536	002	
	50MG	A085005	001	
SANDOZ	25MG	A083899	001	
	25MG	A087565	001	Mar 09, 1982
	50MG	A084912	001	
	50MG	A085219	001	
SOLVAY	25MG	A085323	001	
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
TG UNITED LABS	25MG	A085683	001	
	50MG	A083965	001	
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
	50MG	A087587	001	May 03, 1982
WATSON LABS	25MG	A083458	001	
	25MG	A085232	002	
	50MG	A083232	001	
	50MG	A083456	001	
	50MG	A085233	001	
	50MG	A086087	001	
	50MG	A086594	001	
	100MG	A081190	001	Jan 24, 1992
	100MG	A085099	001	
	100MG	A087002	001	
WEST WARD	25MG	A084899	001	
WHITEWORTH TOWN PLSN	25MG	A083809	002	
	50MG	A083809	001	
	100MG	A085347	001	
HYDRO-D				
HALSEY	25MG	A086504	001	
	50MG	A083891	002	
HYDRODIURIL				
MERCK	25MG	N011835	003	
	50MG	N011835	006	
	100MG	N011835	007	
ORETIC				
ABBOTT	25MG	N011971	001	
ZIDE				
SOLVAY	50MG	A083925	001	

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

SANOFI AVENTIS

12.5MG;75MG

N020758 001 Sep 30, 1997

25MG;300MG

N020758 004 Mar 15, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 169 (of 346)

HYDROCHLOROTHIAZIDE; LABETALOL HYDROCHLORIDE

TABLET; ORAL NORMOZIDE SCHERING	25MG;100MG 25MG;200MG 25MG;300MG 25MG;400MG	N019046 001 N019046 002 N019046 003 N019046 004	Apr 06, 1987 Apr 06, 1987 Apr 06, 1987 Apr 06, 1987
TRANDATE HCT GLAXOSMITHKLINE	25MG;100MG 25MG;200MG 25MG;300MG 25MG;400MG	N019174 001 N019174 002 N019174 003 N019174 004	Apr 10, 1987 Apr 10, 1987 Apr 10, 1987 Apr 10, 1987

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET; ORAL LISINOPRIL AND HYDROCHLOROTHIAZIDE SANDOZ	12.5MG;10MG 12.5MG;20MG 25MG;20MG	A075926 001 A075926 002 A075926 003	Jul 01, 2002 Jul 01, 2002 Jul 01, 2002
TEVA	12.5MG;10MG 12.5MG;20MG 25MG;20MG	A075869 001 A075869 002 A075869 003	Jul 01, 2002 Jul 01, 2002 Jul 01, 2002
PRINZIDE MERCK	25MG;20MG	N019778 002	Feb 16, 1989

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 CLONMEL HLTHCARE	15MG;250MG 25MG;250MG 30MG;500MG 50MG;500MG	A072507 001 A072508 001 A072509 001 A072510 001	Jun 02, 1989 Jun 02, 1989 Jun 02, 1989 Jun 02, 1989
IVAX SUB TEVA PHARMS	15MG;250MG 25MG;250MG 30MG;500MG 50MG;500MG	A071458 001 A071459 001 A071460 001 A071461 001	Mar 08, 1988 Mar 08, 1988 Mar 08, 1988 Mar 08, 1988
PAR PHARM	15MG;250MG 25MG;250MG 30MG;500MG 50MG;500MG	A070616 001 A070612 001 A070613 001 A070614 001	Feb 02, 1987 Feb 02, 1987 Feb 02, 1987 Feb 02, 1987
PARKE DAVIS	15MG;250MG 25MG;250MG 30MG;500MG 50MG;500MG	A071897 001 A071898 001 A071899 001 A071900 001	Nov 23, 1987 Nov 23, 1987 Nov 23, 1987 Nov 23, 1987
PUREPAC PHARM	15MG;250MG 25MG;250MG 30MG;500MG 50MG;500MG	A070853 001 A070688 001 A070854 001 A070689 001	Oct 08, 1986 Apr 24, 1986 Oct 08, 1986 Apr 24, 1986
SANDOZ	15MG;250MG 15MG;250MG 25MG;250MG 25MG;250MG 30MG;500MG 50MG;500MG	A070182 001 A070829 001 A070183 001 A070830 001 A070543 001 A070544 001	Jan 15, 1986 Mar 09, 1987 Jan 15, 1986 Mar 09, 1987 Jan 15, 1986 Jan 15, 1986
TEVA	15MG;250MG	A071819 001	Apr 08, 1988

DISCONTINUED DRUG PRODUCT LIST

6 - 170 (of 346)

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

TEVA	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
	25MG;250MG	A071921	001	Aug 29, 1988
	30MG;500MG	A070367	001	Mar 19, 1986
	30MG;500MG	A071069	001	Jan 19, 1989
	30MG;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 SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DUTOPROL

ASTRAZENECA	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

NOVARTIS	50MG;100MG	N018303	003	Dec 31, 1984
----------	------------	---------	-----	--------------

HYDROCHLOROTHIAZIDE; PINDOLOL

TABLET; ORAL

VISKAZIDE

NOVARTIS	25MG;5MG	N018872	001	Jul 22, 1987
	25MG;10MG	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-80/25

AKRIMAX PHARMS	25MG;80MG	N018031	002	
----------------	-----------	---------	-----	--

PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE

DURAMED PHARMS BARR	25MG;40MG	A071126	001	Mar 02, 1987
	25MG;80MG	A071127	001	Mar 02, 1987

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

ACTAVIS ELIZABETH	25MG;40MG	A070851	001	May 15, 1986
	25MG;80MG	A070852	001	May 15, 1986

BARR	25MG;40MG	A070704	001	Oct 01, 1986
	25MG;80MG	A070705	001	Oct 01, 1986

IVAX SUB TEVA PHARMS	25MG;40MG	A071552	001	Dec 01, 1988
	25MG;80MG	A071553	001	Dec 01, 1988

SANDOZ	25MG;40MG	A071060	001	Aug 26, 1987
	25MG;80MG	A071061	001	Aug 26, 1987

WARNER CHILCOTT	25MG;40MG	A071771	001	Jan 26, 1988
	25MG;80MG	A071772	001	Jan 26, 1988

WATSON LABS	25MG;40MG	A071498	001	Dec 18, 1991
-------------	-----------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 171 (of 346)

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL
 PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE
 WATSON LABS 25MG;80MG A071501 001 Dec 18, 1991

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL
 H.R.-50
 WHITEWORTH TOWN PLSN 50MG;0.125MG A085338 001
 HYDROCHLOROTHIAZIDE W/ RESERPINE
 IVAX SUB TEVA PHARMS 25MG;0.1MG A083572 001
 25MG;0.125MG A083571 001
 50MG;0.1MG A083568 001
 50MG;0.125MG A083573 001
 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
 HYDROPRES 25
 MERCK 25MG;0.125MG N011958 002
 HYDROPRES 50
 MERCK 50MG;0.125MG N011958 003
 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
 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
 MUTUAL PHARM 25MG;25MG A087267 001
 PUREPAC PHARM 25MG;25MG A087999 001 Nov 06, 1985
 SANDOZ 25MG;25MG A086881 001
 SUPERPHARM 25MG;25MG A089137 001 Aug 26, 1985
 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

DISCONTINUED DRUG PRODUCT LIST

6 - 172 (of 346)

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
 BARR 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

HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL
 CODAMINE
 ALPHARMA US PHARMS 5MG/5ML;25MG/5ML A075103 001 Sep 29, 2000
 HYCOMINE
 ENDO PHARMS 5MG/5ML;25MG/5ML N019410 001 Aug 17, 1990
 HYCOMINE PEDIATRIC
 ENDO PHARMS 2.5MG/5ML;12.5MG/5ML N019411 001 Aug 17, 1990

HYDROCORTAMATE HYDROCHLORIDE

OINTMENT; TOPICAL
 MAGNACORT
 PFIZER 0.5% N010554 001

HYDROCORTISONE

AEROSOL; TOPICAL
 AEROSEB-HC
 ALLERGAN HERBERT 0.5% A085805 001
 CREAM; TOPICAL
 CORT-DOME
 BAYER PHARMS 0.5% N009585 003
 1% N009585 001
 DERMACORT
 MONARCH PHARMS 1% A083011 002
 ELDECORT
 VALEANT PHARM INTL 1% A080459 001
 2.5% A084055 001
 FLEXICORT
 WESTWOOD SQUIBB 0.5% A087136 003 Apr 08, 1982
 1% A087136 002 Apr 08, 1982
 2.5% A087136 001 Apr 08, 1982
 HC #1
 BAYER PHARMS 0.5% A080438 001
 HC #4
 BAYER PHARMS 1% A080438 002
 HC (HYDROCORTISONE)
 C AND M PHARMA 0.5% A080482 003
 1% A080482 004
 H-CORT
 PHARM ASSOC 0.5% A086823 001
 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

DISCONTINUED DRUG PRODUCT LIST

6 - 173 (of 346)

HYDROCORTISONE

CREAM; TOPICAL				
HYDROCORTISONE				
ALTANA	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	
TEVA	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				
SANOFI AVENTIS US	1%	A080472	003	
	2.5%	A080472	004	
NOGENIC HC				
IVAX PHARMS	1%	A087427	001	Apr 04, 1988
NUTRACORT				
CORIA	0.5%	A080442	002	
	1%	A080442	003	
PENECLORT				
ALLERGAN HERBERT	1%	A088216	001	Jun 06, 1984
PROCTOCORT				
MONARCH PHARMS	1%	A083011	001	
SYNACORT				
MEDICIS	0.5%	A087459	001	
GEL; TOPICAL				
NUTRACORT				
HEALTHPOINT	1%	A084698	001	
PENECLORT				
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	
BALNEOL-HC				
SOLVAY	1%	A088041	001	Dec 03, 1982
BETA-HC				
BETA DERMAC	1%	A089495	001	Jan 25, 1988
CETACORT				
CORIA	0.5%	A080426	002	
	1%	A080426	001	
CORT-DOME				
BAYER PHARMS	0.5%	N009895	003	
	1%	N009895	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 174 (of 346)

HYDROCORTISONE

LOTION; TOPICAL					
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
HYTONE					
SANOFI AVENTIS US	1%		A080473	003	
	2.5%		A080473	004	Nov 30, 1982
NUTRACORT					
CORIA	0.5%		A080443	002	
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	
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	
PENECORT					
ALLERGAN HERBERT	2.5%		A088217	001	Jun 06, 1984
POWDER; FOR RX COMPOUNDING					
H-CORT					
TORCH	100%		A087834	001	Mar 29, 1982
HYDROCORTISONE					
PADDOCK LLC	100%		A088082	001	Apr 08, 1983
SOLUTION; TOPICAL					
PENECORT					
ALLERGAN HERBERT	1%		A088214	001	Jun 06, 1984
TEXACORT					
MISSION PHARMA	1%		A080425	001	
TABLET; ORAL					
CORTRIL					
PFIZER	10MG		N009127	005	
	20MG		N009127	003	

DISCONTINUED DRUG PRODUCT LIST

6 - 175 (of 346)

HYDROCORTISONE

TABLET; ORAL

HYDROCORTISONE

BARR	20MG	A083999	001	
ELKINS SINK	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	
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
PARKE DAVIS	1%	A089914	001	Jan 03, 1989
PUREPAC PHARM	0.5%	A086050	001	
	1%	A086052	001	

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
-------	------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 176 (of 346)

HYDROCORTISONE ACETATE

OINTMENT; TOPICAL CORTEF ACETATE	PHARMACIA AND UPJOHN 1% 2.5%	N008917 002 N008917 001
-------------------------------------	---------------------------------	----------------------------

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL NEO-CORTEF	PHARMACIA AND UPJOHN 1%;EQ 3.5MG BASE/GM 2.5%;EQ 3.5MG BASE/GM	A061049 001 A061049 002
OINTMENT; OPHTHALMIC NEO-CORTEF	PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/GM 1.5%;EQ 3.5MG BASE/GM	A060610 001 A060610 002
OINTMENT; TOPICAL NEO-CORTEF	PHARMACIA AND UPJOHN 0.5%;EQ 3.5MG BASE/GM 1%;EQ 3.5MG BASE/GM 2.5%;EQ 3.5MG BASE/GM	A060751 001 A060751 002 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 1.5%;EQ 3.5MG BASE/ML	A060612 002 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%	BOCA PHARMA 1%;1%	A089440 001 May 17, 1988
LOTION; TOPICAL PRAMOSONE	FERNDALE LABS 0.5%;1%	A083213 002

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
---------------------------------------	-----------------------	-------------

DISCONTINUED DRUG PRODUCT LIST

6 - 177 (of 346)

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	A085929	001	
	EQ 250MG BASE/VIAL	A085930	001	
	EQ 500MG BASE/VIAL	A085931	001	
	EQ 1GM BASE/VIAL	A085932	001	
<u>HYDROCORTISONE SODIUM SUCCINATE</u>				
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
	EQ 1GM BASE/VIAL	A088670	001	Jun 08, 1984
BAXTER HLTHCARE	EQ 100MG BASE/VIAL	A086619	001	
	EQ 250MG BASE/VIAL	A087567	001	
	EQ 500MG BASE/VIAL	A087568	001	
	EQ 1GM BASE/VIAL	A087569	001	
INTL MEDICATION	EQ 100MG BASE/VIAL	A087532	001	Mar 19, 1982
WATSON LABS	EQ 100MG BASE/VIAL	A084737	002	
	EQ 100MG BASE/VIAL	A084738	001	
	EQ 250MG BASE/VIAL	A084737	001	
	EQ 500MG BASE/VIAL	A084747	001	
	EQ 1GM BASE/VIAL	A084748	001	

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

TEVA PHARMS	0.2%	A074489	001	Aug 12, 1998
-------------	------	---------	-----	--------------

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 & 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

WATSON LABS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062521	001	Jul 11, 1985
-------------	-------------------------------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 178 (of 346)

HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC OTOBIOOTIC			
SCHERRING	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

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

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 HYDROMORPHONE HYDROCHLORIDE			
WATSON LABS	10MG/ML	A074317	001 Aug 23, 1995

HYDROXOCOBALAMIN

INJECTABLE; INJECTION ALPHAREDISOL			
MERCK	1MG/ML	A080778	001
CYANOKIT			
MERCK SANTE SAS	2.5GM/VIAL (5GM/KIT)	N022041	002 Dec 15, 2006
HYDROXOCOBALAMIN			
ABRAXIS PHARM	1MG/ML	A084921	001
WATSON LABS	1MG/ML	A085528	001

DISCONTINUED DRUG PRODUCT LIST

6 - 179 (of 346)

HYDROXOCOBALAMIN

INJECTABLE; INJECTION
 HYDROXOMIN
 BEL MAR 1MG/ML A084629 001

HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS; OPHTHALMIC
 PAREDRINE
 PHARMICS 1% N000004 004

HYDROXYPROGESTERONE CAPROATE

INJECTABLE; INJECTION
 DELALUTIN
 BRISTOL MYERS SQUIBB 125MG/ML N010347 004
 125MG/ML N016911 001
 250MG/ML N010347 002
 250MG/ML N016911 002
 HYDROXYPROGESTERONE CAPROATE
 AKORN 125MG/ML N018004 001
 WATSON LABS 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
 DURAMED PHARMS BARR 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
 APP PHARMS 25MG/ML A088184 001 Mar 31, 1983
 50MG/ML A088185 001 Mar 31, 1983
 BAXTER HLTHCARE 25MG/ML A085551 001
 HOSPIRA 25MG/ML A087416 001
 50MG/ML A086821 001
 50MG/ML A087546 001
 PHARMAFAIR 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

DISCONTINUED DRUG PRODUCT LIST

6 - 180 (of 346)

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

WATSON LABS	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	
SYRUP; ORAL				
ATARAX				
ROERIG	10MG/5ML	N010485	001	
HYDROXYZINE HYDROCHLORIDE				
ACTAVIS MID ATLANTIC	10MG/5ML	A086880	001	
ALPHARMA US PHARMS	10MG/5ML	A088785	001	Feb 03, 1988
KV PHARM	10MG/5ML	A087730	001	Jul 01, 1982
TABLET; ORAL				
ATARAX				
PFIZER	10MG	N010392	001	
	25MG	N010392	004	
	50MG	N010392	006	
	100MG	N010392	005	
HYDROXYZINE HYDROCHLORIDE				
ABLE	10MG	A040559	001	Jul 22, 2004
	25MG	A040562	001	Jul 22, 2004
	50MG	A040563	001	Jul 22, 2004
ACTAVIS TOTOWA	10MG	A040600	001	Dec 28, 2004
	10MG	A089071	001	Jul 22, 1986
	25MG	A040602	001	Dec 28, 2004
	25MG	A089072	001	Jul 22, 1986
	50MG	A040604	001	Dec 28, 2004
	50MG	A089073	001	Jul 22, 1986
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
	100MG	A087862	001	Apr 18, 1983
PLIVA	100MG	A081054	001	Sep 25, 1995
PUREPAC PHARM	10MG	A088120	001	Sep 25, 1984
	25MG	A088121	001	Sep 25, 1984
	50MG	A088122	001	Sep 25, 1984
QUANTUM PHARMICS	10MG	A088540	001	Oct 22, 1985
	25MG	A088551	001	Oct 22, 1985
	50MG	A088529	001	Oct 22, 1985
SANDOZ	10MG	A087246	002	
	10MG	A087869	001	Dec 20, 1982
	25MG	A085247	001	
	25MG	A087870	001	Dec 20, 1982
	50MG	A087245	001	
	50MG	A087871	001	Dec 20, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 181 (of 346)

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

SUPERPHARM	10MG	A088794	001	Dec 05, 1984
	25MG	A088795	001	Dec 05, 1984
	50MG	A088796	001	Dec 05, 1984
USL PHARMA	10MG	A089121	001	Mar 20, 1986
	25MG	A089122	001	Mar 20, 1986
	50MG	A089123	001	Mar 20, 1986
VINTAGE	10MG	A087602	001	Jan 22, 1982
	25MG	A087603	001	Jan 22, 1982
	50MG	A087604	001	Jan 22, 1982
WATSON LABS	10MG	A086827	001	
	25MG	A086829	001	
	50MG	A086836	001	

HYDROXYZINE PAMOATE

CAPSULE; ORAL

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 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
HY-PAM "25"				
TEVA	EQ 25MG HCL	A088713	001	Mar 04, 1985
VISTARIL				
PFIZER	EQ 100MG HCL	N011459	006	

IBANDRONATE SODIUM

TABLET; ORAL

BONIVA

ROCHE

EQ 2.5MG BASE

N021455 001 May 16, 2003

IBUPROFEN

CAPSULE; ORAL

MIDOL

BAYER

200MG

A070626 001 Sep 02, 1987

200MG

A071002 001 Sep 02, 1987

SUSPENSION; ORAL

CHILDREN'S ADVIL

WYETH CONS

100MG/5ML

N019833 002 Sep 19, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 182 (of 346)

IBUPROFEN

SUSPENSION; ORAL IBU					
ABBOTT	100MG/5ML		N019784	001	Dec 18, 1989
MOTRIN					
MCNEIL CONSUMER	100MG/5ML		N019842	001	Sep 19, 1989
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
IBUPRIN					
PLIVA	200MG		A071773	001	Jul 16, 1987
IBUPROFEN					
ABBOTT	600MG		A070556	001	Jun 14, 1985
	800MG		A071264	001	Jul 25, 1986
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		A071144	001	Jan 20, 1987
	200MG		A071154	001	Oct 27, 1987
	200MG		A072040	001	Apr 29, 1988
	200MG		A072901	001	Dec 19, 1991
	200MG		A072903	001	Dec 19, 1991
	400MG		A071145	001	Sep 23, 1986
	600MG		A071146	001	Sep 23, 1986
	800MG		A071769	001	May 08, 1987
LEDERLE	400MG		A070629	001	Sep 19, 1986
	600MG		A070630	001	Sep 19, 1986
LEINER	300MG		A071266	001	Oct 15, 1986
MCNEIL	400MG		A070081	001	Jun 16, 1986
	600MG		A070476	001	Jun 16, 1986
MUTUAL PHARM	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
MYLAN	200MG		A071870	001	May 05, 1988
	400MG		A070045	001	Sep 24, 1985
	600MG		A070057	001	Sep 24, 1985
	800MG		A071999	001	Dec 03, 1987
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

DISCONTINUED DRUG PRODUCT LIST

6 - 183 (of 346)

IBUPROFEN

TABLET; ORAL

IBUPROFEN

PUREPAC PHARM	300MG	A071123	001	Sep 19, 1986
	400MG	A071124	001	Sep 19, 1986
	600MG	A071125	001	Sep 19, 1986
	800MG	A071964	001	Feb 01, 1988
SANDOZ	200MG	A070733	001	Sep 19, 1986
	200MG	A071807	001	Feb 25, 1988
	200MG	A074525	001	Dec 15, 1995
	200MG	A074533	001	Dec 15, 1995
	300MG	A070734	001	Jun 12, 1986
	400MG	A070735	001	Jun 12, 1986
	400MG	A072064	001	Jan 14, 1988
	600MG	A070736	001	Jun 12, 1986
	600MG	A072065	001	Jan 14, 1988
	800MG	A071938	001	Jan 14, 1988
	800MG	A072169	001	Dec 11, 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
WATSON LABS	200MG	A071765	001	Sep 04, 1987
	200MG	A071905	001	Mar 08, 1988
	300MG	A071338	001	Dec 01, 1986
	400MG	A070038	001	Sep 06, 1985
	600MG	A070041	001	Sep 06, 1985
	800MG	A071547	001	Jul 02, 1987
	800MG	A071911	001	Oct 13, 1987
IBUPROHM				
OHM LABS	400MG	A070469	001	Aug 29, 1985
IBU-TAB				
ALRA	800MG	A071965	001	Aug 11, 1988
MEDIPREN				
MCNEIL	200MG	A070475	001	Feb 06, 1986
	200MG	A071215	001	Jun 26, 1986
MIDOL				
BAYER	200MG	A070591	001	Sep 02, 1987
	200MG	A071001	001	Sep 02, 1987
MOTRIN				
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
NUPRIN				
BRISTOL MYERS	200MG	A072035	001	Feb 16, 1988
	200MG	A072036	001	Feb 16, 1988
MCNEIL	200MG	N019012	001	May 18, 1984
	200MG	N019012	002	Jul 29, 1987
RUFEN				
BASF	600MG	N018197	002	Mar 05, 1984
TABLET, CHEWABLE; ORAL				
MOTRIN				
MCNEIL PED	50MG	N020135	001	Nov 16, 1994
	100MG	N020135	002	Nov 16, 1994

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

COMBUNOX

FOREST LABS

400MG;5MG

N021378 001 Nov 26, 2004

DISCONTINUED DRUG PRODUCT LIST

6 - 184 (of 346)

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN

PHARMACIA AND UPJOHN	5MG/VIAL
	10MG/VIAL
	20MG/VIAL

N050661	002	Sep 27, 1990
N050661	001	Sep 27, 1990
N050661	003	Apr 25, 1995

IDOXURIDINEOINTMENT; OPHTHALMIC
STOXIL

GLAXOSMITHKLINE 0.5%

N015868 001

SOLUTION/DROPS; OPHTHALMIC
STOXIL

GLAXOSMITHKLINE 0.1%

N013934 001

IFOSFAMIDE; MESNAINJECTABLE; INJECTION
IFEX/MESNEX KIT

BAXTER HLTHCARE	1GM/VIAL;100MG/ML
	3GM/VIAL;100MG/ML

N019763	003	Oct 10, 1992
N019763	004	Oct 10, 1992

ILOPROSTSOLUTION; INHALATION
VENTAVIS

ACTELION PHARMS LTD 20MCG/2ML (10MCG/ML)

N021779 001 Dec 29, 2004

IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

NOVARTIS	EQ 50MG BASE
	EQ 100MG BASE

N021335	001	May 10, 2001
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

ACTAVIS TOTOWA 10MG

A040753 001 Feb 28, 2008

25MG

A040752 001 Feb 28, 2008

50MG

A040751 001 Feb 28, 2008

LEDERLE 10MG

A086269 001

25MG

A086267 001

50MG

A086268 001

PAR PHARM 25MG

A089497 001 Jul 14, 1987

50MG

A088276 001 Oct 21, 1983

ROXANE 10MG

A083799 001

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

DISCONTINUED DRUG PRODUCT LIST

6 - 185 (of 346)

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

WATSON LABS	10MG 25MG 25MG 50MG 50MG	A085875 001 A084252 002 A085878 001 A085221 001 A085877 001
WEST WARD	25MG 50MG	A088222 001 May 26, 1983 A088223 001 May 26, 1983
JANIMINE		
ABBOTT	10MG 25MG 50MG	N017895 001 N017895 002 N017895 003
PRAMINE		
ALRA	10MG 25MG 50MG	A083827 001 A083827 002 A083827 003
PRESAMINE		
SANOFI AVENTIS US	10MG 25MG 50MG	N011836 006 N011836 003 N011836 007

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

CADISTA PHARMS	1.25MG 2.5MG	A075201 001 Dec 04, 1998 A075201 002 Dec 04, 1998
TEVA	1.25MG 1.25MG 2.5MG 2.5MG	A074498 002 Feb 12, 1998 A074665 001 Apr 04, 1997 A074498 001 Oct 31, 1996 A074665 002 Apr 04, 1997
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 333MG BASE	N020685 005 Dec 17, 1998
-------------------	---------------	--------------------------

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

AKORN	10MG/VIAL 40MG/VIAL 50MG/VIAL	N011525 003 N011525 004 N011525 002
-------	-------------------------------------	---

DISCONTINUED DRUG PRODUCT LIST

6 - 186 (of 346)

INDOMETHACIN

CAPSULE; ORAL

INDOCIN

IROKO PHARMS

25MG

N016059 001

50MG

N016059 002

INDO-LEMMON

TEVA

25MG

A070266 001

Nov 07, 1985

50MG

A070267 001

Nov 07, 1985

INDOMETHACIN

ABLE

25MG

A076666 001

Dec 17, 2003

50MG

A076666 002

Dec 17, 2003

DURAMED PHARMS BARR

25MG

A070326 001

Oct 18, 1985

50MG

A070327 001

Oct 18, 1985

HALSEY

25MG

A070782 001

Jun 03, 1987

50MG

A070635 001

Jun 03, 1987

IVAX SUB TEVA PHARMS

25MG

N018730 001

May 04, 1984

50MG

N018730 002

May 04, 1984

MUTUAL PHARM

25MG

A070067 001

Oct 03, 1986

25MG

A070899 001

Feb 09, 1987

50MG

A070068 001

Oct 03, 1986

50MG

A070900 001

Feb 09, 1987

MYLAN

50MG

N018858 002

Apr 20, 1984

PARKE DAVIS

25MG

N018806 001

Nov 23, 1984

50MG

N018806 002

Nov 23, 1984

PIONEER PHARMS

25MG

A070813 001

Aug 11, 1986

50MG

A070592 001

Aug 11, 1986

PLIVA

25MG

A071148 001

Mar 18, 1987

50MG

A071149 001

Mar 18, 1987

ROXANE

25MG

A070353 001

Jun 18, 1985

50MG

A070354 001

Jun 18, 1985

SUPERPHARM

25MG

A070487 001

Oct 10, 1986

50MG

A070488 001

Oct 10, 1986

TEVA

25MG

A071342 001

Apr 18, 1988

50MG

A071343 001

Apr 18, 1988

VINTAGE

25MG

N018829 002

Aug 06, 1984

50MG

A070651 001

Mar 05, 1986

WATSON LABS

25MG

N018829 001

Aug 06, 1984

25MG

A070529 001

Oct 18, 1985

25MG

A070784 001

Aug 20, 1986

25MG

A072996 001

Jul 31, 1991

50MG

N018690 001

Jul 31, 1984

50MG

A070530 001

Oct 18, 1985

50MG

A070785 001

Aug 20, 1986

50MG

A071635 001

May 18, 1987

50MG

A072997 001

Jul 31, 1991

50MG

N018690 002

Jul 31, 1984

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

75MG

N018185 001

Feb 23, 1982

INDOMETHACIN

75MG

A076114 001

Feb 06, 2002

ABLE

75MG

A072410 001

Mar 15, 1989

SUPPOSITORY; RECTAL

INDOCIN

50MG

N017814 001

Aug 13, 1984

SUSPENSION; ORAL

INDOMETHACIN

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

DISCONTINUED DRUG PRODUCT LIST

6 - 187 (of 346)

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
---	--------------	---------	-----

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 3MG/INH	N021868	001 002	Jan 27, 2006 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

DISCONTINUED DRUG PRODUCT LIST

6 - 188 (of 346)

INSULIN SUSP ISOPHANE BEEF

INJECTABLE; INJECTION

NPH INSULIN

NOVO NORDISK INC 40 UNITS/ML
 100 UNITS/MLN017929 001
N017929 003INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION

NPH ILETIN I (BEEF-PORK)

LILLY 40 UNITS/ML
 100 UNITS/MLN017936 001
N017936 002INSULIN 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
NPH ILETIN II (PORK)
LILLY 100 UNITS/ML
NPH PURIFIED PORK ISOPHANE INSULIN
NOVO NORDISK INC 100 UNITS/MLN018194 001
N018345 001
N018623 001INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

INSULATARD NPH HUMAN

NOVO NORDISK INC 100 UNITS/ML
NOVOLIN N
NOVO NORDISK INC 100 UNITS/MLN019449 001 May 30, 1986
N019065 001 Jan 23, 1985INSULIN SUSP PROTAMINE ZINC BEEF/PORK

INJECTABLE; INJECTION

PROTAMINE ZINC & ILETIN I (BEEF-PORK)
LILLY 40 UNITS/ML
 100 UNITS/MLN017932 001
N017932 002INSULIN SUSP PROTAMINE ZINC PURIFIED BEEF

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II
LILLY 100 UNITS/ML
PROTAMINE ZINC INSULIN
BRISTOL MYERS SQUIBB 40 UNITS/ML
 100 UNITS/MLN018476 001
N017928 001
N017928 003INSULIN 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/MLN017998 001
N017998 003INSULIN ZINC SUSP EXTENDED BEEF

INJECTABLE; INJECTION

ULTRALENTE INSULIN
NOVO NORDISK INC 100 UNITS/ML

N017997 003

DISCONTINUED DRUG PRODUCT LIST

6 - 189 (of 346)

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 N019571 001 Jun 10, 1987
 100 UNITS/ML 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

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

DISCONTINUED DRUG PRODUCT LIST

6 - 190 (of 346)

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

IODIPAMIDE SODIUM

INJECTABLE; INJECTION
 CHOLOGRAFIN SODIUM
 BRACCO 20% N009321 001

IODOHIPPURATE SODIUM I-123

INJECTABLE; INJECTION
 NEPHROFLOW
 GE HEALTHCARE 1mCi/ML N018289 001 Dec 28, 1984

IODOHIPPURATE SODIUM I-131

INJECTABLE; INJECTION
 HIPPURAN I 131
 MALLINCKRODT 0.25mCi/ML N016666 001
 HIPPUTOPE
 BRACCO 1-2mCi/VIAL N015419 002
 IODOHIPPURATE SODIUM I 131
 PHARMALUCENCE 0.2mCi/ML N017313 001

IODOXAMATE MEGLUMINE

INJECTABLE; INJECTION
 CHOLOVUE
 BRACCO 9.9% N018077 001
 40.3% N018076 001

IOFETAMINE HYDROCHLORIDE I-123

INJECTABLE; INJECTION
 SPECTAMINE
 IMP 1mCi/ML N019432 001 Dec 24, 1987

IOHEXOL

INJECTABLE; INJECTION
 OMNIPAQUE 210
 GE HEALTHCARE 45.3% N018956 006 Jun 30, 1989
 SOLUTION; URETHRAL
 OMNIPAQUE 70
 GE HEALTHCARE 15.1% N018956 007 Jun 01, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 191 (of 346)

IOPAMIDOL

INJECTABLE; 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
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				
ABBOTT	61%	A074638	001	Apr 30, 1997
COOK IMAGING	61%	A074881	003	Jul 28, 2000
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
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 ACID

TABLET; ORAL

TELEPAQUE

GE HEALTHCARE	500MG	N008032	001
---------------	-------	---------	-----

IOPHENIDYLATE

INJECTABLE; INJECTION

PANTOPAQUE

ALCON	100%	N005319	001
-------	------	---------	-----

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
--------------	-------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 192 (of 346)

IOTHALAMATE SODIUM I-125

INJECTABLE; INJECTION
 GLOFIL-125
 ISOTEX 250-300uCi/ML N017279 001

IOTROLAN

INJECTABLE; INTRATHECAL
 OSMOVIST 190 40.6% N019580 001 Dec 07, 1989
 BAYER HLTHCARE
 OSMOVIST 240 51.3% N019580 002 Dec 07, 1989
 BAYER HLTHCARE

IOVERSOL

INJECTABLE; INJECTION
 OPTIRAY 160 34% N019710 003 Dec 30, 1988
 MALLINCKRODT
 OPTIRAY 240 51% N020923 001 May 28, 1998
 MALLINCKRODT
 OPTIRAY 320 68% N020923 002 May 29, 1998
 MALLINCKRODT

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
 PHARMASCIENCE INC 0.02% A075507 001 Jan 19, 2001
 ROXANE 0.02% A075867 001 Jul 22, 2002

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

IRON DEXTRAN

INJECTABLE; INJECTION
 IRON DEXTRAN
 SANOFI AVENTIS US EQ 50MG IRON/ML N010787 002

IRON SUCROSE

INJECTABLE; INTRAVENOUS
 VENOFEER
 LUITPOLD EQ 75MG BASE/3.75ML (EQ 20MG BASE/ML) N021135 003 Mar 29, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 193 (of 346)

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
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			
MARSAM PHARMS LLC	99.9%	A074393	001 May 12, 1995

DISCONTINUED DRUG PRODUCT LIST

6 - 194 (of 346)

ISOFLUROPHATE

OINTMENT; OPHTHALMIC
FLOROPRYL
MERCK 0.025% N010656 001

ISONIAZID

INJECTABLE; INJECTION				
NYDRAZID				
SANDOZ	100MG/ML	N008662	001	
RIMIFON				
ROCHE	25MG/ML	N008420	002	
	100MG/ML	N008420	003	
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	
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	
MUTUAL PHARM	100MG	A080136	001	
	300MG	A083633	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
WATSON LABS	50MG	A080522	001	
	100MG	A080401	001	
	100MG	A080523	001	
	100MG	A085790	001	
	300MG	A083178	001	
	300MG	A085784	001	
WHITEWORTH TOWN PLSN	100MG	A080120	002	
LANIAZID				
LANNETT	50MG	A080140	001	
	100MG	A080140	002	
	300MG	A089776	001	Jun 13, 1988
NYDRAZID				
BRISTOL MYERS SQUIBB	100MG	N008392	003	

DISCONTINUED DRUG PRODUCT LIST

6 - 195 (of 346)

ISONIAZID

TABLET; ORAL STANOZIDE EVERYLIFE	100MG 300MG	A080126 001 A080126 002
--	----------------	----------------------------

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
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
DEY	0.5%	A086764 001 Jan 04, 1982
PARKE DAVIS	0.25%	A085994 001
	0.5%	A085540 001
ISUPREL SANOFI AVENTIS US	0.5% 1%	N006327 002 N006327 003
VAPO-ISO FISONS	0.5%	N016813 001
TABLET; RECTAL, SUBLINGUAL ISUPREL SANOFI AVENTIS US	10MG 15MG	N006328 001 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 ABBOTT	10% 25%	N006905 003 N006905 002

ISOSORBIDE

SOLUTION; ORAL ISMOTIC ALCON	100GM/220ML	N017063 001
------------------------------------	-------------	-------------

DISCONTINUED DRUG PRODUCT LIST

6 - 196 (of 346)

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL ISORDIL	40MG	N012882	002	Jul 29, 1988
WYETH AYERST		A086166	002	Sep 19, 1986
TABLET; ORAL ISOSORBIDE DINITRATE		A086169	001	Sep 19, 1986
MUTUAL PHARM	5MG	A086167	001	Sep 19, 1986
	10MG	A087564	001	Sep 18, 1986
	20MG	A089190	001	Feb 17, 1987
	30MG	A089191	001	Feb 17, 1987
SUPERPHARM	5MG	A089192	001	Feb 17, 1987
	10MG			
	20MG			
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				
MUTUAL PHARM	2.5MG	A084204	001	Sep 18, 1986
	5MG	A086168	001	Sep 18, 1986
	10MG	A087545	001	Sep 18, 1986
SANDOZ	2.5MG	A086225	001	Feb 19, 1988
	5MG	A086222	001	Feb 19, 1988
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 MONONITRATE

TABLET; ORAL ISMO	20MG	N019091	001	Dec 30, 1991
PROMIUS PHARMA				
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				
IVAX SUB TEVA PHARMS	30MG	A075448	002	Aug 07, 2001
	60MG	A075448	001	Jun 19, 2000
	120MG	A075448	003	Aug 07, 2001

ISOTRETINOIN

CAPSULE; ORAL ACCUTANE	10MG	N018662	002	May 07, 1982
HOFFMANN LA ROCHE	20MG	N018662	004	Mar 28, 1983
	40MG	N018662	003	May 07, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 197 (of 346)

ISRADIPINE

CAPSULE; ORAL DYNACIRC	SMITHKLINE BEECHAM	2.5MG 5MG	N019546 001 Dec 20, 1990
			N019546 002 Dec 20, 1990

ITRACONAZOLE

INJECTABLE; INJECTION SPORANOX	JANSSEN PHARMS	10MG/ML	N020966 001 Mar 30, 1999
-----------------------------------	----------------	---------	--------------------------

IVERMECTIN

TABLET; ORAL STROMECTOL	MERCK	6MG	N050742 001 Nov 22, 1996
----------------------------	-------	-----	--------------------------

KANAMYCIN SULFATE

CAPSULE; ORAL KANTREX	APOTHECON	EQ 500MG BASE EQ 500MG BASE EQ 500MG BASE	A060516 001 A061911 001 A062726 001 Mar 06, 1987
--------------------------	-----------	---	--

INJECTABLE; INJECTION KANAMYCIN	BAXTER HLTHCARE	EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A062324 001 A062324 002 A062324 003
------------------------------------	-----------------	--	---

KANAMYCIN SULFATE ABRAXIS PHARM		EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A062504 001 Apr 05, 1984 A062504 002 Apr 05, 1984 A062504 003 Apr 05, 1984
------------------------------------	--	--	--

INTL MEDICATION		EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A062466 001 Sep 30, 1983 A062466 002 Sep 30, 1983
-----------------	--	--------------------------------------	--

LOCH		EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A063021 001 Jul 31, 1992 A063022 001 Jul 31, 1992 A063025 001 Jul 31, 1992
------	--	--	--

PHARMAFAIR		EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A062668 001 May 07, 1987 A062672 001 May 07, 1987 A062669 001 May 07, 1987
------------	--	--	--

SOLOPAK		EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A062605 003 Feb 26, 1986 A062605 001 Feb 26, 1986 A062605 002 Feb 26, 1986
---------	--	--	--

WARNER CHILCOTT WATSON LABS		EQ 1GM BASE/3ML EQ 1GM BASE/3ML	A063092 001 Oct 11, 1989 A062520 003 May 09, 1985
--------------------------------	--	------------------------------------	--

KANTREX APOTHECON		EQ 75MG BASE/2ML EQ 75MG BASE/2ML EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 500MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML EQ 1GM BASE/3ML EQ 1GM BASE/3ML	A061655 003 A061901 003 A062564 001 Sep 21, 1984 A061655 001 A061901 001 A062564 002 Sep 21, 1984 A061655 002 A061901 002 A062564 003 Sep 21, 1984
----------------------	--	--	--

KLEBCIL KING PHARMS		EQ 75MG BASE/2ML EQ 500MG BASE/2ML EQ 1GM BASE/3ML	A062170 001 A062170 002 A062170 003
------------------------	--	--	---

KETOCONAZOLE

CREAM; TOPICAL NIZORAL	JANSSEN PHARMA	2%	N019084 001 Dec 31, 1985
---------------------------	----------------	----	--------------------------

DISCONTINUED DRUG PRODUCT LIST

6 - 198 (of 346)

KETOCONAZOLE

SUSPENSION; ORAL NIZORAL	JANSEN PHARMA	100MG/5ML	A070767	001	Nov 07, 1986
TABLET; ORAL KETOCONAZOLE	AAIPHARMA LLC	200MG	A075341	001	Jul 27, 1999
	TEVA	200MG	A074971	001	Jun 15, 1999

KETOPROFEN

CAPSULE; ORAL KETOPROFEN	SANDOZ	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 ORUVAIL	WYETH PHARMS INC	100MG	N019816	003	Feb 08, 1995
		150MG	N019816	002	Feb 08, 1995
		200MG	N019816	001	Sep 24, 1993
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	AMPHASTAR PHARM	15MG/ML	A076209	001	Jul 21, 2004
		30MG/ML	A076209	002	Jul 21, 2004
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
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
TORADOL					
ROCHE PALO	15MG/ML		N019698	001	Nov 30, 1989
	30MG/ML		N019698	002	Nov 30, 1989
TABLET; ORAL KETOROLAC TROMETHAMINE					
ROXANE	10MG		A074790	001	Jun 26, 1997
WATSON LABS	10MG		A074955	001	Sep 19, 1997
TORADOL					
ROCHE PALO	10MG		N019645	001	Dec 20, 1991

KRYPTON, KR-81M

GAS; INHALATION MPI KRYPTON 81M GENERATOR	GE HEALTHCARE	N/A	N018088	001	
--	---------------	-----	---------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 199 (of 346)

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION					
LABETALOL HYDROCHLORIDE					
APOTHECON	5MG/ML		A075355	001	Nov 29, 1999
HOSPIRA	5MG/ML		A075242	001	Sep 30, 1999
NORMODYNE					
SCHERING	5MG/ML		N018686	001	Aug 01, 1984
TRANDATE					
PROMETHEUS LABS	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
MUTUAL PHARM	100MG		A075215	001	Jul 29, 1999
	200MG		A075215	002	Jul 29, 1999
	300MG		A075215	003	Jul 29, 1999
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					
PROMETHEUS LABS	300MG		N018716	003	Aug 01, 1984
	400MG		N018716	004	Aug 01, 1984

LACTULOSE

SOLUTION; ORAL					
CHRONULAC					
SANOFI AVENTIS US	10GM/15ML		N017884	001	
DUPHALAC					
SOLVAY	10GM/15ML		A072372	001	Mar 22, 1989
EVALOSE					
TEVA PHARMS	10GM/15ML		A073497	001	May 28, 1993
LACTULOSE					
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
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					
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

LAMOTRIGINE

TABLET; ORAL					
LAMICTAL					
GLAXOSMITHKLINE	50MG		N020241	006	Dec 27, 1994
	250MG		N020241	004	Dec 27, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 200 (of 346)

LAMOTRIGINE

TABLET; ORAL					
LAMOTRIGINE					
MATRIX 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
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
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
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	100MG		N020764	003	Aug 24, 1998
LAMOTRIGINE					
SANDOZ	5MG		A078409	002	Jan 22, 2009
	25MG		A078409	003	Jan 22, 2009

LANSOPRAZOLE

FOR SUSPENSION, DELAYED RELEASE; ORAL					
PREVACID					
TAKEDA PHARMS NA	15MG/PACKET		N021281	001	May 03, 2001
	30MG/PACKET		N021281	002	May 03, 2001
INJECTABLE; INTRAVENOUS					
PREVACID IV					
TAKEDA PHARMS NA	30MG/VIAL		N021566	001	May 27, 2004

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	EQ 250MG BASE		N021468	001	Oct 26, 2004

LAPYRIUM CHLORIDE; UNDECYLUM CHLORIDE IODINE COMPLEX

SOLUTION; TOPICAL					
VIRAC REX					
CHESEBROUGH PONDS	0.5%;1.8%		N011914	001	

LEFLUNOMIDE

TABLET; ORAL					
LEFLUNOMIDE					
SANDOZ	10MG		A077085	001	Sep 13, 2005
	20MG		A077085	002	Sep 13, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 201 (of 346)

LETROZOLE

TABLET; ORAL
 LETROZOLE
 SYNTHON PHARMS 2.5MG 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	A089352 001 A089353 001	Jun 01, 1988 Jun 01, 1988
ABRAXIS PHARM ELKINS SINK	EQ 50MG BASE/VIAL EQ 50MG BASE/VIAL EQ 100MG BASE/VIAL	A088939 001 A070480 001 A081224 001	Dec 01, 1986 Jan 02, 1987 Jun 03, 1994
HOSPIRA	EQ 3MG BASE/ML EQ 50MG BASE/VIAL EQ 100MG BASE/VIAL EQ 350MG BASE/VIAL	N008107 001 N008107 002 N008107 004 N008107 005	May 23, 1988 Apr 05, 1989
PHARMACHEMIE PHARMACHEMIE USA	EQ 350MG BASE/VIAL EQ 50MG BASE/VIAL EQ 100MG BASE/VIAL	A040262 001 A089628 001 A089915 001	Dec 15, 1999 Apr 17, 1997 Apr 17, 1997
LEUCOVORIN CALCIUM PRESERVATIVE FREE HOSPIRA	EQ 10MG BASE/ML	A040147 001	Jun 25, 1997
TEVA PARENTERAL	EQ 10MG BASE/ML	A040332 001	Jun 28, 1999
WELLCOVORIN GLAXOSMITHKLINE	EQ 5MG BASE/ML EQ 25MG BASE/VIAL EQ 50MG BASE/VIAL EQ 100MG BASE/VIAL	A087439 001 A089833 001 A089465 001 A089834 001	Oct 19, 1982 Jan 23, 1989 Jan 23, 1989 Jan 23, 1989
TABLET; ORAL LEUCOVORIN CALCIUM COREPHARMA	EQ 5MG BASE EQ 25MG BASE	A074544 001 A074544 002	Aug 28, 1997 Aug 28, 1997
PAR PHARM	EQ 5MG BASE EQ 25MG BASE	A071600 001 A071598 001	Oct 14, 1987 Oct 14, 1987
PHARMACHEMIE	EQ 5MG BASE EQ 25MG BASE	A073099 001 A073101 001	Mar 28, 1997 Mar 28, 1997
SANDOZ	EQ 15MG BASE	A075327 001	Mar 24, 1999
XANODYNE PHARM	EQ 5MG BASE EQ 10MG BASE EQ 15MG BASE	N018459 001 A071962 001 A071104 001	Jan 30, 1986 Nov 19, 1987 Mar 04, 1987
WELLCOVORIN GLAXOSMITHKLINE	EQ 5MG BASE EQ 25MG BASE	N018342 001 N018342 002	Jul 08, 1983 Jul 08, 1983

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION VIADUR	EQ 65MG BASE	N021088 001	Mar 03, 2000
ORTHO MCNEIL JANSSEN			
INJECTABLE; INJECTION LEUPROLIDE ACETATE GENZYME	1MG/0.2ML	A075721 001	Nov 29, 2001
LUPRON			
ABBOTT LABS	1MG/0.2ML	N019010 001	Apr 09, 1985
LUPRON DEPOT-PED			
ABBOTT ENDOCRINE	3.75MG/VIAL, 7.5MG/VIAL 7.5MG/VIAL, 7.5MG/VIAL	N020263 003 N020263 004	Apr 16, 1993 Apr 16, 1993

DISCONTINUED DRUG PRODUCT LIST

6 - 202 (of 346)

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

TABLET; ORAL
 LEVETIRACETAM
 MYLAN 250MG A078731 001 Feb 10, 2009
 500MG A078731 002 Feb 10, 2009
 750MG A078731 003 Feb 10, 2009
 1GM A078731 004 Feb 10, 2009
 WATSON LABS FLORIDA 250MG A077408 001 Mar 02, 2009
 500MG A077408 002 Mar 02, 2009
 750MG A077408 003 Mar 02, 2009

LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC
 BETAXON
 ALCON PHARMS LTD EQ 0.5% BASE N021114 001 Feb 23, 2000

LEVOBUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION
 CHIROCAINE
 PURDUE PHARMA LP EQ 2.5MG BASE/ML N020997 001 Aug 05, 1999
 EQ 5MG BASE/ML N020997 002 Aug 05, 1999
 EQ 7.5MG BASE/ML N020997 003 Aug 05, 1999

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC
 LIVOSTIN
 NOVARTIS EQ 0.05% BASE N020219 001 Nov 10, 1993

LEVOCARNITINE

SOLUTION; ORAL
 CARNITOR
 SIGMA TAU 1GM/10ML N018948 002 Apr 27, 1988

LEVODOPA

CAPSULE; ORAL
 BENDOPA
 VALEANT PHARM INTL 100MG N016948 003
 250MG N016948 001
 500MG N016948 002

DOPAR
 SHIRE 100MG N016913 003
 250MG N016913 001
 500MG N016913 002

LARODOPA
 ROCHE 100MG N016912 002
 250MG N016912 001
 500MG N016912 006

TABLET; ORAL
 DOPAR
 SHIRE 250MG N016913 004
 500MG N016913 005

LARODOPA
 ROCHE 100MG N016912 005

DISCONTINUED DRUG PRODUCT LIST

6 - 203 (of 346)

LEVODOPA

TABLET; ORAL LARODOPA ROCHE	250MG 500MG	N016912 003 N016912 004
-----------------------------------	----------------	----------------------------

LEVOMEPRAMAZINE

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-COBEEFRIN EASTMAN KODAK 0.05MG/ML;2%	N012125 002
MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFRIN GRAHAM CHEM 0.05MG/ML;2%	A084850 002 Oct 21, 1983
POLOCAINE W/ LEVONORDEFRIN DENTSPLY PHARM 0.05MG/ML;2%	A089517 001 Apr 14, 1988

LEVONORDEFRIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION RAVOCAIN AND NOVOCAIN W/ NEO-COBEEFRIN EASTMAN KODAK 0.05MG/ML;2%;0.4%	N008592 007
--	-------------

LEVONORGESTREL

IMPLANT; IMPLANTATION LEVONORGESTREL WYETH PHARMS INC 75MG/IMPLANT	N020627 001 Aug 15, 1996
NORPLANT POPULATION COUNCIL 36MG/IMPLANT	N019897 001 Dec 10, 1990
NORPLANT II POPULATION COUNCIL 75MG/IMPLANT	N020544 001 Nov 01, 1996
NORPLANT SYSTEM IN PLASTIC CONTAINER WYETH PHARMS INC 36MG/IMPLANT	N020088 001 Dec 10, 1990
TABLET; ORAL PLAN B TEVA WOMENS 0.75MG	N021045 001 Jul 28, 1999

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

DISCONTINUED DRUG PRODUCT LIST

6 - 204 (of 346)

LEVOOTHYROXINE SODIUM**

TABLET; ORAL					
LEVOLET					
VINTAGE	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	
LEVOXYL					
--> KING PHARMS	--> 0.3MG	N021301	012	May 25, 2001	
NOVOTHYROX					
MERCK KGAA	0.025MG	N021292	001	May 31, 2002	
	0.05MG	N021292	002	May 31, 2002	
	0.075MG	N021292	003	May 31, 2002	
	0.088MG	N021292	004	May 31, 2002	
	0.1MG	N021292	005	May 31, 2002	
	0.112MG	N021292	006	May 31, 2002	
	0.125MG	N021292	007	May 31, 2002	
	0.137MG	N021292	008	May 31, 2002	
	0.15MG	N021292	009	May 31, 2002	
	0.175MG	N021292	010	May 31, 2002	
	0.2MG	N021292	011	May 31, 2002	
	0.3MG	N021292	012	May 31, 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					
GRAHAM CHEM	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		

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION					
ALPHACAIN HYDROCHLORIDE					
CARLISLE	2%	A084721	001		
LIDOCAINE HYDROCHLORIDE					
ABBOTT	10%	A087980	001	Feb 02, 1983	
	20%	A089362	001	May 25, 1988	

DISCONTINUED DRUG PRODUCT LIST

6 - 205 (of 346)

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE

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	
BAXTER HLTHCARE	1%	A080407	001	
	2%	A080407	002	
BEL MAR	1%	A080710	001	
	2%	A080760	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	
GRAHAM CHEM	2%	A080504	001	
HOSPIRA	1%	A040013	001	Jun 23, 1995
	1.5%	A088330	001	May 17, 1984
	2%	A088331	001	May 17, 1984
INTL MEDICATION	1%	A083173	001	
	1%	N017701	002	
	2%	A083173	002	
	2%	N017701	001	
	1GM/VIAL	N018543	001	
	2GM/VIAL	N018543	002	
LYPHOMED	1%	A080390	001	
	2%	A080390	002	
MILES	1%	A080414	001	
	2%	A080414	002	
WATSON LABS	1%	A080377	001	
	1%	A083627	001	
	2%	A080377	002	
	2%	A083627	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	0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	800MG/100ML	N018967	003	Mar 30, 1984
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

DISCONTINUED DRUG PRODUCT LIST

6 - 206 (of 346)

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION				
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE				
BAXTER HLTHCARE	1%	A084625	001	
	2%	A084625	002	
INTL MEDICATION	4%	N017702	002	
LIDOCATON				
PHARMATON	2%	A084727	001	Aug 17, 1983
XYLOCAINE				
ASTRAZENECA	1%	N010418	005	
	1.5%	N010418	009	
	2%	N010418	007	
INJECTABLE; SPINAL				
XYLOCAINE 1.5% W/ DEXTROSE 7.5%				
APP PHARMS	1.5%	N016297	001	
XYLOCAINE 5% W/ GLUCOSE 7.5%				
ASTRAZENECA	5%	N010496	002	Jul 07, 1982
SOLUTION; ORAL				
LIDOCAINE HYDROCHLORIDE VISCOUS				
ACTAVIS MID ATLANTIC	2%	A086578	001	
INTL MEDICATION	2%	A086389	001	Feb 02, 1982
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	
SYSTEM; INTRADERMAL				
ZINGO				
POWDER PHARMS	0.5MG	N022114	001	Aug 16, 2007

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	
INJECTABLE; INJECTION				
LINCOMYCIN HYDROCHLORIDE				
WATSON LABS	EQ 300MG BASE/ML	A063180	001	Apr 16, 1991

LINDANE

CREAM; TOPICAL				
KWELL				
REED AND CARNICK	1%	A084218	001	
	1%	N006309	001	
LOTION; TOPICAL				
GAMENE				
SOLA BARNES HIND	1%	A084989	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 207 (of 346)

LINDANE

LOTION; TOPICAL KWELL				
REED AND CARNICK	1%	A084218	002	
	1%	N006309	003	
SCABENE STIEFEL	1%	A086769	001	
SHAMPOO; TOPICAL GAMENE				
SOLA BARNES HIND	1%	A084988	001	
KWELL				
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				
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
PRINIVIL				
MERCK	2.5MG	N019558	006	Jan 28, 1994

LITHIUM CARBONATE

CAPSULE; ORAL ESKALITH				
NOVEN THERAP	300MG	N016860	001	
LITHIUM CARBONATE				
ABLE	150MG	A076823	001	Jun 29, 2004
	300MG	A076121	001	Sep 27, 2001
	300MG	A076823	002	Jun 29, 2004
	600MG	A076823	003	Jun 29, 2004
USL PHARMA	300MG	A072542	001	Feb 01, 1989
WATSON LABS	300MG	A070407	001	Mar 19, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 208 (of 346)

LITHIUM CARBONATE

CAPSULE; ORAL LITHONATE SOLVAY	300MG	N016782	001
TABLET; ORAL ESKALITH JDS PHARMS	300MG	N017971	001
LITHANE BAYER PHARMS	300MG	N018833	001 Jul 18, 1985
LITHIUM CARBONATE PFIZER	300MG	N016834	001
LITHOTABS SOLVAY	300MG	N016980	001
TABLET, EXTENDED RELEASE; ORAL ESKALITH CR JDS PHARMS	450MG	N018152	001 Mar 29, 1982
LITHIUM CARBONATE ABLE	300MG	A076382	001 Apr 21, 2003
BARR	300MG	A076170	001 Jun 10, 2002
	450MG	A076366	001 Aug 21, 2003

LITHIUM CITRATE

SYRUP; ORAL LITHONATE SOLVAY	EQ 300MG CARBONATE/5ML	N017672	001
------------------------------------	------------------------	---------	-----

LOMEFLOXACIN HYDROCHLORIDE

TABLET; ORAL MAXAQUIN PHARMACIA	EQ 400MG BASE	N020013	001 Feb 21, 1992
---------------------------------------	---------------	---------	------------------

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL IMODIUM MCNEIL PED	2MG	N017690	001
LOPERAMIDE HYDROCHLORIDE ROXANE	2MG	A073080	001 Nov 27, 1991
SANDOZ	2MG	A072993	001 Aug 28, 1992
TEVA	2MG	A073122	001 Aug 30, 1991
SOLUTION; ORAL IMODIUM JANSSEN PHARMA	1MG/5ML	N019037	001 Jul 31, 1984
LOPERAMIDE HYDROCHLORIDE ALPHARMA US PHARMS	1MG/5ML	A073187	001 Sep 15, 1992
DURAMED PHARMS BARR	1MG/5ML	A074991	001 Dec 29, 1997
TEVA	1MG/5ML	A073478	001 Jun 23, 1995
WATSON LABS	1MG/5ML	A073062	001 May 28, 1993
TABLET; ORAL LOPERAMIDE HYDROCHLORIDE ABLE	2MG	A073528	001 Nov 30, 1993
PERRIGO	2MG	A074194	001 Oct 30, 1992

LORACARB

CAPSULE; ORAL LORABID KING PHARMS	200MG	N050668	001 Dec 31, 1991
	400MG	N050668	002 Apr 05, 1996
FOR SUSPENSION; ORAL LORABID KING PHARMS	100MG/5ML	N050667	001 Dec 31, 1991
	200MG/5ML	N050667	002 Dec 31, 1991

DISCONTINUED DRUG PRODUCT LIST

6 - 209 (of 346)

LORATADINE

SYRUP; ORAL CLARITIN HIVES RELIEF	SCHERRING PLOUGH	1MG/ML	N020641	003	Nov 19, 2003
TABLET; ORAL LORATADINE PERRIGO		10MG	N021512	001	Jun 24, 2004

LORAZEPAM

INJECTABLE; INJECTION LORAZEPAM	AKORN	2MG/ML	A074974	001	Jul 23, 1998
	BAXTER HLTHCARE	2MG/ML	A074496	001	Sep 28, 1998
		4MG/ML	A074496	002	Sep 28, 1998
	DAVA PHARMS INC	2MG/ML	A074793	001	Mar 16, 2000
		4MG/ML	A074793	002	Mar 16, 2000
	HOSPIRA	2MG/ML	A074280	001	May 27, 1994
		4MG/ML	A074280	002	May 27, 1994
		4MG/ML	A074300	003	Mar 19, 1997
	MARSAM PHARMS LLC	1MG/0.5ML	A074551	003	Sep 12, 1996
		2MG/ML	A074535	001	Sep 12, 1996
		2MG/ML	A074551	001	Sep 12, 1996
		4MG/ML	A074535	002	Sep 12, 1996
		4MG/ML	A074551	002	Sep 12, 1996
SOLUTION; ORAL LORAZEPAM	Roxane	0.5MG/5ML	A074648	001	Mar 18, 1997
TABLET; ORAL LORAZ	QUANTUM PHARMICS	0.5MG	A070200	001	Aug 09, 1985
		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
	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
		0.5MG	A072553	001	Mar 29, 1991
		1MG	A070473	001	Dec 10, 1985
		1MG	A072554	001	Mar 29, 1991
		2MG	A070474	001	Dec 10, 1985
		2MG	A072555	001	Mar 29, 1991
	PAR PHARM	0.5MG	A070675	001	Dec 01, 1986
		1MG	A070676	001	Dec 01, 1986
		2MG	A070677	001	Dec 01, 1986
	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
		2MG	A071088	001	Mar 23, 1987
		2MG	A071110	001	Jul 24, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 210 (of 346)

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS; OPHTHALMIC
 LOTEMAX
 PHARMOS 0.5% N020841 001 Mar 09, 1998

LOVASTATIN

TABLET; ORAL
 MEVACOR
 MERCK 10MG N019643 002 Mar 28, 1991
 TABLET, EXTENDED RELEASE; ORAL
 ALTOPREV
 ANDRX LABS LLC 10MG N021316 001 Jun 26, 2002

LOXAPINE HYDROCHLORIDE

CONCENTRATE; ORAL
 LOXITANE C
 WATSON LABS EQ 25MG BASE/ML N017658 001
 INJECTABLE; INJECTION
 LOXITANE IM
 WATSON LABS EQ 50MG BASE/ML N018039 001

LOXAPINE SUCCINATE

CAPSULE; ORAL
 LOXAPINE SUCCINATE
 ACTAVIS TOTOWA EQ 5MG BASE A076868 001 Aug 04, 2005
 EQ 10MG BASE A076868 002 Aug 04, 2005
 EQ 25MG BASE A076868 003 Aug 04, 2005
 EQ 50MG BASE A076868 004 Aug 04, 2005
 LOXITANE
 WATSON LABS EQ 5MG BASE N017525 001
 EQ 10MG BASE N017525 002
 EQ 25MG BASE N017525 003
 EQ 50MG BASE N017525 004
 TABLET; ORAL
 LOXITANE
 WATSON LABS EQ 10MG BASE N017525 006
 EQ 25MG BASE N017525 007
 EQ 50MG BASE N017525 008

LYPRESSIN

SOLUTION; NASAL
 DIAPID
 NOVARTIS 0.185MG/ML N016755 001

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 N019006 001 Apr 04, 1984
 G/100ML;530MG/100ML;500MG/100ML;12MG/10
 OML

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 N018252 001
 /100ML;500MG/100ML

SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER
 HOSPIRA INC 14MG/100ML;37MG/100ML;222MG/100ML;526MG N018406 001
 /100ML;502MG/100ML

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA INC 30MG/100ML;37MG/100ML;222MG/100ML;526MG N018406 002 Jul 08, 1982
 /100ML;502MG/100ML

DISCONTINUED DRUG PRODUCT LIST

6 - 211 (of 346)

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

SOLUTION; IRRIGATION

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, CHEWABLE; ORAL

ZEGERID

SANTARUS	700MG;20MG;600MG 700MG;40MG;600MG	N021850	001	Mar 24, 2006
		N021850	002	Mar 24, 2006

MANGAFODIPIR TRISODIUM

INJECTABLE; INJECTION

TESLASCAN

GE HEALTHCARE	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%

HOSPIRA	10GM/100ML	N016269	002
---------	------------	---------	-----

MILES	10GM/100ML	N016472	002
-------	------------	---------	-----

MANNITOL 15%

HOSPIRA	15GM/100ML	N016269	003
---------	------------	---------	-----

MILES	15GM/100ML	N016472	005
-------	------------	---------	-----

MANNITOL 20%

HOSPIRA	20GM/100ML	N016269	004
---------	------------	---------	-----

MILES	20GM/100ML	N016472	004
-------	------------	---------	-----

MANNITOL 25%

ABRAXIS PHARM	12.5GM/50ML	A086754	001
---------------	-------------	---------	-----

ASTRAZENECA	12.5GM/50ML	A089239	001
-------------	-------------	---------	-----

	12.5GM/50ML	A089240	001
--	-------------	---------	-----

HOSPIRA	12.5GM/50ML	N016269	005
---------	-------------	---------	-----

MERCK	12.5GM/50ML	N005620	001
-------	-------------	---------	-----

WATSON LABS	12.5GM/50ML	A087460	001
-------------	-------------	---------	-----

		Jun 27, 1983
--	--	--------------

MANNITOL 5%

HOSPIRA	5GM/100ML	N016269	001
---------	-----------	---------	-----

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
----------	------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 212 (of 346)

MAPROTILINE HYDROCHLORIDE

TABLET; ORAL LUDIOMIL NOVARTIS	50MG 75MG	N017543 002 N017543 003	Sep 30, 1982
<u>MAPROTILINE HYDROCHLORIDE</u>			
AM THERAP	25MG 50MG 75MG	A072129 001 A072130 001 A072131 001	Jan 14, 1988 Jan 14, 1988 Jan 14, 1988
WATSON LABS	25MG 25MG 50MG 50MG 75MG 75MG	A071943 001 A072162 001 A071944 001 A072163 001 A071945 001 A072164 001	Dec 30, 1987 Jun 01, 1988 Dec 30, 1987 Jun 01, 1988 Dec 30, 1987 Jun 01, 1988

MASOPROCOL

CREAM; TOPICAL ACTINEX UNIV AZ CANCER CTR	10%	N019940 001	Sep 04, 1992
---	-----	-------------	--------------

MAZINDOL

TABLET; ORAL MAZANOR WYETH AYERST	1MG 2MG	N017980 002 N017980 001	
<u>SANOREX</u>			
NOVARTIS	1MG 2MG	N017247 001 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 MECLIZINE HYDROCHLORIDE ABC HOLDING	12.5MG 25MG	A085253 001 A085252 001	
<u>ANABOLIC</u>			
BUNDY	12.5MG 25MG	A084382 001 A084872 001	
IVAX SUB TEVA PHARMS	12.5MG 12.5MG 25MG	A083784 001 A084975 001 A084657 001	
KV PHARM	12.5MG 25MG	A085524 001 A085523 001	

DISCONTINUED DRUG PRODUCT LIST

6 - 213 (of 346)

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

PLIVA	12.5MG	A088732	001	Dec 11, 1985
	25MG	A088734	001	Dec 11, 1985
SUPERPHARM	12.5MG	A089113	001	Aug 20, 1985
	25MG	A089114	001	Aug 20, 1985
UDL	12.5MG	A088256	001	Jun 13, 1983
	25MG	A088257	001	Jun 13, 1983
VANGARD	12.5MG	A087877	001	Apr 20, 1982
	25MG	A087620	001	Jan 04, 1982
WATSON LABS	12.5MG	A085195	001	
	12.5MG	A085269	001	
TABLET, CHEWABLE; ORAL				
ANTIVERT				
PFIZER	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

A071640 001 Aug 11, 1987

EQ 100MG BASE

A070401 001 Nov 25, 1986

EQ 100MG BASE

A071641 001 Aug 11, 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

TABLET; ORAL

AMEN

AMARIN PHARMS

10MG

A083242 001

CURRETAB

SOLVAY

10MG

A085686 001

CYCRIN

ESI

2.5MG

A081239 001 Oct 30, 1992

DISCONTINUED DRUG PRODUCT LIST

6 - 214 (of 346)

MEDROXYPROGESTERONE ACETATE

TABLET; ORAL CYCRIN					
ESI	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	

MEDRYSONE

SUSPENSION; OPHTHALMIC HMS					
ALLERGAN	1%	N016624	003		

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL LARIAM					
ROCHE	250MG	N019591	001	May 02, 1989	
MEFLOQUINE HYDROCHLORIDE					
US ARMY WALTER REED	250MG	N019578	001	May 02, 1989	

MEGESTROL ACETATE

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

TABLET; ORAL MELOXICAM					
MUTUAL PHARM	7.5MG	A077935	001	Jul 19, 2006	
	15MG	A077935	002	Jul 19, 2006	
ROXANE	7.5MG	A077925	001	Jul 19, 2006	
	15MG	A077925	002	Jul 19, 2006	
YABAO BIOPHARMS	7.5MG	A077933	001	Jul 19, 2006	
	15MG	A077933	002	Jul 19, 2006	

MEMANTINE HYDROCHLORIDE

TABLET; ORAL MEMANTINE					
SUN PHARMA GLOBAL	5MG	A090058	001	May 05, 2010	
	10MG	A090058	002	May 05, 2010	
MEMANTINE HYDROCHLORIDE					
DR REDDYS LABS LTD	5MG	A090048	001	Apr 14, 2010	
	10MG	A090048	002	Apr 14, 2010	
TEVA PHARMS	5MG	A090052	001	Oct 25, 2011	
	10MG	A090052	002	Oct 25, 2011	

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		

DISCONTINUED DRUG PRODUCT LIST

6 - 215 (of 346)

MENADIOL SODIUM DIPHOSPHATE

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 150 IU/VIAL;150 IU/VIAL	N020328 001 Sep 01, 1994 N020328 002 Sep 01, 1994
MENOTROPINS FERRING	75 IU/VIAL;75 IU/VIAL 150 IU/VIAL;150 IU/VIAL	A073598 001 Jan 30, 1997 A073599 001 Jan 30, 1997
PERGONAL SERONO	75 IU/AMP;75 IU/AMP 150 IU/AMP;150 IU/AMP	N017646 001 N017646 002 May 20, 1985
REPRONEX FERRING	150 IU/VIAL;150 IU/VIAL	N021047 002 Aug 27, 1999

MEPENZOLATE BROMIDE

SOLUTION; ORAL CANTIL SANOFI AVENTIS US	25MG/5ML	N010679 004
---	----------	-------------

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION DEMEROL SANOFI AVENTIS US	25MG/ML 50MG/ML 75MG/ML 100MG/ML	N005010 007 N005010 002 N005010 009 N005010 003
MEPERIDINE HYDROCHLORIDE ABBOTT	25MG/ML 50MG/ML 50MG/ML 75MG/ML 100MG/ML	A080388 001 A080385 001 A080387 001 A080389 001 A080386 001
ASTRAZENECA	25MG/ML 50MG/ML 50MG/ML 75MG/ML 100MG/ML 100MG/ML 100MG/ML	A089781 001 Mar 31, 1989 A089782 001 Mar 31, 1989 A089783 001 Mar 31, 1989 A089784 001 Mar 31, 1989 A089785 001 Mar 31, 1989 A089786 001 Mar 31, 1989 A089787 001 Mar 31, 1989 A089788 001 Mar 31, 1989
BAXTER HLTHCARE	25MG/ML 50MG/ML 75MG/ML 100MG/ML	A088279 001 Jun 15, 1984 A088280 001 Jun 15, 1984 A088281 001 Jun 15, 1984 A088282 001 Jun 15, 1984
INTL MEDICATION	10MG/ML	A086332 001
PARKE DAVIS	50MG/ML 75MG/ML 100MG/ML	A080364 002 A080364 003 A080364 001
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE HOSPIRA	10MG/ML	A040305 001 Mar 10, 1999
MALLINCKRODT	10MG/ML	A040163 001 May 12, 1997

DISCONTINUED DRUG PRODUCT LIST

6 - 216 (of 346)

MEPERIDINE HYDROCHLORIDE

SYRUP; ORAL				
DEMEROL				
SANOFI AVENTIS US	50MG/5ML		N005010	005
TABLET; ORAL				
MEPERIDINE HYDROCHLORIDE				
DURAMED PHARMS BARR	50MG		A040318	001 Oct 05, 1999
	100MG		A040318	002 Oct 05, 1999
MUTUAL PHARM	50MG		A080448	001
	100MG		A080448	002
WYETH AYERST	50MG		A080454	001

MEPERIDINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION				
MEPERGAN				
BAXTER HLTHCARE CORP	25MG/ML; 25MG/ML		N011730	001

MEPHENTERMINE SULFATE

INJECTABLE; INJECTION				
WYAMINE SULFATE				
BAXTER HLTHCARE CORP	EQ 15MG BASE/ML		N008248	002
	EQ 30MG BASE/ML		N008248	001

MEPHENYTOIN

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				
GRAHAM CHEM	3%		A083559	001
INTL MEDICATION	1%		A087509	001 Oct 05, 1982

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		N011284	001
	400MG		N011284	002
TABLET; ORAL				
AMOSENE				
FERNDALE LABS	400MG		A084030	001
BAMATE				
ALRA	200MG		A080380	001
	400MG		A080380	002
EQUANIL				
WYETH AYERST	200MG		N010028	005
	400MG		N010028	004
MEPRIAM				
TEVA	400MG		N016069	001

DISCONTINUED DRUG PRODUCT LIST

6 - 217 (of 346)

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

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
MALLARD	400MG	N015072	002
MK LABS	200MG	N014368	004
	400MG	N014368	002
MUTUAL PHARM	200MG	A080699	001
	400MG	A080699	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	A080655	001
	400MG	N014547	001
SCHERER LABS	400MG	A083343	001
SOLVAY	200MG	A084435	001
STANLABS PHARM	200MG	N014474	002
	400MG	N014474	004
TABLICAPS	400MG	A083494	001
USL PHARMA	200MG	A087825	001
	400MG	A087826	001
VALEANT PHARM INTL	200MG	N015139	006
	400MG	N015139	005
VANGARD	400MG	A088011	001
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
	400MG	N014359	001

DISCONTINUED DRUG PRODUCT LIST

6 - 218 (of 346)

MEPROBAMATE

TABLET; ORAL TRANMEP SOLVAY	400MG 400MG	A084369 001 N016249 001
-----------------------------------	----------------	----------------------------

MERSALYL SODIUM; THEOPHYLLINE

INJECTABLE; INJECTION MERSALYL-THEOPHYLLINE WATSON LABS	100MG/ML;50MG/ML	A084875 001
---	------------------	-------------

MESALAMINE

SUPPOSITORY; RECTAL CANASA AXCAN	500MG	N021252 001 Jan 05, 2001
ROWASA ALAVEN PHARM	500MG	N019919 001 Dec 18, 1990

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 WATSON LABS	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 WATSON LABS	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
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

DISCONTINUED DRUG PRODUCT LIST

6 - 219 (of 346)

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				
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
NEPHRON	0.4%		A071855	001 Jul 14, 1988
	0.6%		A071726	001 Jul 14, 1988
WOCKHARDT	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				
MORTON GROVE	10MG/5ML		A071656	001 Oct 13, 1987
TEVA	10MG/5ML		A072761	001 Feb 27, 1992
TEVA PHARMS	10MG/5ML		A073034	001 Aug 30, 1991
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
	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

METARAMINOL BITARTRATE

INJECTABLE; 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

DISCONTINUED DRUG PRODUCT LIST

6 - 220 (of 346)

METARAMINOL BITARTRATE

INJECTABLE; INJECTION
METARAMINOL BITARTRATE

GD SEARLE LLC EQ 20MG BASE/ML A086418 002

METAXALONE

TABLET; ORAL
SKELAXIN
KING PHARMS 400MG

N013217 001

METFORMIN HYDROCHLORIDE

TABLET; ORAL
GLUCOPHAGE
BRISTOL MYERS SQUIBB 625MG
750MG

N020357 003 Nov 05, 1998
N020357 004 Nov 05, 1998

METFORMIN HYDROCHLORIDE

BARR 500MG
850MG
1GM

A075971 001 Jan 25, 2002
A075971 002 Jan 25, 2002
A075971 003 Jan 25, 2002

IPCA LABS LTD 500MG
850MG
1GM

A078422 001 Aug 06, 2007
A078422 002 Aug 06, 2007
A078422 003 Aug 06, 2007

IVAX SUB TEVA PHARMS 500MG
625MG
750MG

A075975 001 Jan 24, 2002
A075975 004 Jan 24, 2002
A075975 005 Jan 24, 2002

TEVA 500MG
850MG
1GM

A075975 002 Jan 24, 2002
A075975 003 Jan 24, 2002
A076328 001 Dec 16, 2002

TABLET, EXTENDED RELEASE; ORAL
METFORMIN HYDROCHLORIDE

ACTAVIS ELIZABETH 500MG
750MG

A076450 001 Oct 01, 2004
A076878 001 Apr 13, 2005

BARR 500MG

A076496 001 Nov 25, 2005

IVAX SUB TEVA PHARMS 500MG

A076545 001 Dec 01, 2003

MUTUAL PHARM 500MG

A077124 001 Dec 21, 2005

SANDOZ 500MG

A076223 001 Dec 14, 2004

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL
AVANDAMET
SB PHARMCO 500MG;EQ 1MG BASE

N021410 001 Oct 10, 2002

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

SYRUP; ORAL
DOLOPHINE HYDROCHLORIDE
ROXANE 10MG/30ML

N006134 004

TABLET; ORAL
METHADONE HYDROCHLORIDE

ROXANE 5MG
10MG
40MG
SANDOZ 5MG

A088108 001 Mar 08, 1983
A088109 001 Mar 08, 1983
A074081 001 Apr 28, 1995
A040241 001 May 29, 1998

DISCONTINUED DRUG PRODUCT LIST

6 - 221 (of 346)

METHADONE HYDROCHLORIDE

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 ABLE	5MG	A040529	001 Feb 25, 2004
REXAR	5MG	A084931	001
	10MG	A084931	002
TEVA	5MG	A086359	001
TABLET, EXTENDED RELEASE; ORAL DESOXYN LUNDBECK INC	5MG	N005378	004
	10MG	N005378	003
	15MG	N005378	005

METHANTHELINE BROMIDE

TABLET; ORAL BANTHINE SHIRE	50MG	N007390	001
-----------------------------------	------	---------	-----

METHARBITAL

TABLET; ORAL GEMONIL ABBOTT	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

DISCONTINUED DRUG PRODUCT LIST

6 - 222 (of 346)

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				
CEDAR PHARMS	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	

METHOCARBAMOL

INJECTABLE; INJECTION				
METHOCARBAMOL				
MARSAM PHARMS LLC	100MG/ML	A089849	001	Dec 27, 1991
TABLET; ORAL				
DELAXIN				
FERNDALE LABS	500MG	A085454	001	
FORBAXIN				
FOREST LABS	750MG	A085136	001	
METHOCARBAMOL				
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	
MUTUAL PHARM	500MG	A084488	001	
	750MG	A084486	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 223 (of 346)

METHOCARBAMOLTABLET; ORAL
METHOCARBAMOL

ROXANE	500MG	A088646	001	Feb 29, 1984
	750MG	A088647	001	Feb 29, 1984
SANDOZ	500MG	A087283	001	
	750MG	A087282	001	
SOLCO HLTHCARE	500MG	A086989	001	
	750MG	A086988	001	
SOLVAY	500MG	A084448	001	
	750MG	A084449	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	
	750MG	A083605	002	

METHOHEXITAL SODIUMINJECTABLE; INJECTION
BREVITAL SODIUM

JHP PHARMS	5GM/VIAL	N011559	003
------------	----------	---------	-----

METHOTREXATE SODIUMINJECTABLE; 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

APP PHARMS	EQ 25MG BASE/ML	A040265	001	Feb 26, 1999
	EQ 1GM BASE/VIAL	A040266	001	Feb 26, 1999
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
---------	------------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 224 (of 346)

METHOTREXATE SODIUM

INJECTABLE; INJECTION			
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

METHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION			
VASOXYL			
GLAXOSMITHKLINE	10MG/ML	N006772	002
	20MG/ML	N006772	001

METHOXALEN

CAPSULE; ORAL			
METHOXALEN			
SANDOZ	10MG	A087781	001 Jun 08, 1982

METHSCOPOLAMINE BROMIDE

TABLET; ORAL			
METHSCOPOLAMINE BROMIDE			
PVT FORM	2.5MG	A080970	001

METHYCLOTHIAZIDE

TABLET; ORAL			
AQUATENSEN			
MEDPOINTE PHARM HLC	5MG	N017364	001
METHYCLOTHIAZIDE			
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
SANDOZ	2.5MG	A089835	001 Aug 18, 1988
	5MG	A089837	001 Aug 18, 1988
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

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

METHYLDOPA

SUSPENSION; ORAL			
ALDOMET			
MERCK	250MG/5ML	N018389	001

DISCONTINUED DRUG PRODUCT LIST

6 - 225 (of 346)

METHYLDOPA

TABLET; ORAL

ALDOMET

MERCK	125MG	N013400	003
	250MG	N013400	001
	500MG	N013400	002

METHYLDOPA

ACCORD HLTH	125MG	A070070	003	Oct 15, 1985
DURAMED PHARMS BARR	250MG	A071006	001	Dec 16, 1986
	500MG	A071009	001	Dec 16, 1986
HALSEY	125MG	A071751	001	Mar 28, 1988
	250MG	A071752	001	Mar 28, 1988
	500MG	A071753	001	Mar 28, 1988
MUTUAL PHARM	125MG	A070073	001	Oct 09, 1986
	250MG	A070060	001	Oct 09, 1986
	500MG	A070074	001	Oct 09, 1986
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
	250MG	N018934	001	Jun 29, 1984
	500MG	N018934	002	Jun 29, 1984
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	Jun 03, 1986
BAXTER HLTHCARE	50MG/ML	A070291	001	Jul 01, 1986
HOSPIRA	50MG/ML	A070691	001	Jun 19, 1987
	50MG/ML	A070698	001	Jun 15, 1987
	50MG/ML	A070699	001	Jun 15, 1987
	50MG/ML	A070849	001	Jun 19, 1987
MARSAM PHARMS LLC	50MG/ML	A071812	001	Dec 22, 1987
SMITH AND NEPHEW	50MG/ML	A070841	001	Jan 02, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 226 (of 346)

METHYLPHENIDATE HYDROCHLORIDE

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
	10MG	A040321	002	Feb 05, 2002
	20MG	A040321	003	Feb 05, 2002
TABLET, EXTENDED RELEASE; ORAL				
	METHYLPHENIDATE HYDROCHLORIDE			
ABLE	20MG	A076032	001	May 09, 2001
ACTAVIS ELIZABETH	20MG	A075450	001	Dec 21, 2001

METHYLPREDNISOLONE

TABLET; ORAL

MEDROL

PHARMACIA AND UPJOHN	2MG	N011153	002	
	24MG	N011153	005	
METHYLPREDNISOLONE				
HEATHER	4MG	A085650	001	
PAR PHARM	16MG	A089207	001	Apr 25, 1988
	24MG	A089208	001	Apr 25, 1988
	32MG	A089209	001	Apr 25, 1988
SANDOZ	4MG	A087341	001	
WATSON LABS	4MG	A086161	001	Feb 09, 1982
	16MG	A086159	001	Feb 09, 1982

METHYLPREDNISOLONE ACETATE

ENEMA; RECTAL

MEDROL

PHARMACIA AND UPJOHN	40MG/BOT	N018102	001	
----------------------	----------	---------	-----	--

INJECTABLE; INJECTION

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	

M-PREDROL

BEL MAR

AKORN	40MG/ML	A086666	001	
	80MG/ML	A087135	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

DISCONTINUED DRUG PRODUCT LIST

6 - 227 (of 346)

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

HOSPIRA	EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL	A085853 001 A085855 001 A085854 001 A089173 001 Aug 18, 1987 A085852 001 A089174 001 Aug 18, 1987
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
ELKINS SINK	EQ 40MG BASE/VIAL	A086906 001
INTL MEDICATION	EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A087812 001 Feb 09, 1983 A087813 001 Feb 09, 1983 A087851 001 Feb 09, 1983 A087852 001 Feb 09, 1983
TEVA PARENTERAL	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	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

N003240 004

10MG

N003240 005

METHYLTESTOSTERONE

IMPAK LABS 10MG

A084287 001

LILLY 10MG

A080256 001

PUREPAC PHARM 10MG

A080308 001

10MG

A080475 001

DISCONTINUED DRUG PRODUCT LIST

6 - 228 (of 346)

METHYLTESTOSTERONE

TABLET; Buccal/Sublingual				
METHYLTESTOSTERONE				
PVT FORM	5MG	A083836	001	
TABLICAPS	10MG	A085125	001	
USL PHARMA	10MG	A080271	001	
TABLET; ORAL				
METANDREN				
NOVARTIS	10MG	N003240	001	
	25MG	N003240	003	
METHYLTESTOSTERONE				
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
LILLY	25MG	A080256	002	
PARKE DAVIS	10MG	A084244	001	
	25MG	A084241	001	
PUREPAC PHARM	10MG	A080309	001	
	10MG	A080475	002	
	25MG	A080310	001	
	25MG	A080475	003	
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

DISCONTINUED DRUG PRODUCT LIST

6 - 229 (of 346)

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

HOSPIRA	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
SMITH AND NEPHEW	EQ 5MG BASE/ML	A070623	001	Mar 02, 1987
	EQ 10MG BASE/2ML	A070622	001	Mar 02, 1987
REGLAN				
BAXTER HLTHCARE CORP	EQ 10MG BASE/ML	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
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	A072384	001	Jun 02, 1988
	EQ 10MG BASE	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 HLTHCARE	EQ 10MG BASE	A072639	001	May 09, 1991
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
	EQ 10MG BASE	A071536	001	Apr 28, 1993
PAR PHARM	EQ 10MG BASE	A070342	001	Mar 25, 1986
SANDOZ	EQ 5MG BASE	A072436	001	Jun 22, 1989
	EQ 5MG BASE	A074478	001	Oct 05, 1995
	EQ 10MG BASE	A070850	001	Feb 03, 1987
	EQ 10MG BASE	A072215	001	Jan 30, 1990
	EQ 10MG BASE	A074478	002	Oct 05, 1995
SCHERING	EQ 10MG BASE	A070598	001	Feb 02, 1987
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				
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

DISCONTINUED DRUG PRODUCT LIST

6 - 230 (of 346)

METOLAZONE

TABLET; ORAL DIULO					
GD SEARLE LLC	2.5MG	N018535	001		
	5MG	N018535	002		
	10MG	N018535	003		
METOLAZONE					
ROXANE	10MG	A076482	002	Apr 29, 2004	
WATSON LABS	10MG	A076891	001	Jul 21, 2004	
MYKROX					
UCB INC	0.5MG	N019532	001	Oct 30, 1987	

METOPROLOL FUMARATE

TABLET, EXTENDED RELEASE; ORAL LOPRESSOR					
NOVARTIS	EQ 100MG TARTRATE	N019786	001	Dec 27, 1989	
	EQ 200MG TARTRATE	N019786	002	Dec 27, 1989	
	EQ 300MG TARTRATE	N019786	003	Dec 27, 1989	
	EQ 400MG TARTRATE	N019786	004	Dec 27, 1989	

METOPROLOL TARTRATE

TABLET; ORAL METOPROLOL TARTRATE					
APOTHECON	50MG	A074258	001	Jan 27, 1994	
	100MG	A074258	002	Jan 27, 1994	
MYLAN	50MG	A073666	001	Dec 21, 1993	
	100MG	A073666	002	Dec 21, 1993	
PUREPAC PHARM	50MG	A074380	001	Jul 29, 1994	
	100MG	A074380	002	Jul 29, 1994	
SOLCO HLTHCARE	50MG	A074453	001	Apr 27, 1995	
	100MG	A074453	002	Apr 27, 1995	
TEVA	50MG	A074143	001	Sep 30, 1994	
	100MG	A074143	002	Sep 30, 1994	
TEVA PHARMS	50MG	A074333	001	Jan 27, 1994	
	100MG	A074333	002	Jan 27, 1994	

METRIZAMIDE

INJECTABLE; INJECTION AMIPAQUE					
GE HEALTHCARE	2.5GM/VIAL	N017982	003	Sep 12, 1983	
	3.75GM/VIAL	N017982	001		
	6.75GM/VIAL	N017982	002		
	13.5GM/VIAL	N017982	004	Sep 12, 1983	

METRONIDAZOLE

CAPSULE; ORAL METRONIDAZOLE					
ABLE	375MG	A076505	001	Nov 13, 2003	
INJECTABLE; INJECTION 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	
ELKINS SINK	500MG/100ML	N018907	001	Mar 30, 1984	
INTL MEDICATION	500MG/100ML	A070004	001	May 08, 1985	
WATSON LABS	500MG/100ML	A070042	001	Dec 20, 1984	
	500MG/100ML	A070170	001	Apr 01, 1986	
TABLET; ORAL METROMIDOL					
LABS AF	250MG	A074523	001	Oct 24, 1996	
	500MG	A074523	002	Oct 24, 1996	

DISCONTINUED DRUG PRODUCT LIST

6 - 231 (of 346)

METRONIDAZOLE

TABLET; ORAL

METRONIDAZOLE

ABLE	250MG	A076519	001	Jun 27, 2003
	500MG	A076519	002	Jun 27, 2003
HALSEY	250MG	A070021	001	Apr 02, 1985
	500MG	A070593	001	Feb 27, 1986
IVAX SUB TEVA PHARMS	250MG	N018517	001	
	500MG	N018517	002	May 05, 1982
LNK	250MG	N019029	001	Apr 10, 1984
MUTUAL PHARM	250MG	N018818	001	Feb 16, 1983
	500MG	N018818	002	Feb 16, 1983
PAR PHARM	250MG	N018845	001	Aug 18, 1983
	500MG	N018930	001	Aug 18, 1983
SANDOZ	250MG	N018620	001	Mar 04, 1982
	250MG	N018740	001	Oct 22, 1982
	500MG	N018620	002	Jun 02, 1983
	500MG	N018740	002	Oct 22, 1982
SUPERPHARM	250MG	A070008	001	Dec 11, 1984
	500MG	A070009	001	Dec 11, 1984
WATSON LABS	250MG	N018599	001	Sep 17, 1982
	500MG	N018599	002	Feb 13, 1984
WORLD GEN	250MG	A070040	001	Jan 29, 1985
	500MG	A070039	001	Jan 29, 1985
PROTOSTAT				
ORTHO MCNEIL PHARM	250MG	N018871	001	Mar 02, 1983
	500MG	N018871	002	Mar 02, 1983
SATRIC				
SAVAGE LABS	250MG	A070029	001	Mar 19, 1985
	500MG	A070731	001	Jun 08, 1987
TABLET, EXTENDED RELEASE; ORAL				
METRONIDAZOLE				
ABLE	750MG	A076462	001	Jun 25, 2003

METRONIDAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

FLAGYL I.V.

PFIZER	EQ 500MG BASE/VIAL	N018353	001	
METRONIDAZOLE HYDROCHLORIDE				
ABRAXIS PHARM	EQ 500MG BASE/VIAL	A070295	001	Oct 15, 1985

METYRAPONE

TABLET; ORAL

METOPIRONE

NOVARTIS

250MG

N012911 001

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

SANDOZ

150MG

A074450 001 May 16, 1996

200MG

A074450 002 May 16, 1996

250MG

A074450 003 May 16, 1996

MEXITIL

BOEHRINGER INGELHEIM 150MG

N018873 002 Dec 30, 1985

200MG

N018873 003 Dec 30, 1985

250MG

N018873 004 Dec 30, 1985

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

DISCONTINUED DRUG PRODUCT LIST

6 - 232 (of 346)

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

BAYER PHARMS	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

ASTRAZENECA	EQ 5MG BASE/ML	A075263	001	Jun 26, 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
--	----------------	---------	-----	--------------

VERSED

HLR	EQ 1MG BASE/ML	N018654	002	May 26, 1987
-----	----------------	---------	-----	--------------

	EQ 5MG BASE/ML	N018654	001	Dec 20, 1985
--	----------------	---------	-----	--------------

SYRUP; ORAL

VERSED

ROCHE	EQ 2MG BASE/ML	N020942	001	Oct 15, 1998
-------	----------------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 233 (of 346)

MILRINONE LACTATE

INJECTABLE; INJECTION					
MILRINONE LACTATE					
BAXTER HLTHCARE CORP	EQ 1MG BASE/ML		A075852	001	May 28, 2002
BIONICHE PHARMA	EQ 1MG BASE/ML		A076428	001	Jun 16, 2003
HOSPIRA	EQ 1MG BASE/ML		A075830	001	May 28, 2002
	EQ 1MG BASE/ML		A075884	001	May 28, 2002
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER					
BAXTER HLTHCARE	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)		A075510	001	May 28, 2002
	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)		A076259	001	Aug 08, 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					
DYNACIN					
MEDICIS	EQ 50MG BASE		A063066	001	Aug 14, 1990
MINOCIN					
TRIAK PHARMS	EQ 50MG BASE		N050315	002	
	EQ 100MG BASE		N050315	001	
TRIAK PHARMS LLC	EQ 75MG BASE		N050649	003	Feb 12, 2001
INJECTABLE; INJECTION					
MINOCIN					
LEDERLE	EQ 100MG BASE/VIAL		A062139	001	
SUSPENSION; ORAL					
MINOCIN					
TRIAK PHARMS	EQ 50MG BASE/5ML		N050445	001	
TABLET; ORAL					
MINOCYCLINE HYDROCHLORIDE					
TRIAK PHARMS	EQ 50MG BASE		N050451	003	Aug 10, 1982
	EQ 100MG BASE		N050451	002	Aug 10, 1982
TABLET, EXTENDED RELEASE; ORAL					
MINOCYCLINE HYDROCHLORIDE					
LUPIN LTD	EQ 55MG BASE		A091424	002	Nov 30, 2011

MINOXIDIL

SOLUTION; TOPICAL					
MINOXIDIL (FOR MEN)					
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)					
SIGHT PHARMS	2%		A074743	001	Oct 18, 1996
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

DISCONTINUED DRUG PRODUCT LIST

6 - 234 (of 346)

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076308	001	Jun 20, 2003
	30MG	A076308	002	Jun 20, 2003
	45MG	A076308	003	Jun 20, 2003
ACTAVIS TOTOWA	15MG	A076241	001	Jun 25, 2003
	30MG	A076241	002	Jun 25, 2003
	45MG	A076241	003	Jun 25, 2003
IVAX SUB TEVA PHARMS	15MG	A076244	001	Dec 22, 2003
	30MG	A076244	002	Dec 22, 2003
	45MG	A076244	003	Dec 22, 2003
ROXANE	15MG	A076270	001	Jun 19, 2003
	30MG	A076270	002	Jun 19, 2003
	45MG	A076270	003	Jun 19, 2003
SANDOZ	15MG	A076189	001	Jun 19, 2003
	30MG	A076189	002	Jun 19, 2003
	45MG	A076189	003	Jun 19, 2003
TABLET, ORALLY DISINTEGRATING; ORAL				
MIRTAZAPINE				
ACTAVIS TOTOWA	15MG	A076689	001	Aug 31, 2005
	30MG	A076689	002	Aug 31, 2005
	45MG	A076689	003	Aug 31, 2005

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	
	20MG/VIAL	N050450	002	
BRISTOL MYERS	5MG/VIAL	A062336	001	
	20MG/VIAL	A062336	002	
	40MG/VIAL	A062336	003	Mar 10, 1988

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

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

ABBOTT	EQ 2MG BASE/ML	N020098	001	Jan 22, 1992
MIVACRON IN DEXTROSE 5%	IN PLASTIC CONTAINER			
ABBOTT	EQ 0.5MG BASE/ML	N020098	002	Jan 22, 1992
	EQ 50MG BASE/100ML	N020098	003	Jan 22, 1992
MIVACURIUM CHLORIDE				
PISGAH LABS	EQ 2MG BASE/ML	A078562	001	Apr 30, 2009

MOLINDONE HYDROCHLORIDE

CAPSULE; ORAL

MOBAN

ENDO PHARMS	5MG	N017111	001
	10MG	N017111	002
	25MG	N017111	003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS	20MG/ML	N017938	001
-------------	---------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 235 (of 346)

MOLINDONE HYDROCHLORIDE

TABLET; ORAL

MOBAN

ENDO PHARMS

5MG

N017111 004

10MG

N017111 005

25MG

N017111 006

50MG

N017111 007

100MG

N017111 008

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

N019753 001 Jun 19, 1990

250MG

N019753 002 Jun 19, 1990

300MG

N019753 003 Jun 19, 1990

MORPHINE SULFATE

INJECTABLE; INJECTION

MORPHINE SULFATE

HOSPIRA

0.5MG/ML

N019917 001 Oct 30, 1992

MALLINCKRODT

1MG/ML

N020631 001 Jul 03, 1996

2MG/ML

N020631 002 Jul 03, 1996

INJECTABLE, LIPOSOMAL; EPIDURAL

DEPODUR

EKR THERAP

15MG/1.5ML (10MG/ML)

N021671 002 May 18, 2004

20MG/2ML (10MG/ML)

N021671 003 May 18, 2004

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

WATSON LABS

100MG

A075656 001 Jan 30, 2001

MOXALACTAM DISODIUM

INJECTABLE; INJECTION

MOXAM

LILLY

EQ 250MG BASE/VIAL

N050550 001

EQ 500MG BASE/VIAL

N050550 002

EQ 1GM BASE/VIAL

N050550 003

EQ 2GM BASE/VIAL

N050550 004

EQ 10GM BASE/VIAL

N050550 008

MYCOPHENOLATE MOFETIL

TABLET; ORAL

MYCOPHENOLATE MOFETIL

DR REDDYS LABS LTD 500MG

A090464 001 Sep 13, 2010

NABUMETONE

TABLET; ORAL

NABUMETONE

ACTAVIS ELIZABETH

500MG

A079093 001 Feb 27, 2009

750MG

A079093 002 Feb 27, 2009

COPELEY PHARM

750MG

A075179 001 Jun 06, 2000

SANDOZ

500MG

A075590 001 Feb 25, 2002

750MG

A075590 002 Feb 25, 2002

DISCONTINUED DRUG PRODUCT LIST

6 - 236 (of 346)

NABUMETONE

TABLET; ORAL RELAFEN	SMITHKLINE BEECHAM	500MG 750MG	N019583 001	Dec 24, 1991
			N019583 002	Dec 24, 1991

NADOLOL

TABLET; ORAL CORGARD	KING PHARMS	120MG 160MG	N018063 003 N018063 004	
NADOLOL				
	IVAX SUB TEVA PHARMS	120MG 160MG	A074255 002 A074255 003	Jan 24, 1996 Jan 24, 1996
	TEVA PHARMS	80MG 120MG 160MG	A074368 001 A074368 002 A074368 003	Aug 31, 1994 Aug 31, 1994 Aug 31, 1994

NAFCILLIN 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	A061984 001 A061984 002 A061984 003 A061984 005	
MARSAM PHARMS LLC		EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 1.5GM BASE/VIAL EQ 2GM BASE/VIAL EQ 4GM BASE/VIAL EQ 10GM BASE/VIAL	A062844 001 A062844 002 A062844 003 A062844 004 A062844 005 A063008 001	Oct 26, 1988 Oct 26, 1988 Oct 26, 1988 Oct 26, 1988 Oct 26, 1988 Sep 29, 1988
SANDOZ NALLPEN		EQ 500MG BASE/VIAL	A062527 001	Aug 02, 1984
GLAXOSMITHKLINE		EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL EQ 10GM BASE/VIAL	A061999 001 A061999 002 A062755 001 A061999 003 A062755 002 A061999 004	
UNIPEN				
WYETH AYERST		EQ 500MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL EQ 4GM BASE/VIAL EQ 10GM BASE/VIAL EQ 20GM BASE/VIAL	A062717 001 N050320 001 A062717 002 A062717 004 N050320 003 N050320 004 N050320 005 N050320 006	Dec 16, 1986 Dec 16, 1986
UNIPEN IN PLASTIC CONTAINER WYETH AYERST		EQ 1GM BASE/VIAL	N050320 002	
TABLET; ORAL UNIPEN				
WYETH AYERST		EQ 500MG BASE	N050462 001	

DISCONTINUED DRUG PRODUCT LIST

6 - 237 (of 346)

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE

ABRAXIS PHARM	10MG/ML	A070751	001	Jul 02, 1986
	20MG/ML	A070752	001	Sep 24, 1986
NALBUPHINE HYDROCHLORIDE				
ABBOTT	1.5MG/ML	N020200	001	Mar 12, 1993
	20MG/ML	A070917	001	Feb 03, 1989
ASTRAZENECA	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
BARR	10MG/ML	A072075	001	Apr 10, 1989
	20MG/ML	A074471	001	Mar 19, 1998
NUBAIN		A074471	002	Mar 19, 1998
ENDO PHARMS	10MG/ML	N018024	001	
	20MG/ML	N018024	002	May 27, 1982

NALIDIXIC ACID

SUSPENSION; ORAL

NEGRAM

SANOFI AVENTIS US	250MG/5ML	N017430	001	
TABLET; ORAL				
NALIDIXIC ACID				
MUTUAL PHARM	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
NEGRAM				
SANOFI AVENTIS US	250MG	N014214	002	
	500MG	N014214	004	
	1GM	N014214	005	

NALMEFENE HYDROCHLORIDE

INJECTABLE; INJECTION

REVEX

BAXTER HLTHCARE CORP	EQ 0.1MG BASE/ML	N020459	001	Apr 17, 1995
	EQ 1MG BASE/ML	N020459	002	Apr 17, 1995

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

BAXTER HLTHCARE	0.4MG/ML	A070298	001	Sep 24, 1986
	0.4MG/ML	A070299	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
	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
	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

DISCONTINUED DRUG PRODUCT LIST

6 - 238 (of 346)

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HYDROCHLORIDE

ASTRAZENECA	0 .4MG /ML 0 .4MG /ML 0 .4MG /ML 0 .4MG /ML 0 .4MG /ML 1MG /ML 1MG /ML 1MG /ML	A072086 001 A072087 001 A072088 001 A072089 001 A072090 001 A072091 001 A072092 001 A072093 001	Apr 11, 1989 Apr 11, 1989
BAXTER HLTHCARE	0 .02MG /ML 1MG /ML 1MG /ML 1MG /ML	A071272 001 A071273 001 A071274 001 A071287 001	May 24, 1988 May 24, 1988 May 24, 1988 May 24, 1988
HOSPIRA	0 .02MG /ML 0 .02MG /ML 0 .02MG /ML 0 .4MG /ML	A070171 001 A070252 001 A070253 001 A070255 001	Sep 24, 1986 Jan 16, 1987 Jan 16, 1987 Jan 07, 1987
INTL MEDICATION	0 .4MG /ML 1MG /ML	A070417 001 A072115 001	Sep 24, 1986 Apr 27, 1988
MARSAM PHARMS LLC	0 .4MG /ML	A071811 001	Jul 19, 1988
SMITH AND NEPHEW	0 .02MG /ML 0 .4MG /ML 0 .4MG /ML	A071671 001 A071681 001 A071682 001	Nov 17, 1987 Nov 17, 1987 Nov 17, 1987
SOLOPAK	0 .02MG /ML 0 .4MG /ML	A071672 001 A071683 001	Nov 17, 1987 Nov 17, 1987
WATSON LABS	0 .4MG /ML	A071339 001	Nov 18, 1987
NARCAN			
BRISTOL MYERS SQUIBB	0 .4MG /ML 1MG /ML 1MG /ML	A071083 001 A071084 001 A071311 001	Jul 28, 1988 Jul 28, 1988 Jul 28, 1988
ENDO PHARMS	0 .02MG /ML 0 .4MG /ML 1MG /ML	N016636 002 N016636 001 N016636 003	
			Jun 14, 1982

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN NX

SANOFI AVENTIS US EQ 0 .5MG BASE;EQ 50MG BASE N018733 001 Dec 16, 1982

NANDROLONE DECANOATE

INJECTABLE; INJECTION

DECA-DURABOLIN

ORGANON USA INC	50MG /ML 100MG /ML 200MG /ML	N013132 001 N013132 002 N013132 003	Jun 12, 1986 Jun 12, 1986 Jun 12, 1986
NANDROLONE DECANOATE			
ABRAXIS PHARM	100MG /ML 200MG /ML	A088290 001 A088317 001	Oct 03, 1983 Oct 14, 1983
AKORN	100MG /ML	A087519 001	Sep 28, 1983
WATSON LABS	50MG /ML 50MG /ML 100MG /ML 100MG /ML 200MG /ML	A086385 001 A087598 001 A088554 001 A086598 001 A087599 001 A088128 001	Jan 13, 1984 Oct 06, 1983 Feb 10, 1986 Jan 13, 1984 Oct 06, 1983 Dec 05, 1983

NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

DURABOLIN

ORGANON USA INC 25MG /ML N011891 001
50MG /ML N011891 002

DISCONTINUED DRUG PRODUCT LIST

6 - 239 (of 346)

NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

NANDROLONE PHENPROPIONATE

WATSON LABS	25MG/ML	A086386	001	Jun 17, 1983
	50MG/ML	A087488	001	Jun 17, 1983

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

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
----------	------	---------	-----	--------------

NAPROXEN

TABLET; ORAL

NAPROXEN

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

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

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

TEVA	250MG	A074129	001	Dec 21, 1993
	375MG	A074129	002	Dec 21, 1993
	500MG	A074129	003	Dec 21, 1993

TEVA PHARMS	250MG	A074207	001	Dec 21, 1993
	375MG	A074207	002	Dec 21, 1993
	500MG	A074207	003	Dec 21, 1993

WATSON LABS	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

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

ABLE	EQ 250MG BASE	A076544	001	Aug 22, 2003
	EQ 500MG BASE	A076544	002	Aug 22, 2003

HAMILTON PHARMS	EQ 250MG BASE	A074106	001	Aug 31, 1993
	EQ 500MG BASE	A074106	002	Aug 31, 1993

DISCONTINUED DRUG PRODUCT LIST

6 - 240 (of 346)

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

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 250MG BASE	A074162	001	Dec 21, 1993
	EQ 500MG 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 500MG BASE	A074195	002	Dec 21, 1993
TABLET, EXTENDED RELEASE; ORAL				
NAPROXEN SODIUM				
WATSON LABS FLORIDA	EQ 375MG BASE	A075416	002	Apr 23, 2003
	EQ 500MG BASE	A075416	001	Aug 27, 2002

NEDOCROMIL SODIUM

AEROSOL, METERED; INHALATION

TILADE

KING PHARMS 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

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	A076072	001	Sep 16, 2003
	50MG	A076302	001	Sep 16, 2003
	100MG	A076072	002	Sep 16, 2003
	100MG	A076302	002	Sep 16, 2003
	150MG	A076072	003	Sep 16, 2003
	150MG	A076302	003	Sep 16, 2003
	200MG	A076072	004	Sep 16, 2003
	200MG	A076302	004	Sep 16, 2003
	250MG	A076072	005	Sep 16, 2003
	250MG	A076302	005	Sep 16, 2003

DISCONTINUED DRUG PRODUCT LIST

6 - 241 (of 346)

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

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

NEOMYCIN SULFATESOLUTION; ORAL
MYCIFRADIN

PHARMACIA AND UPJOHN EQ 87.5MG BASE/5ML N050285 001

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 SULFATECREAM; 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; PREDNISOLONE ACETATEOINTMENT; 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

NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATEOINTMENT; OPHTHALMIC
NEO-HYDELTRASOL

MERCK EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE N050378 001

NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDECREAM; 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

DISCONTINUED DRUG PRODUCT LIST

6 - 242 (of 346)

NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL					
NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE					
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	
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	
WEST WARD	500MG		A083718	001	
NICOLAR					
SANOFI AVENTIS US	500MG		A083823	001	
TABLET, EXTENDED RELEASE; ORAL					
NIACIN					
BARR	500MG		A076378	001	Apr 26, 2005
	750MG		A076378	002	Apr 26, 2005
	1GM		A076250	001	Apr 14, 2005
NIASPAN					
ABBOTT	375MG		N020381	001	Jul 28, 1997
NIASPAN TITRATION STARTER PACK					
ABBOTT	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, EXTENDED RELEASE; ORAL					
CARDENE SR					
EKR THERAP	45MG		N020005	002	Feb 21, 1992

NICLOSAMIDE

TABLET, CHEWABLE; ORAL					
NICLOCIDE					
BAYER PHARMS	500MG		N018669	001	May 14, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 243 (of 346)

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					
WATSON LABS	EQ 2MG BASE		A076568	001	Jul 29, 2004
	EQ 4MG BASE		A076569	002	Jul 29, 2004

NIFEDIPINE

CAPSULE; ORAL					
ADALAT					
BAYER PHARMS	10MG		N019478	001	Nov 27, 1985
	20MG		N019478	002	Sep 17, 1986
NIFEDIPINE					
CATALENT	20MG		A074045	001	Apr 30, 1992
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

NILUTAMIDE

TABLET; ORAL					
NILANDRON					
SANOFI AVENTIS US	50MG		N020169	001	Sep 19, 1996

NIMODIPINE

CAPSULE; ORAL					
NIMOTOP					
BAYER PHARMS	30MG		N018869	001	Dec 28, 1988

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL					
SULAR					
SHIONOGI INC	10MG		N020356	001	Feb 02, 1995
	20MG		N020356	002	Feb 02, 1995
	30MG		N020356	003	Feb 02, 1995
	40MG		N020356	004	Feb 02, 1995

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	
	100MG		A080017	002	
NITROFURANTOIN					
ELKINS SINK	50MG		A080003	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 244 (of 346)

NITROFURANTOIN

TABLET; ORAL

NITROFURANTOIN

ELKINS SINK	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 SODIUM

INJECTABLE; INJECTION

IVADANTIN

PROCTER AND GAMBLE	EQ 180MG BASE/VIAL	N012402	001
--------------------	--------------------	---------	-----

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; 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
NITROFURANTOIN MACROCRYSTALLINE				
WATSON LABS	50MG	A070248	001	Jun 24, 1988
	100MG	A070249	001	Jun 24, 1988

NITROFURAZONE

CREAM; 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

AMBIK	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
--------------	-------------	---------	-----	--------------

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

MYLAN TECHNOLOGIES	0.1MG/HR	A074992	004	Nov 12, 1999
	0.2MG/HR	A074992	003	Nov 12, 1999
	0.4MG/HR	A074992	002	Nov 12, 1999
	0.6MG/HR	A074992	001	Nov 12, 1999

DISCONTINUED DRUG PRODUCT LIST

6 - 245 (of 346)

NITROGLYCERIN

FILM, EXTENDED RELEASE; TRANSDERMAL TRANSDERM-NITRO					
NOVARTIS	0.1MG/HR 0.2MG/HR 0.4MG/HR 0.6MG/HR 0.8MG/HR		N020144	001 002 003 004 005	Feb 27, 1996 Feb 27, 1996 Feb 27, 1996 Feb 27, 1996 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 A071159	001 001	Jan 05, 1982 Feb 28, 1990
NITROGLYCERIN					
ABRAXIS PHARM	5MG/ML 5MG/ML		A070077 A071203	001 001	Dec 13, 1985 May 08, 1987
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 A070634	001 001	Jun 19, 1986 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 A070863 N018588 A070871 A070872	001 001 002 001 001	
TRIDIL					
HOSPIRA	0.5MG/ML 5MG/ML		N018537 N018537	002 001	Jun 16, 1983

NIZATIDINE

CAPSULE; ORAL NIZATIDINE					
IVAX SUB TEVA PHARMS	150MG 300MG		A075461	001 002	Jul 08, 2002 Jul 08, 2002

NONOXYNOL-9

AEROSOL; VAGINAL DELFEN					
PERSONAL PRODS	12.5%		N014349	002	

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	

DISCONTINUED DRUG PRODUCT LIST

6 - 246 (of 346)

NORETHINDRONE ACETATE

TABLET; ORAL
 NORLUTATE
 PARKE DAVIS 5MG N012184 002

NORFLOXACIN

SOLUTION/DROPS; OPHTHALMIC
 CHIBROXIN
 MERCK 0.3% N019757 001 Jun 17, 1991

NORGESTREL

TABLET; ORAL
 OVRETTE
 WYETH PHARMS 0.075MG N017031 001

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL
 AVENTYL HYDROCHLORIDE
 LILLY EQ 10MG BASE N014684 001
 EQ 25MG BASE N014684 002
 NORTRIPTYLINE HYDROCHLORIDE
 SANDOZ EQ 10MG BASE A074054 001 Dec 31, 1992
 EQ 10MG BASE A074835 001 Jun 30, 1997
 EQ 25MG BASE A074054 002 Dec 31, 1992
 EQ 25MG BASE A074835 002 Jun 30, 1997
 EQ 50MG BASE A074054 003 Dec 31, 1992
 EQ 50MG BASE A074835 003 Jun 30, 1997
 EQ 75MG BASE A074054 004 Dec 31, 1992
 EQ 75MG BASE A074835 004 Jun 30, 1997
 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
 PAMELOR
 MALLINCKRODT INC EQ 10MG BASE/5ML N018012 001

NYSTATIN

CREAM; TOPICAL
 CANDEX
 BAYER PHARMS 100,000 UNITS/GM A061810 001
 MYCOSTATIN
 BRISTOL MYERS SQUIBB 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
 NYSTATIN
 TARO 100,000 UNITS/GM A062457 001 Jul 28, 1983
 TEVA 100,000 UNITS/GM A061966 001
 LOTION; TOPICAL
 CANDEX
 BAYER PHARMS 100,000 UNITS/ML N050233 001
 OINTMENT; TOPICAL
 MYCOSTATIN
 WESTWOOD SQUIBB 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
 BRISTOL MYERS SQUIBB 200,000 UNITS N050619 001 Apr 09, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 247 (of 346)

NYSTATIN

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 BRISTOL MYERS SQUIBB	100,000 UNITS/GM	A060578	001	
SUPPOSITORY; VAGINAL NYsert WARNER CHILCOTT	100,000 UNITS	N050478	001	
SUSPENSION; ORAL MYCOSTATIN APOTHECON	100,000 UNITS/ML	A061533	001	
NYSTATIN ALPHARMA US PHARMS	100,000 UNITS/ML	A062571	001	Oct 29, 1985
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
ROXANE	100,000 UNITS/ML	A062832	001	Dec 27, 1991
TEVA	100,000 UNITS/ML	A062670	001	Jun 18, 1987
	100,000 UNITS/ML	A062776	001	Dec 17, 1987
NYSTEX SAVAGE LABS	100,000 UNITS/ML	A062519	001	Jul 06, 1984
TABLET; ORAL MYCOSTATIN BRISTOL MYERS SQUIBB	500,000 UNITS	A060574	001	
NILSTAT LEDERLE	500,000 UNITS	A061151	001	
NYSTATIN HERITAGE PHARMS INC	500,000 UNITS	A062474	001	Dec 22, 1983
QUANTUM PHARMICS	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 BRISTOL MYERS SQUIBB	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 PHARMICS	100,000 UNITS	A062509	001	Apr 03, 1984
SANDOZ	100,000 UNITS	A061965	001	
TEVA	100,000 UNITS	A062502	001	Dec 23, 1983
WATSON LABS	100,000 UNITS	A062176	001	

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL MYCOLOG-II APOTHECON	100,000 UNITS/GM; 0.1% 100,000 UNITS/GM; 0.1%	A060576	002	May 01, 1985
MYCO-TRIACET II TEVA	100,000 UNITS/GM; 0.1%	A062606	001	May 15, 1985
MYKACET ACTAVIS MID ATLANTIC	100,000 UNITS/GM; 0.1%	A061954	002	Sep 20, 1985
MYTREX F SAVAGE LABS	100,000 UNITS/GM; 0.1%	A062367	001	May 28, 1985
		A062597	001	Oct 08, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 248 (of 346)

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

NYSTATIN AND TRIAMCINOLONE 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-TRIAMCINOLONE ACETONIDE				
PHARMADERM	100,000 UNITS/GM;0.1%	A062596	001	Oct 08, 1985

OINTMENT; TOPICAL

MYCOLOG-II

APOTHECON	100,000 UNITS/GM;0.1%	A060572	001	Jun 28, 1985
-----------	-----------------------	---------	-----	--------------

MYCO-TRIACET II

TEVA	100,000 UNITS/GM;0.1%	A062045	002	Nov 26, 1985
------	-----------------------	---------	-----	--------------

MYKACET

ACTAVIS MID ATLANTIC	100,000 UNITS/GM;0.1%	A062733	001	Mar 09, 1987
----------------------	-----------------------	---------	-----	--------------

MYTREX F

SAVAGE LABS	100,000 UNITS/GM;0.1%	A062601	001	Oct 09, 1985
-------------	-----------------------	---------	-----	--------------

NYSTATIN AND TRIAMCINOLONE 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-TRIAMCINOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM;0.1%	A062603	001	Oct 09, 1985
------------	-----------------------	---------	-----	--------------

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
---------	---------	---------	-----	--------------

TABLET; ORAL

FLOXIN

JANSSEN PHARMS	200MG	N019735	001	Dec 28, 1990
	300MG	N019735	002	Dec 28, 1990
	400MG	N019735	003	Dec 28, 1990

OFLOXACIN

LARKEN LABS	200MG	A076093	001	Sep 02, 2003
	300MG	A076093	002	Sep 02, 2003
	400MG	A076093	003	Sep 02, 2003

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

HOSPIRA	EQ 2MG BASE/ML	A076695	001	Dec 26, 2006
---------	----------------	---------	-----	--------------

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

HOSPIRA	EQ 0.64MG BASE/ML	A076978	001	Feb 26, 2007
---------	-------------------	---------	-----	--------------

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

HOSPIRA	EQ 2MG BASE/ML	A076696	001	Dec 26, 2006
---------	----------------	---------	-----	--------------

ZOFTRAN AND DEXTROSE IN PLASTIC CONTAINER

GLAXOSMITHKLINE	EQ 0.64MG BASE/ML	N020403	001	Jan 31, 1995
-----------------	-------------------	---------	-----	--------------

ORPHENADRINE CITRATE

TABLET, EXTENDED RELEASE; ORAL

NORFLEX

MEDICIS	100MG	N012157	001	
---------	-------	---------	-----	--

ORPHENADRINE CITRATE

ASCOT	100MG	A088067	001	Apr 06, 1983
-------	-------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 249 (of 346)

ORPHENADRINE CITRATE

TABLET, EXTENDED RELEASE; ORAL ORPHENADRINE CITRATE		
SANDOZ	100MG	A085046 001
WATSON LABS	100MG	A084303 001

ORPHENADRINE 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		
APOTHECON	EQ 250MG BASE	A061450 002
	EQ 500MG BASE	A061450 001
TEVA	EQ 250MG BASE	A062222 001
	EQ 500MG BASE	A062222 002
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
PROSTAPHLIN APOTHECON	EQ 250MG BASE/5ML	N050194 001
INJECTABLE; INJECTION BACTOCILL		
GLAXOSMITHKLINE	EQ 500MG BASE/VIAL	A061334 009 Mar 26, 1982
	EQ 1GM BASE/VIAL	A061334 006 Mar 26, 1982
	EQ 1GM BASE/VIAL	A062736 001 Dec 19, 1986
	EQ 2GM BASE/VIAL	A061334 007 Mar 26, 1982
	EQ 2GM BASE/VIAL	A062736 002 Dec 19, 1986
	EQ 4GM BASE/VIAL	A061334 008 Mar 26, 1982
	EQ 10GM BASE/VIAL	A061334 010
OXACILLIN SODIUM APOTHECON		
	EQ 250MG BASE/VIAL	N050195 001
	EQ 500MG BASE/VIAL	N050195 002
	EQ 1GM BASE/VIAL	N050195 003
	EQ 2GM BASE/VIAL	N050195 004
	EQ 4GM BASE/VIAL	N050195 005
ELKINS SINK	EQ 250MG BASE/VIAL	A062711 001 Feb 03, 1989
	EQ 500MG BASE/VIAL	A062711 002 Feb 03, 1989
	EQ 1GM BASE/VIAL	A062711 003 Feb 03, 1989
	EQ 2GM BASE/VIAL	A062711 004 Feb 03, 1989
	EQ 4GM BASE/VIAL	A062711 005 Feb 03, 1989
	EQ 10GM BASE/VIAL	A062711 006 Feb 03, 1989
IBI	EQ 125MG BASE/VIAL	A062798 003 Dec 11, 1995
	EQ 250MG BASE/VIAL	A062798 004 Dec 11, 1995
	EQ 500MG BASE/VIAL	A062798 005 Dec 11, 1995
	EQ 1GM BASE/VIAL	A062798 001 Dec 11, 1995

DISCONTINUED DRUG PRODUCT LIST

6 - 250 (of 346)

OXACILLIN SODIUM

INJECTABLE; INJECTION OXACILLIN SODIUM					
IBI	EQ 2GM BASE/VIAL	A062798	002	Dec 11, 1995	
MARSAM PHARMS LLC	EQ 10GM BASE/VIAL	A062984	001	Sep 29, 1988	
SANDOZ	EQ 250MG BASE/VIAL	A061490	001		
	EQ 500MG BASE/VIAL	A061490	002		
WATSON LABS	EQ 250MG BASE/VIAL	A062856	001	Oct 26, 1988	
	EQ 500MG BASE/VIAL	A062856	002	Oct 26, 1988	
	EQ 1GM BASE/VIAL	A062856	003	Oct 26, 1988	
	EQ 2GM BASE/VIAL	A062856	004	Oct 26, 1988	
	EQ 4GM BASE/VIAL	A062856	005	Oct 26, 1988	

OXALIPLATIN

INJECTABLE; IV (INFUSION) ELOXATIN					
SANOFI AVENTIS US	50MG/VIAL	N021492	001	Aug 09, 2002	
	100MG/VIAL	N021492	002	Aug 09, 2002	

OXAMNIQUINE

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	

OXAPROZIN

TABLET; ORAL OXAPROZIN					
ACTAVIS ELIZABETH	600MG	A075843	001	Oct 03, 2001	
MYLAN	600MG	A075851	001	Aug 17, 2001	
SANDOZ	600MG	A075842	001	Apr 12, 2001	
	600MG	A075850	001	Apr 27, 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	
MUTUAL PHARM	10MG	A071026	002	Aug 10, 1987	
	15MG	A071026	003	Aug 10, 1987	
	30MG	A071026	001	Aug 10, 1987	
MYLAN	10MG	A071713	001	Oct 20, 1987	
	15MG	A071714	001	Oct 20, 1987	
	30MG	A071715	001	Oct 20, 1987	
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 251 (of 346)

OXAZEPAM

TABLET; ORAL					
OXAZEPAM					
MUTUAL PHARM	15MG		A070683	001	Jan 16, 1987
PARKE DAVIS	15MG		A071508	001	Feb 02, 1987
WATSON LABS	15MG		A071494	001	Apr 21, 1987
SERAX					
ALPHARMA US PHARMS	15MG		N015539	008	

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

OXTRIPTYLLINE

SOLUTION; ORAL					
CHOLEDYL					
PARKE DAVIS	100MG/5ML		N009268	012	Nov 27, 1984
OXTRIPTYLLINE					
MORTON GROVE	100MG/5ML		A088243	001	Dec 05, 1983
SYRUP; ORAL					
CHOLEDYL					
PARKE DAVIS	50MG/5ML		N009268	011	
OXTRIPTYLLINE PEDIATRIC					
MORTON GROVE	50MG/5ML		A088242	001	Dec 05, 1983
TABLET, DELAYED RELEASE; ORAL					
CHOLEDYL					
PARKE DAVIS	100MG		N009268	003	
	200MG		N009268	007	
OXTRIPTYLLINE					
WATSON LABS	100MG		A087866	001	Aug 25, 1983
	200MG		A087835	001	Aug 25, 1983

OXYBUTYNIN CHLORIDE

SYRUP; ORAL					
DITROPAN					
ORTHO MCNEIL JANSSEN	5MG/5ML		N018211	001	
TABLET; ORAL					
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					
OXYCONTIN					
PURDUE PHARMA LP	10MG		N020553	001	Dec 12, 1995
	15MG		N020553	006	Sep 18, 2006
	20MG		N020553	002	Dec 12, 1995
	30MG		N020553	007	Sep 18, 2006
	40MG		N020553	003	Dec 12, 1995
	60MG		N020553	008	Sep 18, 2006
	80MG		N020553	004	Jan 06, 1997
	160MG		N020553	005	Mar 15, 2000
ROXICODONE					
ROXANE	10MG		N020932	001	Oct 26, 1998
	30MG		N020932	002	Oct 26, 1998

DISCONTINUED DRUG PRODUCT LIST

6 - 252 (of 346)

OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION OPANA			
ENDO PHARMS	1.5MG/ML	N011707	001
SUPPOSITORY; RECTAL NUMORPHAN			
ENDO PHARMS	5MG	N011738	004
TABLET, EXTENDED RELEASE; ORAL OPANA ER			
ENDO PHARMS	7.5MG	N021610	005 Feb 29, 2008
	15MG	N021610	006 Feb 29, 2008

OXPHENBUTAZONE

TABLET; ORAL OXYPHENBUTAZONE			
WATSON LABS	100MG	A088399	001 Sep 17, 1984
TANDEARIL			
NOVARTIS	100MG	N012542	004 Sep 03, 1982

OXPHENCYCLIMINE HYDROCHLORIDE

TABLET; ORAL DARICON			
PFIZER	10MG	N011612	001

OXYPHENONIUM 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			
IMPAX LABS	EQ 250MG BASE	A060760	001
PROTER	EQ 250MG BASE	A060869	001
PUREPAC PHARM	EQ 250MG BASE	A060634	001
WEST WARD	EQ 250MG BASE	A060770	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

DISCONTINUED DRUG PRODUCT LIST

6 - 253 (of 346)

OXYTOCIN

INJECTABLE; INJECTION					
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
OXYTOCIN 5 USP UNITS IN DEXTROSE 5%					
ABBOTT	1USP UNITS/100ML	N019185	001	Mar 29,	1985
SYNTOCINON					
NOVARTIS	10USP UNITS/ML	N018245	001		
SOLUTION; NASAL					
SYNTOCINON					
NOVARTIS	40USP UNITS/ML	N012285	001		

PACLITAXEL

INJECTABLE; INJECTION					
PACLITAXEL					
HOSPIRA	6MG/ML	A076233	001	Aug 01,	2002
PLIVA LACHEMA	6MG/ML	A077413	001	Mar 12,	2008
TEVA PARENTERAL	6MG/ML	A075297	001	Jan 25,	2002
TAXOL					
BRISTOL MYERS SQUIBB	6MG/ML	N020262	001	Dec 29,	1992

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

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION					
AREDIA					
NOVARTIS	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

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					
ASTRAZENECA	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
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
PAVULON					
ORGANON USA INC	1MG/ML	N017015	002		
	2MG/ML	N017015	001		

DISCONTINUED DRUG PRODUCT LIST

6 - 254 (of 346)

PARAMETHADIONE

CAPSULE; ORAL PARADIONE			
ABBOTT	150MG	N006800	003
	300MG	N006800	001
SOLUTION; ORAL PARADIONE			
ABBOTT	300MG/ML	N006800	002

PARAMETHASONE ACETATE

TABLET; ORAL HALDRONE			
LILLY	1MG	N012772	005
	2MG	N012772	006

PARGYLINE HYDROCHLORIDE

TABLET; ORAL EUTONYL			
ABBOTT	10MG	N013448	002
	25MG	N013448	003
	50MG	N013448	004

PAROMOMYCIN SULFATE

CAPSULE; ORAL HUMATIN			
KING PFIZER	EQ 250MG BASE	A062310	001
PARKADEALE	EQ 250MG BASE	A060521	001
SYRUP; ORAL HUMATIN			
PARKE DAVIS	EQ 125MG BASE/5ML	A060522	001

PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL PAXIL			
GLAXOSMITHKLINE	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
TABLET; ORAL PAROXETINE HYDROCHLORIDE			
ACTAVIS ELIZABETH	EQ 10MG BASE	A076968	001 Jun 21, 2010
	EQ 20MG BASE	A076968	002 Jun 21, 2010
	EQ 30MG BASE	A076968	003 Jun 21, 2010
	EQ 40MG BASE	A076968	004 Jun 21, 2010
ROXANE	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, 2007
SANDOZ	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
TEVA 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, 2007
PAXIL			
GLAXOSMITHKLINE	EQ 50MG BASE	N020031	004 Dec 29, 1992

PEMOLINE

TABLET; ORAL CYLERT			
ABBOTT	18.75MG	N016832	001

DISCONTINUED DRUG PRODUCT LIST

6 - 255 (of 346)

PEMOLINE

TABLET; ORAL CYLERT					
ABBOTT	37.5MG	N016832	002		
	75MG	N016832	003		
PEMOLINE					
ACTAVIS TOTOWA	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 TOTOWA	37.5MG	A075678	001	Jul 26, 2000	
TEVA PHARMS	37.5MG	A075555	001	Feb 18, 2000	

PENBUTOLOL SULFATE

TABLET; ORAL LEVATOL					
SCHWARZ PHARMA	10MG	N018976	001	Dec 30, 1987	

PENICILLAMINE

CAPSULE; ORAL CUPRIMINE					
ATON	125MG	N019853	002		

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		

DISCONTINUED DRUG PRODUCT LIST

6 - 256 (of 346)

PENICILLIN G POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN-2

TEVA	250,000 UNITS/5ML	A060307	003
PENTIDS '200'	200,000 UNITS/5ML	A062149	001
APOTHECON			
PENTIDS '400'	400,000 UNITS/5ML	A062149	002
APOTHECON			
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

MARSAM PHARMS LLC

	1,000,000 UNITS/VIAL	A062991	Sep 13, 1988
	5,000,000 UNITS/VIAL	A062991	Sep 13, 1988
	10,000,000 UNITS/VIAL	A062991	Sep 13, 1988
	20,000,000 UNITS/VIAL	A062991	Sep 13, 1988

PARKE DAVIS

	1,000,000 UNITS/VIAL	A062003	001
	5,000,000 UNITS/VIAL	A062003	002

PFIZER

SANDOZ

	1,000,000 UNITS/VIAL	A065079	001
			Aug 30, 2002

PFIZERPEN

	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

	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

DISCONTINUED DRUG PRODUCT LIST

6 - 257 (of 346)

PENICILLIN G POTASSIUM

TABLET; ORAL 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; IM-IV PENICILLIN G SODIUM				
MARSAM PHARMS LLC	5,000,000 UNITS/VIAL		A063014	001 Sep 13, 1988
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

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
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

DISCONTINUED DRUG PRODUCT LIST

6 - 258 (of 346)

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PEN-VEE K

WYETH AYERST	EQ 125MG BASE/5ML	A060007 001
	EQ 250MG BASE/5ML	A060007 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

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

PEN-VEE K

WYETH AYERST	EQ 125MG BASE	A060006 001
	EQ 250MG BASE	A060006 002
	EQ 500MG BASE	A060006 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

VEETIDS '250'

APOTHECON	EQ 250MG BASE	A061164 001
-----------	---------------	-------------

DISCONTINUED DRUG PRODUCT LIST

6 - 259 (of 346)

PENICILLIN V POTASSIUM

TABLET; ORAL VEETIDS '250'	EQ 250MG BASE	A062156	002
APOTHECON VEETIDS '500'	EQ 500MG BASE	A061164	002
APOTHECON	EQ 500MG BASE	A062156	001

PENTAGASTRIN

INJECTABLE; INJECTION PEPTAVLON	WYETH AYERST	0 .25MG/ML	N017048	001
------------------------------------	--------------	------------	---------	-----

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION NEBUPENT	APP PHARMS	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

PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL TALWIN 50	SANOFI AVENTIS US	EQ 50MG BASE	N016732	001
---------------------------	-------------------	--------------	---------	-----

PENTETATE CALCIUM TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS PENTETATE CALCIUM TRISODIUM	HAMELN PHARMS	EQ 1GM BASE/5ML (EQ 200MG BASE/ML)	N021749	001	Aug 11, 2004
--	---------------	------------------------------------	---------	-----	--------------

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE; INJECTION YTTERBIUM YB 169 DTPA	3M	2mCi/ML	N017518	001
--	----	---------	---------	-----

PENTETATE ZINC TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS PENTETATE ZINC TRISODIUM	HAMELN PHARMS	EQ 1GM BASE/5ML (EQ 200MG BASE/ML)	N021751	001	Aug 11, 2004
---	---------------	------------------------------------	---------	-----	--------------

PENTOBARBITAL

ELIXIR; ORAL NEMBUTAL	OVATION PHARMS	18 .2MG/5ML	A083244	001
--------------------------	----------------	-------------	---------	-----

PENTOBARBITAL SODIUM

CAPSULE; ORAL NEMBUTAL SODIUM	OVATION PHARMS	30MG	A084095	001
		50MG	A084093	001
		100MG	A083245	001
PENTOBARBITAL SODIUM				
LANNETT		50MG	A085937	001
		100MG	A085915	001
VITARINE		100MG	A083284	001
WHITEWORTH TOWN PLSN		100MG	A083338	001
SODIUM PENTOBARBITAL				
ANABOLIC		100MG	A084590	001
ELKINS SINK		100MG	A083368	001

DISCONTINUED DRUG PRODUCT LIST

6 - 260 (of 346)

PENTOBARBITAL SODIUM

CAPSULE; ORAL				
SODIUM PENTOBARBITAL				
EVERYLIFE	100MG	A083259	001	
HALSEY	100MG	A084677	001	
IVAX SUB TEVA PHARMS	50MG	A083461	001	
	100MG	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 SINK	50MG/ML	A083270	001	
SODIUM PENTOBARBITAL				
WYETH AYERST	50MG/ML	A083261	001	
SUPPOSITORY; RECTAL				
NEMBUTAL				
OVATION 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 TARTRATE

INJECTABLE; INJECTION				
ANSOLYSSEN				
WYETH AYERST	10MG/ML	N009372	001	

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL				
PENTOXIFYLLINE				
ACTAVIS ELIZABETH	400MG	A074878	001	Jul 09, 1997
HERITAGE PHARMS INC	400MG	A074877	001	Jul 08, 1997
TEVA	400MG	A075199	001	Sep 03, 1999

PERFLUBRON

LIQUID; ORAL				
IMAGENT				
ALLIANCE PHARM	100%	N020091	001	Aug 13, 1993

PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE

PASTE; TOPICAL				
SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS				
US ARMY	50%;50%	N021084	001	Feb 17, 2000

PERGOLIDE MESYLATE

TABLET; 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

DISCONTINUED DRUG PRODUCT LIST

6 - 261 (of 346)

PERGOLIDE MESYLATE

TABLET; ORAL PERMAX	VALEANT PHARM INTL	EQ 0.25MG BASE EQ 1MG BASE	N019385 002 Dec 30, 1988
			N019385 003 Dec 30, 1988

PERMETHRIN

LOTION; TOPICAL NIX	GLAXOSMITHKLINE	1%	N019435 001 Mar 31, 1986
------------------------	-----------------	----	--------------------------

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	IVAX PHARMS	2MG 4MG 8MG 16MG	A089707 001 Sep 10, 1987 A089708 001 Sep 10, 1987 A089456 001 Sep 10, 1987 A089457 001 Sep 10, 1987
TRILAFON	SCHERING	2MG 4MG 8MG 16MG	N010775 001 N010775 002 N010775 003 N010775 004
TABLET, EXTENDED RELEASE; ORAL TRILAFON	SCHERING	8MG	N011361 002

PHENACEMIDE

TABLET; ORAL PHENURONE	ABBOTT	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 PENAZOPYRIDINE 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 35MG	A086523 001 A086524 001
----------------------------	---------	--------------	----------------------------

DISCONTINUED DRUG PRODUCT LIST

6 - 262 (of 346)

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL				
PHENAZINE				
MAST MM	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
CAPSULE, EXTENDED RELEASE; ORAL				Sep 22, 1982
MELFIAT-105				
NUMARK	105MG		A087487	001
PHENDIMETRAZINE TARTRATE				Oct 13, 1982
GRAHAM DM	105MG		A087214	001
	105MG		A088020	001
	105MG		A088028	001
	105MG		A088062	001
	105MG		A088063	001
	105MG		A088111	001
SANDOZ	105MG		A087378	001
SPRX-105				
NUMARK	105MG		A088024	001
X-TROZINE L.A.				Dec 22, 1982
SHIRE RICHWOOD	105MG		A087371	001
TABLET; ORAL				Aug 24, 1982
ADPHEN				
FERNDALE LABS	35MG		A083655	001
ALPHAZINE				
SANDOZ	35MG		A085034	001
CAM-METRAZINE				
ABC HOLDING	35MG		A085511	001
CAMALL	35MG		A085756	001
TG UNITED LABS	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

DISCONTINUED DRUG PRODUCT LIST

6 - 263 (of 346)

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

PHENDIMETRAZINE TARTRATE

BARR	35MG	A084831	001	
	35MG	A084834	001	
	35MG	A084835	001	
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	
	35MG	A086365	001	
	35MG	A086370	001	
SOLVAY	35MG	A083993	001	
TG UNITED LABS	35MG	A085761	001	
	35MG	A085941	001	Jun 27, 1983
USL PHARMA	35MG	A083805	001	
	35MG	A084398	001	
	35MG	A084399	001	
VITARINE	35MG	A085519	001	
	35MG	A086005	001	
	35MG	A086106	001	
WATSON LABS	35MG	A085767	001	
	35MG	A085768	001	
	35MG	A085770	001	
	35MG	A085773	001	
PLEGINE				
WYETH AYERST	35MG	N012248	001	
STATOBEX				
TEVA	35MG	A086013	001	
STATOBEX-G				
TEVA	35MG	A085095	001	
X-TROZINE				
SHIRE RICHWOOD	35MG	A086550	001	
	35MG	A086551	001	
	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

DISCONTINUED DRUG PRODUCT LIST

6 - 264 (of 346)

PHENMETRAZINE HYDROCHLORIDE

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
<u>PHENTERMINE HYDROCHLORIDE</u>		
ABC HOLDING	30MG	A085411 001
ABLE	15MG	A040497 001 Mar 13, 2003
	30MG	A040403 001 Aug 30, 2001
	30MG	A040427 001 Aug 30, 2001
ACTAVIS TOTOWA	15MG	A040460 001 Jan 14, 2003
	30MG	A040227 001 Jun 18, 1997
	30MG	A040448 001 Jan 22, 2003
	37.5MG	A040228 001 Jun 19, 1997
CAMALL	15MG	A086735 001
	30MG	A087226 001
DURAMED PHARMS BARR	30MG	A088948 001 Apr 25, 1986
IVAX PHARMS	30MG	A086329 001
MUTUAL PHARM	37.5MG	A040527 001 Oct 23, 2003
SANDOZ	30MG	A087208 001
	30MG	A087223 001
	37.5MG	A088414 001 Oct 19, 1983
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
TG UNITED LABS	18.75MG	A088576 001 May 23, 1984
	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
USL PHARMA	30MG	A084487 001 Apr 09, 1982
	30MG	A088430 001 Mar 27, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 265 (of 346)

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL					
<u>PHENTERMINE HYDROCHLORIDE</u>					
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		
<u>PHENTERMINE HYDROCHLORIDE</u>					
ABLE	37.5MG	A040402	001	Aug 30,	2001
ACTAVIS TOTOWA	37.5MG	A040190	001	May 30,	1997
IVAX PHARMS	8MG	A085553	001		
SANDOZ	8MG	A085671	001		
	8MG	A085689	001		
TG UNITED LABS	8MG	A083923	001		
	8MG	A085319	001		
	37.5MG	A087805	001	Dec 06,	1982
	37.5MG	A088596	001	Apr 04,	1984
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		

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL					
<u>IONAMIN</u>					
UCB INC	EQ 15MG BASE	N011613	004		
	EQ 30MG BASE	N011613	002		
<u>PHENTERMINE RESIN 30</u>					
QUANTUM PHARMICS	EQ 30MG BASE	A089120	001	Feb 04,	1988

PHENYL AMINOSALICYLATE

POWDER; ORAL					
<u>PHENY-PAS-TEBAMIN</u>					
PHARM RES ASSOC	50%	N011695	002		
<u>TABLET; ORAL</u>					
PHENY-PAS-TEBAMIN					
PHARM RES ASSOC	500MG	N011695	003		

PHENYLBUTAZONE

CAPSULE; ORAL					
<u>AZOLID</u>					
SANOFI AVENTIS US	100MG	A087260	001		
BUTAZOLIDIN					
NOVARTIS	100MG	N008319	009		
<u>PHENYLBUTAZONE</u>					
IVAX PHARMS	100MG	A088218	001	Jun 24,	1983
MUTUAL PHARM	100MG	A088994	001	Dec 04,	1985
SANDOZ	100MG	A087774	001	Jun 16,	1982
WATSON LABS	100MG	A087756	001	Dec 17,	1982
TABLET; ORAL					
AZOLID					
SANOFI AVENTIS US	100MG	A087091	001		
BUTAZOLIDIN					
NOVARTIS	100MG	N008319	008		

DISCONTINUED DRUG PRODUCT LIST

6 - 266 (of 346)

PHENYLBUTAZONE

TABLET; ORAL					
PHENYLBUTAZONE					
MUTUAL PHARM	100MG		A088863	001	Dec 04, 1985
SANDOZ	100MG		A084339	001	
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

PHENYTOIN SODIUM

CAPSULE; ORAL					
DIPHENYLAN SODIUM					
LANNETT	30MG PROMPT		A080857	001	
	100MG PROMPT		A080857	002	
EXTENDED PHENYTOIN SODIUM					
BARR	100MG EXTENDED		A040435	001	Jun 20, 2003
PLIVA	100MG EXTENDED		A089441	001	Dec 18, 1986
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					
IVAX SUB TEVA PHARMS	100MG PROMPT		A080259	001	
WATSON LABS	100MG PROMPT		A080905	001	
INJECTABLE; INJECTION					
DILANTIN					
PARKE DAVIS	50MG/ML		N010151	001	
PHENYTOIN SODIUM					
APP PHARMS	50MG/ML		A089003	001	May 31, 1985
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					
BIOVAIL TECHNOLOGIES	1MG/0.5ML		N012223	002	

DISCONTINUED DRUG PRODUCT LIST

6 - 267 (of 346)

PHYTONADIONE

INJECTABLE; INJECTION				
AQUAMEPHYTON				
KONAKION	BIOVAIL TECHNOLOGIES	10MG/ML	N012223	001
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

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	GENPHARM	5MG	A074013	001 Sep 24, 1992
		10MG	A074018	001 Sep 24, 1992
IVAX SUB TEVA PHARMS	5MG		A073687	001 Feb 26, 1993
		10MG	A073687	002 Feb 26, 1993
MUTUAL PHARM	5MG		A074063	001 Jan 27, 1994
		10MG	A074063	002 Jan 27, 1994
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
SANDOZ	5MG		A073608	001 Mar 29, 1993
		10MG	A073609	001 Mar 29, 1993
TEVA	5MG		A073661	001 Oct 31, 1993
		10MG	A074123	001 Apr 17, 1997
		10MG	A073661	002 Oct 31, 1993
			A074123	002 Apr 17, 1997
VISKEN				
NOVARTIS	5MG		N018285	001 Sep 03, 1982
		10MG	N018285	002 Sep 03, 1982

PIPECURONIUM BROMIDE

INJECTABLE; INJECTION				
ARDUAN	ORGANON USA INC	10MG/VIAL	N019638	001 Jun 26, 1990

PIPERACETAZINE

TABLET; ORAL				
QUIDE	DOW PHARM	10MG	N013615	001
		25MG	N013615	002

DISCONTINUED DRUG PRODUCT LIST

6 - 268 (of 346)

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			
IMPAK LABS	EQ 250MG BASE	A080874	001

PIPOBROMANTABLET; ORAL
VERCYTE

ABBOTT	10MG	N016245	001
	25MG	N016245	002

PIRBUTEROL ACETATEAEROSOL, METERED; INHALATION
MAXAIR

MEDICIS	EQ 0.2MG BASE/INH	N019009	001	Dec 30, 1986
---------	-------------------	---------	-----	--------------

PIROXICAMCAPSULE; ORAL
PIROXICAM

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
MUTUAL PHARM	20MG	A073536	001	Mar 12, 1993
MYLAN	10MG	A074043	001	Sep 22, 1992
	20MG	A074043	002	Sep 22, 1992
ROXANE	10MG	A073651	001	Feb 26, 1993
	20MG	A073651	002	Feb 26, 1993
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

DISCONTINUED DRUG PRODUCT LIST

6 - 269 (of 346)

PLICAMYCIN

INJECTABLE; INJECTION
 MITHRACIN
 PFIZER 2.5MG/VIAL N050109 001

POLYESTRADIOL PHOSPHATE

INJECTABLE; INJECTION
 ESTRADURIN
 WYETH AYERST 40MG/AMP N010753 001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL
 POLYETHYLENE GLYCOL 3350
 TEVA PHARMS 17GM/SCOOPFUL A077445 001 May 04, 2006

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

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
 MEDA PHARMS 120GM/PACKET;1.49GM/PACKET;3.36GM/PACKET N018983 005 Oct 26, 1984
 T;2.92GM/PACKET;11.36GM/PACKET
 227.1GM/PACKET;2.82GM/PACKET;6.36GM/PAC N018983 004 Oct 26, 1984
 KET;5.53GM/PACKET;21.5GM/PACKET
 360GM/PACKET;4.47GM/PACKET;10.08GM/PACK N018983 006 Oct 26, 1984
 ET;8.76GM/PACKET;34.08GM/PACKET

FOR SUSPENSION; ORAL

CO-LAV
 BOCA PHARMA 240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/ A073428 001 Jan 28, 1992
 BOT;22.72GM/BOT

COLOVAGE
 DYNAPHARM 227.1GM/PACKET;2.82GM/PACKET;6.36GM/PAC A071320 001 Apr 20, 1988
 KET;5.53GM/PACKET;21.5GM/PACKET

E-Z-EM PREP LYTE
 E Z EM 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/ A071278 001 Nov 21, 1988
 BOT;22.74GM/BOT

GLYCOPREP
 GOLDLINE 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/ A072319 001 Dec 23, 1988
 BOT;22.74GM/BOT

GO-EVAC
 BOCA PHARMA 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/ A073433 001 Apr 28, 1992
 BOT;22.74GM/BOT

PEG-LYTE
 SANDOZ 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/ A073098 001 Aug 31, 1993
 BOT;22.74GM/BOT

POLYMYXIN B SULFATE

INJECTABLE; INJECTION
 AEROSPORIN
 GLAXOSMITHKLINE EQ 500,000 U BASE/VIAL A062036 001
 POWDER; FOR RX COMPOUNDING
 POLYMYXIN B SULFATE
 PADDOCK LLC 100,000,000 UNITS/BOT A062455 001 Jul 27, 1983

POLYTHIAZZIDE

TABLET; ORAL
 RENESE
 PFIZER 1MG N012845 001
 2MG N012845 002
 4MG N012845 003

DISCONTINUED DRUG PRODUCT LIST

6 - 270 (of 346)

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 20MEO/PACKET N019561 003 Aug 26, 1988

INJECTABLE; INJECTION
POTASSIUM CHLORIDE
ABRAXIS PHARM 2MEq/ML A080204 001

ABRAKAT PHARM 2MEQ/ML A080204 001
 2MEQ/ML A084290 001
 2MEQ/ML A086713 001
 2MEQ/ML A086714 001
 2MEQ/ML A087787 001 Apr 20, 1982

AKORN 2MEO/ML A088286 001 Sep 05, 1985

APP PHARMS	2MEQ/ML	A087817	001	Oct 20, 1982
BAXTER HLTHCARE	2MEQ/ML	A080203	001	
CD SEARLE, LLC	1MEQ/ML	A086219	001	

GD SEARCS LLC 1MEQ/ML A086219 001
2MEQ/ML A086219 002
2MEQ/ML A086220 002

HOSPIRA	3MEQ/ML	A086219	003
	3MEQ/ML	A086220	001
	4MEQ/ML	A086219	004
	1MEQ/ML	A080205	003
	1MEQ/ML	A083245	003

1MEQ/ML	A083345	003
1.5MEQ/ML	A083345	001
2MEQ/ML	A083345	002
3MEQ/ML	A083345	004

LILLY	2.4MEQ/ML	A080205	004
	3.2MEQ/ML	A080205	005
	2MEQ/ML	N007865	002

LUITPOLD	2MEQ/ML	A080221	001
	2MEQ/ML	A080736	001
	2MEQ/ML	A087584	001
	2MEQ/ML	A087585	001

DISCONTINUED DRUG PRODUCT LIST

6 - 271 (of 346)

POTASSIUM CHLORIDEINJECTABLE; INJECTION
POTASSIUM CHLORIDE

MILES	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
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+8				
FUTURE PAK	8MEQ	A070998	001	Jan 25, 1993
KAON CL				
SAVAGE LABS	6 . 7MEQ	N017046	001	
POTASSIUM CHLORIDE				
COPLEY PHARM	8MEQ	A070618	001	Sep 09, 1987
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

UNIV TX 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
--------	--------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 272 (of 346)

POTASSIUM IODIDE

TABLET; ORAL
THYRO-BLOCK
MEDPOINTE PHARM HLC 130MG N018307 001

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
---	--------	---------	-----	--------------

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
CLONMEL HLTHCARE	EQ 1MG BASE	A072705	001	May 16, 1989
	EQ 2MG BASE	A072706	001	May 16, 1989
	EQ 5MG BASE	A072707	001	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
SANDOZ	EQ 1MG BASE	A072576	001	May 16, 1989
	EQ 2MG BASE	A072577	001	May 16, 1989
	EQ 5MG BASE	A072578	001	May 16, 1989
WATSON LABS	EQ 1MG BASE	A072352	001	May 16, 1989
	EQ 2MG BASE	A072333	001	May 16, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 273 (of 346)

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL PRAZOSIN HYDROCHLORIDE					
WATSON LABS	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	0.5%		N010209	002	
SCHERING					
SYRUP; ORAL PREDNISOLONE					
IVAX SUB TEVA PHARMS	15MG/5ML		A040287	001	May 28, 1999
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					
BARR	5MG		A084426	002	
BUNDY	5MG		A083675	001	
ELKINS SINK	5MG		A080625	001	
EVERYLIFE	1MG		A084439	001	
	2.5MG		A084439	002	
	5MG		A084439	003	
FERRANTE	2.5MG		A080562	001	
	5MG		A080562	002	
HEATHER	5MG		A080326	001	
IMPAK 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	
PERRIGO	5MG		A084542	001	
PHOENIX LABS NY	5MG		A080322	001	
PUREPAC PHARM	5MG		A080325	001	
PVT FORM	5MG		A080211	001	
ROXANE	5MG		A080327	002	
SANDOZ	5MG		A080339	001	
	5MG		A084773	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 274 (of 346)

PREDNISOLONE

TABLET; ORAL				
PREDNISOLONE				
WATSON LABS	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
SUSPENSION/DROPS; OPHTHALMIC				Jan 17, 2008
ECONOPRED				
ALCON	0.125%		N017468	001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC				
CETAPRED				
ALCON	0.25%;10%		A087771	001
METIMYD				Aug 06, 1993
SCHERING	0.5%;10%		N010210	002
PREDSULFAIR				Sep 09, 1984
PHARMAFAIR	0.5%;10%		A088032	001
VASOCIDIN				Apr 15, 1983
NOVARTIS	0.5%;10%		A088791	001
SUSPENSION; OPHTHALMIC				Oct 05, 1984
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
PREDSULFAIR				Jul 29, 1983
PHARMAFAIR	0.5%;10%		A088007	001
PREDSULFAIR II				Apr 19, 1983
PHARMAFAIR	0.2%;10%		A088837	001
SULPHRIN				Dec 24, 1985
BAUSCH AND LOMB	0.5%;10%		A088089	001
				Dec 28, 1982

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE; INJECTION				
HYDELTRASOL				
MERCK	EQ 20MG PHOSPHATE/ML		N011583	002

DISCONTINUED DRUG PRODUCT LIST

6 - 275 (of 346)

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE; INJECTION			
PREDNISOLONE SODIUM PHOSPHATE			
WATSON LABS	EQ 20MG PHOSPHATE/ML	A080517	001
OINTMENT; OPHTHALMIC, OTIC			
HYDELTRASOL			
MERCK	EQ 0.25% PHOSPHATE	N011028	001
SOLUTION; ORAL			
PREDNISOLONE SODIUM PHOSPHATE			
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
METRETTON			
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 UNIVERSAL	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

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

DISCONTINUED DRUG PRODUCT LIST

6 - 276 (of 346)

PREDNISONE

TABLET; ORAL				
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
BUNDY	5MG		A083676	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
IMPAX 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
	50MG		A086596	001
NYLOS	5MG		A085115	001
PANRAY	1MG		A080350	001
	2.5MG		A080350	002
	5MG		A080350	003
PERRIGO	5MG		A083059	001
PHARMAVITE	5MG		A084662	002

DISCONTINUED DRUG PRODUCT LIST

6 - 277 (of 346)

PREDNISONE

TABLET; ORAL

PREDNISONE

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	
	5MG	A084774	001	
	10MG	A089983	001	Jan 12, 1989
	20MG	A085813	001	
	50MG	A089984	001	Jan 12, 1989
SCHERER LABS	5MG	A080371	001	
SPERTI	1MG	A080359	001	
	2.5MG	A080359	002	
	5MG	A080359	003	
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
WEST WARD	50MG	A088465	001	Jun 01, 1984
WHITEWORTH TOWN PLSN	2.5MG	A084913	001	
	5MG	A080343	001	
	10MG	A089028	001	Jul 24, 1986
	20MG	A084913	002	
SERVISONE				
LEDERLE	5MG	A080223	001	

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST

ASTRAZENECA	1%	N014763	004
	2%	N014763	005
	3%	N014763	003

CITANEST PLAIN

ASTRAZENECA	4%	N014763	007
-------------	----	---------	-----

PRIMIDONE

SUSPENSION; ORAL

MYSOLINE

NURO PHARMA	250MG/5ML	N010401	001
-------------	-----------	---------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 278 (of 346)

PRIMIDONE

TABLET; ORAL PRIMIDONE WATSON LABS	250MG	A085052 001
--	-------	-------------

PROBENECID

TABLET; ORAL BENEMID MERCK	500MG	N007898 004
PROBENECID 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
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
SANDOZ	250MG	A089219 001 Jul 01, 1986
	375MG	A089220 001 Jul 01, 1986
	500MG	A089221 001 Jul 01, 1986
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
	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
BAXTER HLTHCARE	100MG/ML	A089029 001 Apr 17, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 279 (of 346)

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

BAXTER HLTHCARE	500MG/ML	A089030	001	Apr 17, 1986
HOSPIRA	500MG/ML	A089537	001	Aug 25, 1987
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	A089528	001	May 03, 1988
	500MG/ML	A089529	001	May 03, 1988
WATSON LABS	100MG/ML	A087079	001	
	500MG/ML	A087080	001	
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				
COPLEY PHARM	500MG	A088974	001	Jul 22, 1985
	750MG	A089438	001	Mar 23, 1987
	1GM	A040111	001	Dec 13, 1996
INWOOD LABS	500MG	A089840	001	Mar 06, 1989
PLIVA	250MG	A088958	001	Dec 02, 1985
	500MG	A088959	001	Dec 02, 1985
SANDOZ	250MG	A089369	001	Aug 14, 1987
	500MG	A089284	001	Jun 23, 1986
	500MG	A089370	001	Jan 09, 1987
	750MG	A089371	001	Aug 14, 1987
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	

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVOCAIN

HOSPIRA

1%

A085362 003

2%

A085362 004

10%

A086797 001

PROCAINE HYDROCHLORIDE

ABRAXIS PHARM

1%

A080384 002

1%

A080421 001

2%

A080384 003

2%

A080421 002

DISCONTINUED DRUG PRODUCT LIST

6 - 280 (of 346)

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE

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 HYDROCHLORIDE

INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	40MG/VIAL;100MG/VIAL	N050276	001
	40MG/VIAL;250MG/VIAL	N050276	003
TETRACYN			
PFIZER	40MG/VIAL;100MG/VIAL	A060285	002
	40MG/VIAL;250MG/VIAL	A060285	003

PROCAINE MERETHOXYLLINE; THEOPHYLLINE

INJECTABLE; INJECTION

DICURIN PROCAINE

LILLY	100MG/ML;50MG/ML	N008869	001
-------	------------------	---------	-----

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPazine

GLAXOSMITHKLINE	2.5MG	N011127	003
	5MG	N011127	001
	25MG	N011127	002
PROCHLORPERAZINE			
ABLE	2.5MG	A040407	001 Jul 11, 2001
	5MG	A040407	002 Jul 11, 2001
	25MG	A040407	003 Jul 11, 2001

PROCHLORPERAZINE EDISYLATE

CONCENTRATE; ORAL

COMPazine

GLAXOSMITHKLINE	EQ 10MG BASE/ML	N011276	001
PROCHLORPERAZINE			
ALPHARMA US PHARMS	EQ 10MG BASE/ML	A087153	001 Jun 08, 1982
PROCHLORPERAZINE EDISYLATE			
MORTON GROVE	EQ 10MG BASE/ML	A088598	001 Oct 25, 1984
INJECTABLE; INJECTION			
COMPazine			
GLAXOSMITHKLINE	EQ 5MG BASE/ML	N010742	002
PROCHLORPERAZINE			
BAXTER HLTHCARE	EQ 5MG BASE/ML	A087759	001 Oct 01, 1982
PROCHLORPERAZINE EDISYLATE			
BAXTER HLTHCARE	EQ 5MG BASE/ML	A089523	001 May 03, 1988
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 - 281 (of 346)

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION					
PROCHLORPERAZINE EDISYLATE					
WATSON LABS	EQ 5MG BASE/ML		A089605	001	Jul 08, 1987
	EQ 5MG BASE/ML		A089606	001	Jul 08, 1987
WYETH AYERST	EQ 5MG BASE/ML		A086348	001	
SYRUP; ORAL					
COMPАЗINE					
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					
COMPАЗINE					
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					
COMPАЗINE					
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					
ABBOTT LABS	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 282 (of 346)

PROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE

WATSON LABS	25MG/ML	A084510	001
	50MG/ML	A084517	001
SPARINE			
BAXTER HLTHCARE CORP	25MG/ML	N010349	008
	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
BIONICHE PHARMA	25MG/ML	A040471	001 Nov 21, 2002
HOSPIRA	50MG/ML	A040372	002 Jun 08, 2000
	50MG/ML	A083838	002
MARSAM PHARMS LLC	25MG/ML	A089463	001 May 02, 1988
	50MG/ML	A089477	001 May 02, 1988
SANDOZ	25MG/ML	A040593	001 Nov 08, 2006
	50MG/ML	A040593	002 Nov 08, 2006
WATSON LABS	25MG/ML	A083532	001
	25MG/ML	A084591	001
	50MG/ML	A080629	002
	50MG/ML	A083532	002

ZIPAN-25

ALTANA	25MG/ML	A083997	001
--------	---------	---------	-----

ZIPAN-50

ALTANA	50MG/ML	A083997	002
--------	---------	---------	-----

SUPPOSITORY; RECTAL

PHENERGAN

SHIONOGI 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

DISCONTINUED DRUG PRODUCT LIST

6 - 283 (of 346)

PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL				
PROMETH PLAIN				
ACTAVIS MID ATLANTIC	6.25MG/5ML	A085953	001	
PROMETHAZINE				
CENCI	6.25MG/5ML	A089013	001	Sep 20, 1985
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				
WYETH PHARMS INC	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
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	
MUTUAL PHARM	12.5MG	A084555	001	
	25MG	A084554	001	
	50MG	A084557	001	
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	
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	
	12.5MG	A085986	001	
	25MG	A083204	001	
	25MG	A085684	001	
	50MG	A083403	001	
	50MG	A085664	001	
REMSED				
BRISTOL MYERS SQUIBB	25MG	A083176	002	
	50MG	A083176	001	

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 - 284 (of 346)

PROPANTHELINE BROMIDE

TABLET; ORAL PRO-BANTHINE SHIRE	15MG	N008732	002	
PROPANTHELINE BROMIDE ASCOT	15MG	A087663	001	Oct 25, 1982
HEATHER	15MG	A085780	001	
IMPAK LABS	15MG	A084541	002	
MYLAN	15MG	A083706	001	
PAR PHARM	15MG	A088377	001	Dec 08, 1983
PVT FORM	15MG	A080977	001	
ROXANE	7.5MG	A080927	001	
SANDOZ	15MG	A080928	001	
TABLICAPS	15MG	A084428	001	
WATSON LABS	15MG	A083029	002	
	15MG	A083151	001	

PROPARACAINe HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC KAINAIR PHARMAFAIR	0.5%	A088087	001	Jun 07, 1983
PARACAINe OPTOPICS	0.5%	A087681	001	Aug 05, 1982
PROPARACAINe HYDROCHLORIDE SOLA BARNES HIND	0.5%	A084144	001	
	0.5%	A084151	001	

PROPIOLACTONE

SOLUTION; IRRIGATION BETAPRONE FOREST LABS	N/A	N011657	001	
--	-----	---------	-----	--

PROPIOMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION LARGON BAXTER HLTHCARE CORP	20MG/ML	N012382	002	
---	---------	---------	-----	--

PROPOFOL

INJECTABLE; INJECTION DIPRIVAN APP PHARMS	10MG/ML	N019627	001	Oct 02, 1989
PROPOFOL BEDFORD	10MG/ML	A074848	001	Apr 19, 2005
TEVA PARENTERAL	10MG/ML	A075392	001	Sep 19, 2000

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	
IMPAK 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

DISCONTINUED DRUG PRODUCT LIST

6 - 285 (of 346)

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL				
PROPOXYPHENE HYDROCHLORIDE				
MYLAN	65MG	A083299	001	
NEXGEN PHARMA INC	65MG	A083185	001	
PAR PHARM	65MG	A080269	001	
PUREPAC PHARM	65MG	A083278	001	
PVT FORM	32MG	A083464	001	
	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	
PROPOXYPHENE HYDROCHLORIDE 65				
WARNER CHILCOTT	65MG	A083786	001	

PROPOXYPHENE 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
CONCENTRATE; ORAL				
PROPRANOLOL HYDROCHLORIDE INTENSOL				
ROXANE	80MG/ML	A071388	001	May 15, 1987
INJECTABLE; INJECTION				
PROPRANOLOL HYDROCHLORIDE				
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				
AKRIMAX PHARMS	10MG	N016418	001	
	20MG	N016418	003	
	90MG	N016418	010	Oct 18, 1982
PROPRANOLOL HYDROCHLORIDE				
CLONMEL HLTHCARE	10MG	A070125	001	Jul 30, 1985
	20MG	A070126	001	Jul 30, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 286 (of 346)

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

CLONMEL HLTHCARE	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
MUTUAL PHARM	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
MYLAN	60MG	A072275	001	Jun 09, 1989
PAR PHARM	90MG	A071288	001	Oct 22, 1986
PLIVA	90MG	A071977	001	Apr 06, 1988
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

DISCONTINUED DRUG PRODUCT LIST

6 - 287 (of 346)

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

SUPERPHARM	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
	80MG	A070442	001	Sep 15, 1986
WATSON LABS	10MG	A070140	001	Jul 30, 1985
	10MG	A070378	001	Mar 19, 1987
	10MG	A070548	001	Jul 10, 1986
	20MG	A070141	001	Jul 30, 1985
	20MG	A070379	001	Mar 19, 1987
	20MG	A070549	001	Apr 11, 1986
	40MG	A070142	001	Jul 30, 1985
	40MG	A070380	001	Mar 19, 1987
	40MG	A070550	001	Apr 11, 1986
	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

PROPYLIODONE

SUSPENSION; INTRATRACHEAL

DIONOSIL AQUEOUS

GLAXOSMITHKLINE 50%

N009309 001

DIONOSIL OILY

GLAXOSMITHKLINE 60%

N009309 002

PROPYLTIOURACIL

TABLET; ORAL

PROPYLTIOURACIL

ABBOTT	50MG	A084075	001
ANABOLIC	50MG	A080285	001
HALSEY	50MG	A080015	001
IMPAK LABS	50MG	A080159	001
IVAX SUB TEVA PHARMS	50MG	A080215	001
LANNETT	50MG	A080016	001
LILLY	50MG	N006213	001
MUTUAL PHARM	50MG	A083982	001
PERRIGO	50MG	A084543	001
TABLICAPS	50MG	A080840	001
WATSON LABS	50MG	A080932	001
	50MG	A085201	001

PROTAMINE SULFATE

INJECTABLE; INJECTION

PROTAMINE SULFATE

BAXTER HLTHCARE	10MG/ML	A089474	001	Nov 05, 1986
	10MG/ML	A089475	001	Nov 05, 1986
LILLY	10MG/ML	N006460	002	
PHARMACIA AND UPJOHN	50MG/VIAL	N007413	001	
	250MG/VIAL	N007413	002	Aug 02, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 288 (of 346)

PROTEIN HYDROLYSATE

INJECTABLE; INJECTION				
AMINOSOL 5%				
ABBOTT	5%		N005932	012
HYPROTIGEN 5%				Jan 31, 1985
B BRAUN	5%		N006170	003
				Jan 10, 1984

PROTIRELIN

INJECTABLE; INJECTION				
THYPINONE				
ABBOTT	0.5MG/ML		N017638	001
THYREL TRH				
FERRING	0.5MG/ML		N018087	001

PROTOKYLOL HYDROCHLORIDE

TABLET; ORAL				
VENTAIRE				
SANOFI AVENTIS US	2MG		A083459	001

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL				
VIVACTIL				
ODYSSEY PHARMS	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
TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES				Jun 17, 1985
KV PHARM	120MG;5MG		A071798	001
SYRUP; ORAL				Mar 16, 1989
ACTAHIST				
CENCI	30MG/5ML;1.25MG/5ML		A088344	001
HISTAFED				Feb 09, 1984
CENCI	30MG/5ML;1.25MG/5ML		A088283	001
MYFED				Apr 20, 1984
USL PHARMA	30MG/5ML;1.25MG/5ML		A088116	001
TRILITRON				Mar 04, 1983
NEWTRON PHARMS	30MG/5ML;1.25MG/5ML		A088474	001
TABLET; ORAL				Feb 12, 1985
ALLERFED				
PVT FORM	60MG;2.5MG		A088860	001
COPHED				Jan 31, 1985
SANDOZ	60MG;2.5MG		A088602	001
PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE				
SANDOZ	60MG;2.5MG		A088193	001
TRILITRON				May 17, 1983
NEWTRON PHARMS	60MG;2.5MG		A088515	001
TRIPHED				Jan 09, 1985
TEVA	60MG;2.5MG		A088630	001
TRIPROLIDINE AND PSEUDOEPHEDRINE				May 17, 1984
WATSON LABS	60MG;2.5MG		A088318	002
WEST WARD	60MG;2.5MG		A088117	001
TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE				Jan 13, 1984
IVAX SUB TEVA PHARMS	60MG;2.5MG		A085273	001
				Apr 19, 1983
				Dec 12, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 289 (of 346)

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE 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 BARR	30MG	A040512	002	Jul 20, 2005
	60MG	A040512	001	Oct 08, 2003
SOLVAY	30MG	A089572	001	Nov 27, 1990
US ARMY	30MG	N020414	001	Feb 05, 2003

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 & 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 QUESTCOR PHARMS	7.5MG	N018708	003	Feb 26, 1987
--	-------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 290 (of 346)

QUETIAPINE FUMARATE

TABLET; ORAL SEROQUEL	ASTRAZENECA	EQ 150MG BASE	N020639 004	Dec 20, 1998
--------------------------	-------------	---------------	-------------	--------------

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL QUINAPRIL HYDROCHLORIDE	ACTAVIS TOTOWA	EQ 5MG BASE EQ 10MG BASE EQ 20MG BASE EQ 40MG BASE	A076459 001 A076459 002 A076459 003 A076459 004	Dec 22, 2004 Dec 22, 2004 Dec 22, 2004 Dec 22, 2004
---	----------------	---	--	--

QUINESTROL

TABLET; ORAL ESTROVIS	PARKE DAVIS	0.1MG 0.2MG	N016768 002 N016768 003	
--------------------------	-------------	----------------	----------------------------	--

QUINETHAZONE

TABLET; ORAL HYDROMOX	LEDERLE	50MG	N013264 001	
--------------------------	---------	------	-------------	--

QUINETHAZONE; RESERPINE

TABLET; ORAL HYDROMOX R	LEDERLE	50MG; 0.125MG	N013927 001	
----------------------------	---------	---------------	-------------	--

QUINIDINE GLUCONATE

TABLET; ORAL QUINACT	BAYER HLTHCARE	266MG 400MG	A085978 001 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
	HALSEY	324MG	A089476 001	Apr 10, 1987
	ROXANE	324MG	A088431 001	Jan 06, 1984
	SANDOZ	324MG	A089894 001	Dec 15, 1988
	SUPERPHARM	324MG	A089164 001	Nov 21, 1985
	WATSON LABS	324MG	A087785 001	Jan 24, 1983

QUINIDINE POLYGALACTURONATE

TABLET; ORAL CARDIOQUIN	PHARM RES ASSOC	275MG	N011642 002	
----------------------------	-----------------	-------	-------------	--

QUINIDINE SULFATE

CAPSULE; ORAL CIN-QUIN	SOLVAY	200MG 300MG	A085296 001 A085297 001	
QUINIDINE SULFATE	LILLY	200MG	A085103 001	

DISCONTINUED DRUG PRODUCT LIST

6 - 291 (of 346)

QUINIDINE SULFATE

TABLET; ORAL				
CIN-QUIN				
SOLVAY	100MG	A085299	001	
	200MG	A084932	001	
	300MG	A085298	001	
QUINIDINE SULFATE				
BARR	200MG	A084177	001	
CLONMEL HLTHCARE	200MG	A087011	001	
CONTRACT PHARMACAL	200MG	A083808	001	
ELKINS SINK	200MG	A083622	001	
EVERYLIFE	200MG	A083439	001	
HALSEY	200MG	A083583	001	
IMPAX 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	
MUTUAL PHARM	100MG	A081029	001	Apr 14, 1989
PERRIGO	200MG	A085322	001	
PHARMAVITE	200MG	A084627	001	
PUREPAC PHARM	200MG	A084003	001	
ROXANE	200MG	A083640	001	
	300MG	A085632	001	
SANDOZ	200MG	A084631	001	
	200MG	A084914	001	
	300MG	A089839	001	Sep 29, 1988
SCHERER LABS	200MG	A085068	001	
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	
WEST WARD	200MG	A083862	001	
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 SODIUM

TABLET, DELAYED RELEASE; ORAL				
ACIPHEX				
EISAI INC	10MG	N020973	001	May 29, 2002

RAMIPRIL

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

DISCONTINUED DRUG PRODUCT LIST

6 - 292 (of 346)

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
 GLAXOSMITHKLINE 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
 GLAXOSMITHKLINE EQ 50MG BASE/100ML N019593 001 Dec 17, 1986
 TABLET; ORAL
 RANITIDINE HYDROCHLORIDE
 BOEHRINGER INGELHEIM EQ 150MG BASE A074662 001 Aug 29, 1997
 EQ 300MG BASE A074662 002 Aug 29, 1997
 MYLAN EQ 150MG BASE A074552 001 Jul 30, 1998
 EQ 300MG BASE A074552 002 Jul 30, 1998
 RANBAXY EQ 75MG BASE A075132 001 Jan 14, 2000
 EQ 75MG BASE A075254 001 Jan 14, 2000
 EQ 150MG BASE A075000 001 Jan 30, 1998
 EQ 150MG BASE A075439 001 Apr 19, 2000
 EQ 300MG BASE A075000 002 Jan 30, 1998
 EQ 300MG BASE A075439 002 Apr 19, 2000
 SANDOZ EQ 75MG BASE A075519 001 Sep 26, 2002
 TABLET, EFFERVESCENT; ORAL
 ZANTAC 150
 GLAXOSMITHKLINE EQ 150MG BASE N020251 001 Mar 31, 1994
 ZANTAC 75
 BOEHRINGER INGELHEIM EQ 75MG BASE N020745 001 Feb 26, 1998

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

DISCONTINUED DRUG PRODUCT LIST

6 - 293 (of 346)

RAUWOLFIA SERPENTINA ROOT

TABLET; ORAL RAUSERPIN FERNDALE LABS	50MG 100MG	N009926 002 N009926 004
RAUVAL PAL PAK	50MG 100MG	N009108 002 N009108 004
RAUWOLFIA SERPENTINA BUNDY	50MG 100MG	N009477 001 N009477 002
HALSEY	50MG 100MG	A080498 001 A080498 002
IMPAX LABS	50MG 100MG	N009273 001 N009273 002
IVAX SUB TEVA PHARMS	50MG 100MG	N011521 001 N011521 002
PUREPAC PHARM	50MG 100MG	A080842 001 A080842 002
PVT FORM	50MG 100MG	A080583 001 A080583 002
SOLVAY	50MG 100MG	A080500 001 A080500 002
TABLICAPS	50MG 100MG	A083867 001 A083444 001
VALEANT PHARM INTL	50MG 100MG	N009668 001 N009668 002
WATSON LABS	50MG 100MG	A080907 001 A080914 001
WOLFINA FOREST PHARMS	50MG 100MG	N009255 008 N009255 006

RESCINNAMINE

CAPSULE; ORAL CINNASIL PANRAY	0 .5MG	A084736 001
TABLET; ORAL MODERIL PFIZER	0 .25MG 0 .5MG	N010686 003 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 0 .25MG	N009631 002 N009631 004
RAU-SED BRISTOL MYERS SQUIBB	0 .1MG 0 .25MG 0 .5MG 1MG	N009357 001 N009357 004 N009357 006 N009357 008
RESERPINE BARR BELL PHARMA	0 .25MG 0 .1MG 0 .25MG	A080721 002 A083058 001 A083058 002

DISCONTINUED DRUG PRODUCT LIST

6 - 294 (of 346)

RESERPINE

TABLET; ORAL

RESERPINE

BUNDY	0.1MG	N009663 001
	0.25MG	N009663 003
ELKINS SINK	0.1MG	A083145 001
	0.25MG	A083145 002
EVERTYLIFE	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
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
ROXANE	0.1MG	N009859 001
	0.25MG	N009859 002
SOLVAY	0.25MG	A080446 001
TABLICAPS	0.25MG	A085207 001
TEVA	0.1MG	A089020 001
	0.25MG	A089019 001
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
WEST WARD	0.1MG	A080975 001
	0.25MG	A080975 002
	1MG	A080975 003
WHITEWORTH TOWN PLSN	0.1MG	A080723 001
	0.25MG	A080723 002
	1MG	A080723 003
SANDRIL		
LILLY	0.1MG	N009376 004
	0.25MG	N009376 001
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

DISCONTINUED DRUG PRODUCT LIST

6 - 295 (of 346)

RESERPINE

TABLET; ORAL			
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

TABLET; ORAL			
COPEGUS			
ROCHE	400MG		N021511 002 Jun 21, 2005

RIFAMPIN

CAPSULE; ORAL			
RIMACTANE			
PROSAM LABS	300MG		N050429 001

RIMANTADINE HYDROCHLORIDE

SYRUP; ORAL			
FLUMADINE			
FOREST LABS	50MG/5ML		N019650 001 Sep 17, 1993
TABLET; ORAL			
RIMANTADINE HYDROCHLORIDE			
ACTAVIS TOTOWA	100MG		A076375 001 Jan 14, 2003

RISEDRONATE SODIUM

TABLET; ORAL			
ACTONEL			
WARNER CHILCOTT	75MG		N020835 004 Apr 16, 2007

RISPERIDONE

TABLET; ORAL			
RISPERDAL			
JANSSEN PHARMS	5MG		N020272 005 Dec 29, 1993
RISPERIDONE			
ACTAVIS TOTOWA	0 . 25MG		A078071 001 Jun 17, 2009
	0 . 5MG		A078071 002 Jun 17, 2009
	1MG		A078071 003 Jun 17, 2009
	2MG		A078071 004 Jun 17, 2009
	3MG		A078071 005 Jun 17, 2009
	4MG		A078071 006 Jun 17, 2009
CADISTA PHARMS	0 . 25MG		A078828 001 Mar 23, 2009
	0 . 5MG		A078828 002 Mar 23, 2009
	1MG		A078828 003 Mar 23, 2009
	2MG		A078828 004 Mar 23, 2009
	3MG		A078828 005 Mar 23, 2009
	4MG		A078828 006 Mar 23, 2009
RATIOPHARM	0 . 25MG		A077784 001 Jun 08, 2010
	0 . 5MG		A077784 002 Jun 08, 2010
	1MG		A077784 003 Jun 08, 2010
	2MG		A077784 004 Jun 08, 2010
	3MG		A077784 005 Jun 08, 2010
	4MG		A077784 006 Jun 08, 2010
SYNTHON PHARMS	0 . 25MG		A078187 001 Oct 22, 2009

DISCONTINUED DRUG PRODUCT LIST

6 - 296 (of 346)

RISPERIDONE

TABLET; ORAL					
RISPERIDONE					
SYNTHON PHARMS	0 .5MG		A078187	002	Oct 22, 2009
	1MG		A078187	003	Oct 22, 2009
	2MG		A078187	004	Oct 22, 2009
	3MG		A078187	005	Oct 22, 2009
	4MG		A078187	006	Oct 22, 2009

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION					
RITODRINE HYDROCHLORIDE					
ABRAXIS PHARM	10MG/ML		A071188	001	Jul 23, 1987
	15MG/ML		A071189	001	Jul 23, 1987
HOSPIRA	10MG/ML		A071618	001	Feb 28, 1991
	15MG/ML		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		N018580	001	
	15MG/ML		N018580	002	
TABLET; ORAL					
YUTOPAR					
ASTRAZENECA	10MG		N018555	001	

RITONAVIR

CAPSULE; ORAL					
NORVIR					
ABBOTT	100MG		N020680	001	Mar 01, 1996

ROCURONIUM BROMIDE

INJECTABLE; INJECTION					
ZEMURON					
SCHERING	10MG/ML (10MG/ML)		N020214	002	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, EXTENDED RELEASE; ORAL					
REQUIP XL					
SMITHKLINE BEECHAM	EQ 3MG BASE		N022008	002	Jun 13, 2008

ROSE BENGAL SODIUM I-131

INJECTABLE; INJECTION					
ROBENGATOP					
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 297 (of 346)

ROTIGOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL NEUPRO			
UCB INC	2MG/24HR	N021829	001 May 09, 2007
	4MG/24HR	N021829	002 May 09, 2007
	6MG/24HR	N021829	003 May 09, 2007

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

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

DISCONTINUED DRUG PRODUCT LIST

6 - 298 (of 346)

SECOBARBITAL SODIUM

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

AAIPHARMA LLC 5MG A075145 001 Sep 15, 2003

TABLET; ORAL
SELEGILINE HYDROCHLORIDE

ENDO PHARMS	5MG	A074565	001	Aug 02, 1996
IVAX SUB TEVA PHARMS	5MG	A074756	001	Nov 25, 1998
SIEGFRIED	5MG	A074672	001	Apr 01, 1997
SOMERSET	5MG	N019334	001	Jun 05, 1989
TEVA	5MG	A074537	001	Aug 02, 1996
	5MG	A074744	001	Jan 27, 1997

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL
EXSEL

ALLERGAN HERBERT	2.5%	A083892	001
SELENIUM SULFIDE			
ACTAVIS MID ATLANTIC	2.5%	A084394	001
IVAX PHARMS	2.5%	A085777	001
TARO	2.5%	A086209	001

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

SERACTIDE ACETATE

INJECTABLE; INJECTION
ACTHAR GEL-SYNTHETIC

ARMOUR PHARM	40 UNITS/ML	N017861	001
	80 UNITS/ML	N017861	002

SERMORELIN ACETATE

INJECTABLE; INJECTION
GEREF

EMD SERONO	EQ 0.05MG BASE/AMP	N019863	001	Dec 28, 1990
	EQ 0.5MG BASE/VIAL	N020443	001	Sep 26, 1997
	EQ 1MG BASE/VIAL	N020443	002	Sep 26, 1997

DISCONTINUED DRUG PRODUCT LIST

6 - 299 (of 346)

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL SERTRALINE HYDROCHLORIDE					
ROXANE	EQ 20MG BASE/ML		A076934	001	Jun 30, 2006
TABLET; ORAL SERTRALINE HYDROCHLORIDE					
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
ACTAVIS TOTOWA	EQ 25MG BASE		A078175	001	Jul 21, 2010
	EQ 50MG BASE		A078175	002	Jul 21, 2010
	EQ 100MG BASE		A078175	003	Jul 21, 2010
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
MUTUAL PHARM	EQ 25MG BASE		A077818	001	Feb 06, 2007
	EQ 50MG BASE		A077818	002	Feb 06, 2007
	EQ 100MG BASE		A077818	003	Feb 06, 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
ROXANE	EQ 25MG BASE		A076881	001	Feb 06, 2007
	EQ 50MG BASE		A076881	002	Feb 06, 2007
	EQ 100MG BASE		A076881	003	Feb 06, 2007
SANDOZ	EQ 25MG BASE		A077713	001	Feb 06, 2007
	EQ 50MG BASE		A077713	002	Feb 06, 2007
	EQ 100MG BASE		A077713	003	Feb 06, 2007
WATSON LABS	EQ 25MG BASE		A077162	001	Feb 06, 2007
	EQ 50MG BASE		A077162	002	Feb 06, 2007
	EQ 100MG BASE		A077162	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		N020632	001	Nov 22, 1997
	10MG		N020632	002	Nov 22, 1997
	15MG		N020632	003	Nov 22, 1997

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					
ACTAVIS TOTOWA	5MG		A078735	001	Aug 30, 2010
	10MG		A078735	002	Aug 30, 2010
	20MG		A078735	003	Aug 30, 2010
	40MG		A078735	004	Aug 30, 2010

DISCONTINUED DRUG PRODUCT LIST

6 - 300 (of 346)

SIMVASTATIN

TABLET; ORAL SIMVASTATIN					
ACTAVIS TOTOWA	80MG	A078735	005	Aug 30, 2010	
SANDOZ INC	5MG	A077766	001	Dec 20, 2006	
	10MG	A077766	002	Dec 20, 2006	
	20MG	A077766	003	Dec 20, 2006	
	40MG	A077766	004	Dec 20, 2006	
	80MG	A077766	005	Dec 20, 2006	
TABLET, ORALLY DISINTEGRATING; ORAL SIMVASTATIN					
SYNTTHON PHARMS	10MG	N021961	001	Oct 09, 2007	
	20MG	N021961	002	Oct 09, 2007	
	40MG	N021961	003	Oct 09, 2007	
	80MG	N021961	004	Oct 09, 2007	

SIROLIMUS

TABLET; ORAL RAPAMUNE					
WYETH PHARMS INC	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 LLC	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% IN PLASTIC CONTAINER					
ABBOTT	9MG/ML	N019218	001	Jul 13, 1984	
MILES	900MG/100ML	N018502	001		
SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER					
ABRAXIS PHARM	234MG/ML	N019329	001	Apr 22, 1987	
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER					
B BRAUN	3GM/100ML	N019635	003	Mar 09, 1988	
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER					
B BRAUN	5GM/100ML	N019635	004	Mar 09, 1988	
SOLUTION; IRRIGATION					
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER					
BAXTER HLTHCARE	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		

DISCONTINUED DRUG PRODUCT LIST

6 - 301 (of 346)

SODIUM CHROMATE CR-51

INJECTABLE; INJECTION CHROMITOPE SODIUM		
BRACCO	2mCi/VIAL	N013993 002
SODIUM CHROMATE CR 51		
MALLINCKRODT	100uCi/ML	N016708 001

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 IODIDE I-123

CAPSULE; ORAL SODIUM IODIDE I 123		
SYNCOR PHARMS	400uCi	N018671 003 May 27, 1982

SODIUM IODIDE I-131

CAPSULE; ORAL IODOTOPE		
BRACCO	1-130mCi	N010929 001
	1-150mCi	N010929 003
SODIUM IODIDE I 131		
CIS	50uCi	N017316 001
	100uCi	N017316 002
MALLINCKRODT	0.8-100mCi	N016515 002
	15-100uCi	N016517 002
SOLUTION; ORAL 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
HOSPIRA	1.87GM/100ML	N018249 001

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
	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 - 302 (of 346)

SODIUM PHOSPHATE P-32

SOLUTION; INJECTION, ORAL PHOSPHOTOPE		
BRACCO	1-8mCi/VIAL	N010927 001
SODIUM PHOSPHATE P 32		
MALLINCKRODT	0.67mCi/ML	N011777 001
	1.5mCi/VIAL	N011777 002

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL SODIUM POLYSTYRENE SULFONATE		
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 SINN	30%	A080516 001

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION SOTRADECOL		
ELKINS SINN	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	N019107 001 Oct 17, 1985
	10MG/VIAL	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		
CANGENE	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	N019721 001 May 08, 1995
	8MG/VIAL	N019721 002 May 08, 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

DISCONTINUED DRUG PRODUCT LIST

6 - 303 (of 346)

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION SAIZEN					
EMD SERONO	4MG/VIAL 6MG/VIAL		N019764	005	Jan 16, 2007
SEROSTIM	8.8MG/VIAL		N019764	001	Oct 08, 1996
ZORBTIVE					
EMD SERONO	4MG/VIAL 5MG/VIAL 6MG/VIAL		N021597	001	Dec 01, 2003
			N021597	002	Dec 01, 2003
			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			N018512	001	May 27, 1982
---	--	--	---------	-----	--------------

SOTALOL HYDROCHLORIDE

TABLET; ORAL BETAPACE					
BAYER HLTHCARE	320MG		N019865	004	Oct 30, 1992
BETAPACE AF					
BAYER HLTHCARE	40MG 60MG 100MG		N021151	006	Apr 02, 2003
SOTALOL HYDROCHLORIDE			N021151	007	Apr 02, 2003
MUTUAL PHARM	80MG 80MG 120MG 120MG 160MG 160MG 240MG		N021151	005	Mar 14, 2003
WATSON LABS			A075515	001	Oct 15, 2001
	80MG 120MG 160MG 240MG		A076576	001	Apr 08, 2004
			A075515	004	Oct 15, 2001
			A076576	002	Apr 08, 2004
			A075515	002	Oct 15, 2001
			A076576	003	Apr 08, 2004
			A075515	003	Oct 15, 2001
			A075238	001	Jul 13, 2000
			A075238	002	Jul 13, 2000
			A075238	003	Jul 13, 2000
			A075238	004	Jul 13, 2000

SOYBEAN OIL

INJECTABLE; INJECTION 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	

DISCONTINUED DRUG PRODUCT LIST

6 - 304 (of 346)

SPIRAPRIL HYDROCHLORIDE

TABLET; ORAL RENORMAX SCHERING	3MG 6MG 12MG 24MG	N020240 001 N020240 002 N020240 003 N020240 004	Dec 29, 1994 Dec 29, 1994 Dec 29, 1994 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 ZERIT BRISTOL MYERS SQUIBB	5MG	N020412 001	Jun 24, 1994
CAPSULE, EXTENDED RELEASE; ORAL ZERIT XR BRISTOL MYERS SQUIBB	37.5MG	N021453 001	Dec 31, 2002
	50MG	N021453 002	Dec 31, 2002
	75MG	N021453 003	Dec 31, 2002
	100MG	N021453 004	Dec 31, 2002

STERILE WATER FOR INJECTION

LIQUID; N/A BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER ABRAXIS PHARM	100%	A089099 001	Dec 29, 1987
	100%	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	A060684 001	
LILLY	EQ 1GM BASE/VIAL	A060107 001	
	EQ 1GM BASE/2ML	A060404 001	
	EQ 5GM BASE/VIAL	A060107 002	
PFIZER	EQ 1GM BASE/VIAL	A060076 001	
	EQ 1GM BASE/2.5ML	A060111 001	

DISCONTINUED DRUG PRODUCT LIST

6 - 305 (of 346)

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION
 STREPTOMYCIN SULFATE
 PFIZER EQ 5GM BASE/VIAL A060076 002

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION
 ANECTINE
 SANDOZ 50MG/ML N008453 003
 500MG/VIAL N008453 001
 1GM/VIAL N008453 004
 QUELICIN PRESERVATIVE FREE
 HOSPIRA 50MG/ML N008845 002
 SUCCINYLCHOLINE CHLORIDE
 INTL MEDICATION 100MG/VIAL A085400 001 Feb 04, 1982
 ORGANON USA INC 20MG/ML A080997 001
 SUCOSTRIN
 APOTHECON 20MG/ML N008847 001
 100MG/ML 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
 SODIUM SULAMYD
 SCHERING 10% N005963 002
 SULFAIR 10
 PHARMAFAIR 10% A088000 001 Dec 22, 1982
 SOLUTION/DROPS; OPHTHALMIC
 BLEPH-30
 ALLERGAN 30% A080028 002
 ISOPTO CETAMIDE
 ALCON 15% A080020 002
 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 10% A040215 001 May 25, 1999
 30% A040216 001 May 25, 1999
 ALCON PHARMS LTD 30% A089068 001 May 05, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 306 (of 346)

SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC SULFACETAMIDE SODIUM				
PHARMAFAIR 10% SULFAIR 10% PHARMAFAIR 10% SULFAIR FORTE 30% SULFAIR-15 15% PHARMAFAIR 15% SULTEN-10 10%			A088947 001 A087949 001 A088385 001 A088186 001 A087818 001	May 17, 1985 Dec 13, 1982 Oct 13, 1983 May 25, 1983 Feb 03, 1983
BAUSCH AND LOMB	10%			

SULFACYTINE

TABLET; ORAL RENOQUID GLENWOOD	250MG	N017569 001
--------------------------------------	-------	-------------

SULFADIAZINE

TABLET; ORAL SULFADIAZINE		
ABBOTT 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

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

DISCONTINUED DRUG PRODUCT LIST

6 - 307 (of 346)

SULFAMETHOXAZOLE

TABLET; ORAL

SULFAMETHOXAZOLE

ASCOT	500MG	A087662	001	Oct 20, 1982
BARR	500MG	A087189	001	Jul 25, 1983
HEATHER	500MG	A086163	001	
SANDOZ	500MG	A085844	001	
WATSON LABS	500MG	A085053	001	
	1GM	A086000	001	
UROBAK				
SHIONOGI	500MG	A087307	001	

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

BACTRIM

MUTUAL PHARM	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
BAXTER HLTHCARE	80MG/ML;16MG/ML	A070627	001	Dec 29, 1987
	80MG/ML;16MG/ML	A070628	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

SUSPENSION; ORAL

BACTRIM

MUTUAL PHARM	200MG/5ML;40MG/5ML	N017560	001	
BACTRIM PEDIATRIC				
MUTUAL PHARM	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				
TEVA	200MG/5ML;40MG/5ML	A070028	001	Jun 02, 1987
	200MG/5ML;40MG/5ML	N018812	001	Jan 28, 1983
	200MG/5ML;40MG/5ML	N018812	002	Jun 10, 1983
TEVA PHARMS	200MG/5ML;40MG/5ML	A077612	001	Nov 13, 2006

SULFATRIM

ACTAVIS MID ATLANTIC	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
COTRIM D.S.				
TEVA	800MG;160MG	A070048	001	Mar 18, 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
	400MG;80MG	A071016	001	Aug 25, 1986
PLIVA	400MG;80MG	A070215	001	Sep 10, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 308 (of 346)

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

PLIVA	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
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

TEVA 15%

A088718 001 Sep 19, 1985

SUPPOSITORY; VAGINAL

AVC

AZUR PHARMA 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

SULFASALAZINE

HERITAGE PHARMS INC 500MG

A080197 001

MUTUAL PHARM 500MG

A089590 001 Oct 19, 1987

SANDOZ 500MG

A086184 001

SUPERPHARM 500MG

A089339 001 Oct 26, 1987

WATSON LABS 500MG

A084964 001

DISCONTINUED DRUG PRODUCT LIST

6 - 309 (of 346)

SULFASALAZINE

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 BARR 500MG	A084031 001
HEATHER 500MG	A080189 001
IMPAX LABS 500MG	A080109 001
IVAX SUB TEVA PHARMS 500MG	A080142 001
LANNETT 500MG	A080085 001
LEDERLE 500MG	A087649 001
PHARMERAL 500MG	A084385 001
PUREPAC PHARM 500MG	A080087 001
ROXANE 500MG	A080082 001
SANDOZ 500MG	A085628 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 - 310 (of 346)

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		
ABBOTT	165MG	N006044 003

SULFUR

POWDER; TOPICAL BENSULFOID		
POYTHRESS	33.32%	N002918 001

SULINDAC

TABLET; ORAL CLINORIL		
MERCK	150MG	N017911 001
SULINDAC		
HERITAGE PHARMS INC	150MG	A073262 002 Sep 06, 1991
	200MG	A073262 001 Sep 06, 1991
SANDOZ	150MG	A072712 001 Aug 30, 1991
	200MG	A072713 001 Aug 30, 1991
TEVA	150MG	A072972 001 Feb 28, 1992
	200MG	A072973 001 Feb 28, 1992

SUMATRIPTAN

SPRAY; NASAL IMITREX		
GLAXOSMITHKLINE	10MG/SPRAY	N020626 002 Aug 26, 1997

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS SUMATRIPTAN SUCCINATE		
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
TABLET; ORAL SUMATRIPTAN SUCCINATE		
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

SUPROFEN

SOLUTION/DROPS; OPHTHALMIC PROFENAL		
ALCON	1%	N019387 001 Dec 23, 1988

SUTILAINS

OINTMENT; TOPICAL TRAVASE		
ABBOTT	82,000 UNITS/GM	N012828 001

DISCONTINUED DRUG PRODUCT LIST

6 - 311 (of 346)

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	

TALBUTAL

TABLET; ORAL LOTUSATE					
SANOFI AVENTIS US	120MG	N009410	005		

TAMOXIFEN CITRATE

SOLUTION; ORAL SOLTAMOX					
ROSEMONT	EQ 10MG BASE/5ML	N021807	001	Oct 29, 2005	
TABLET; ORAL NOLVADEX					
ASTRAZENECA	EQ 10MG BASE	N017970	001		
	EQ 20MG BASE	N017970	002	Mar 21, 1994	
TAMOXIFEN CITRATE					
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		
TECHNECAN 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	

TECHNETIUM TC-99M ALBUMIN KIT

INJECTABLE; INJECTION TECHNETIUM TC 99M HSA					
GE HEALTHCARE	N/A	N017775	001		

DISCONTINUED DRUG PRODUCT LIST

6 - 312 (of 346)

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 LIDOFENIN KIT

INJECTABLE; INJECTION
TECHNESCAN HIDA
DRAXIMAGE N/A N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION
DRAXIMAGE MDP-10
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
MPI DTPA KIT - CHELATE
GE HEALTHCARE N/A N017255 001

DISCONTINUED DRUG PRODUCT LIST

6 - 313 (of 346)

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION
 TECHNETIUM TC-99M PENTETATE KIT
 GE HEALTHCARE N/A N017264 002

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
 TECHNETIUM TC 99M GENERATOR
 GE HEALTHCARE 830-16600mCi/GENERATOR N017693 001

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 - 314 (of 346)

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE; INJECTION
 CARDIOTEC
 BRACCO N/A N019928 001 Dec 19, 1990

TEGASEROD MALEATE

TABLET; ORAL
 ZELNORM
 NOVARTIS EQ 2MG BASE N021200 001 Jul 24, 2002
 EQ 6MG BASE N021200 002 Jul 24, 2002

TELBIVUDINE

SOLUTION; ORAL
 TYZEKA
 NOVARTIS 100MG/5ML N022154 001 Apr 28, 2009

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
 MUTUAL PHARM 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
 30MG A070384 001 Mar 23, 1987

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL
 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
 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
 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 1MG BASE A074657 001 Apr 28, 2000
 EQ 2MG BASE A074315 002 Dec 31, 1998
 EQ 2MG BASE A074657 002 Apr 28, 2000
 EQ 5MG BASE A074315 003 Dec 31, 1998
 EQ 5MG BASE A074657 003 Apr 28, 2000
 EQ 10MG BASE A074315 004 Dec 31, 1998
 EQ 10MG BASE A074657 004 Apr 28, 2000
 TEVA EQ 1MG BASE A074446 001 May 18, 2000
 EQ 2MG BASE A074446 002 May 18, 2000
 EQ 5MG BASE A074446 003 May 18, 2000
 EQ 10MG BASE A074446 004 May 18, 2000

DISCONTINUED DRUG PRODUCT LIST

6 - 315 (of 346)

TERBINAFINE

GEL; TOPICAL LAMISIL NOVARTIS	1%	N020846 001	Apr 29, 1998
-------------------------------------	----	-------------	--------------

TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL LAMISIL NOVARTIS	1%	N020192 001	Dec 30, 1992
SOLUTION; TOPICAL LAMISIL NOVARTIS	1%	N020749 001	Oct 17, 1997
TABLET; ORAL TERBINAFINE HYDROCHLORIDE GEDEON RICHTER USA ROXANE	EQ 250MG BASE EQ 250MG BASE	A077065 001 A077223 001	Jul 02, 2007 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 AAIPHARMA LLC	1MG/ML	N018571 001	
BRICANYL SANOFI AVENTIS US	1MG/ML	N017466 001	
TABLET; ORAL BRETHINE LEHIGH VALLEY	2.5MG 5MG	N017849 001 N017849 002	
BRICANYL SANOFI AVENTIS US	2.5MG 5MG	N017618 001 N017618 002	

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
---	------------------------	-------------	--------------

TESTOLACTONE

INJECTABLE; INJECTION TESLAC BRISTOL MYERS SQUIBB	100MG/ML	N016119 001	
TABLET; ORAL TESLAC BRISTOL MYERS SQUIBB	50MG 250MG	N016118 001 N016118 002	

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL ANDRODERM WATSON LABS	2.5MG/24HR 5MG/24HR	N020489 001 N020489 002	Sep 29, 1995 May 02, 1997
TESTODERM ALZA	4MG/24HR 6MG/24HR	N019762 001 N019762 002	Oct 12, 1993 Oct 12, 1993

DISCONTINUED DRUG PRODUCT LIST

6 - 316 (of 346)

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL TESTODERM TTS ALZA	5MG/24HR	N020791	001	Dec 18, 1997
GEL; TRANSDERMAL TESTOSTERONE PAR PHARM	1% (2.5GM/PACKET) 1% (5GM/PACKET)	A076744	001	May 23, 2007
WATSON LABS	1% (2.5GM/PACKET) 1% (5GM/PACKET)	A076737	001	Jan 27, 2006
		A076737	002	Jan 27, 2006
INJECTABLE; INJECTION TESTOSTERONE WATSON LABS	25MG/ML 50MG/ML 100MG/ML	A086420	001	May 10, 1983
		A086419	001	Aug 23, 1983
		A086417	001	Jul 07, 1983

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION DEPO-TESTOSTERONE PHARMACIA AND UPJOHN	50MG/ML	A085635	001
TESTOSTERONE CYPIONATE WATSON LABS	100MG/ML 100MG/ML 200MG/ML	A084401	001
		A086029	001
		A084401	002

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION DELATESTRYL ENDO PHARM	200MG/ML	N009165	001
TESTOSTERONE ENANTHATE WATSON LABS	100MG/ML 100MG/ML 200MG/ML	A083667	001
		A085599	001
		A083667	002

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION TESTOSTERONE PROPIONATE BEL MAR	25MG/ML 50MG/ML 100MG/ML	A080741	001
ELKINS SINK	25MG/ML	A080276	001
LILLY	50MG/ML	A080254	002
WATSON LABS	25MG/ML 25MG/ML 50MG/ML 50MG/ML 100MG/ML 100MG/ML	A080188	001
		A085490	001
		A080188	002
		A085490	002
		A080188	003
		A083595	003

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL ACHROMYCIN V HERITAGE PHARMS INC	250MG 500MG	N050278	003
BRISTACYCLINE BRISTOL	250MG 250MG 500MG 500MG	A061658	001
		A061888	001
		A061658	002
		A061888	002
CYCLOPAR WARNER CHILCOTT	250MG 250MG 250MG	A061725	001
		A062175	001
		A062332	001

DISCONTINUED DRUG PRODUCT LIST

6 - 317 (of 346)

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL				
CYCLOPAR				
WARNER CHILCOTT	500MG	A061725	002	
	500MG	A062332	002	
PANMYCIN				
PHARMACIA AND UPJOHN	250MG	A060347	001	
RETEF				
SOLVAY	250MG	A061443	001	
	500MG	A061443	002	
ROBITET				
WYETH AYERST	250MG	A061734	001	
	500MG	A061734	002	
SUMYCIN				
APOTHECON	100MG	A060429	002	
	125MG	A060429	004	
	250MG	A060429	001	
	500MG	A060429	003	
TETRACHEL				
ANGUS	250MG	A060343	001	
	500MG	A060343	003	
TETRACYCLINE HYDROCHLORIDE				
ABBOTT	250MG	A061802	001	
	500MG	A061802	002	
ELKINS SINK	250MG	A060059	001	
FERRANTE	125MG	A060173	001	
	250MG	A060173	002	
HEATHER	250MG	A061148	001	
	500MG	A061148	002	
LABS ATRAL	250MG	A062752	001	Aug 12, 1988
	500MG	A062752	002	Aug 12, 1988
MAST MM	250MG	A062085	001	
MUTUAL PHARM	250MG	A060736	001	
	500MG	A060736	002	
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	
SANDOZ	250MG	A061471	001	
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	
WEST WARD	250MG	A060768	001	
	500MG	A060768	002	
WYETH AYERST	250MG	A061685	001	
	500MG	A061685	002	
TETRACYN				
PFIPHARMECS	250MG	A060082	003	
	500MG	A060082	004	
FIBER, EXTENDED RELEASE; PERIODONTAL				
ACTISITE				
SCHIFF AND CO	12.7MG/FIBER	N050653	001	Mar 25, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 318 (of 346)

TETRACYCLINE HYDROCHLORIDE

FOR SOLUTION; TOPICAL TOPICYCLINE			
SHIRE	2.2MG/ML	N050493	001
INJECTABLE; INJECTION ACHROMYCIN			
LEDERLE	250MG/VIAL	N050273	002
	500MG/VIAL	N050273	003
TETRACYN			
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
TETRACYN			
PFIPHARMECS	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

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE; ORAL TETREX			
BRISTOL	EQ 100MG HCL	A061653	001
	EQ 250MG HCL	A061653	002
	EQ 250MG HCL	A061889	002
	EQ 250MG HCL	N050212	002
	EQ 500MG HCL	A061653	003
	EQ 500MG HCL	A061889	001
	EQ 500MG HCL	N050212	003

THALLOUS 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

THEOPHYLLINE

CAPSULE; ORAL BRONKODYL			
SANOFI AVENTIS US	100MG	A085264	001
	200MG	A085264	002

DISCONTINUED DRUG PRODUCT LIST

6 - 319 (of 346)

THEOPHYLLINE

CAPSULE; ORAL ELIXOPHYLLIN FOREST LABS	100MG 200MG	A085545 001 A083921 001	Jul 31, 1984 Jul 31, 1984
SOMOPHYLLIN-T FISONS	100MG 200MG 250MG	A087155 001 A087155 002 A087155 003	Feb 25, 1985 Feb 25, 1985 Feb 25, 1985
THEOPHYLLINE KV PHARM	100MG 200MG	A085263 001 A085263 002	
SCHERER RP	100MG 200MG 250MG	A084731 002 A084731 001 A084731 003	Nov 07, 1986 Nov 07, 1986 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 250MG	A086826 001 A086826 002	Jan 29, 1985 Jan 29, 1985
SLO-BID SANOFI AVENTIS US	50MG 75MG 100MG 125MG 200MG 300MG	A088269 001 A089539 001 A087892 001 A089540 001 A087893 001 A087894 001	Jan 31, 1985 May 10, 1989 Jan 31, 1985 May 10, 1989 Jan 31, 1985 Jan 31, 1985
SLO-PHYLLIN SANOFI AVENTIS US	60MG 125MG 250MG	A085206 001 A085203 001 A085205 001	May 24, 1982 May 24, 1982 May 24, 1982
SOMOPHYLLIN-CRT GRAHAM DM	50MG 100MG 200MG 250MG 300MG	A087763 001 A087194 001 A088382 001 A087193 001 A088383 001	Feb 27, 1985 Feb 27, 1985 Feb 27, 1985 Feb 27, 1985 Feb 27, 1985
THEOBID WHITBY	260MG	A085983 001	Mar 20, 1985
THEOBID JR. WHITBY	130MG	A087854 001	Mar 20, 1985
THEOCLEAR L.A.-130 SCHWARZ PHARMA	130MG	A086569 001	May 27, 1982
THEOCLEAR L.A.-260 SCHWARZ PHARMA	260MG	A086569 002	May 27, 1982
THEO-DUR SCHERING	50MG 75MG 125MG 200MG	A088022 001 A088015 001 A088016 001 A087995 001	Sep 10, 1985 Sep 10, 1985 Sep 10, 1985 Sep 10, 1985
THEOPHYLLINE CENT PHARMS	125MG 250MG	A088654 001 A088689 001	Feb 12, 1985 Feb 12, 1985
HOSPIRA	100MG 200MG 300MG	A089976 001 A089977 001 A089932 001	Jan 04, 1995 Jan 04, 1995 Jan 04, 1995
SANDOZ	260MG	A087462 001	May 11, 1982

DISCONTINUED DRUG PRODUCT LIST

6 - 320 (of 346)

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE; ORAL					
THEOPHYLLINE-SR					
SCHERER RP	300MG		A088255	001	Jun 12, 1986
THEOPHYL-SR					
ORTHO MCNEIL PHARM	125MG		A086480	001	Feb 08, 1985
	250MG		A086471	001	Feb 08, 1985
THEOVENT					
SCHERING	125MG		A087010	001	Jan 31, 1985
	250MG		A087910	001	Jan 31, 1985
ELIXIR; ORAL					
ELIXOMIN					
CENCI	80MG/15ML		A088303	001	Jan 25, 1984
LANOPHYLLIN					
LANNETT	80MG/15ML		A084578	001	
THEOLIXIR					
PANRAY	80MG/15ML		A084559	001	
THEOPHYL-225					
ORTHO MCNEIL PHARM	112.5MG/15ML		A086485	001	
THEOPHYLLINE					
ALPHARMA US PHARMS	80MG/15ML		A089223	001	May 27, 1988
CENCI	80MG/15ML		A087679	001	Apr 15, 1982
HALSEY	80MG/15ML		A085169	001	
PERRIGO	80MG/15ML		A085952	001	
PHARM ASSOC	80MG/15ML		A086720	001	
PRECISION DOSE	80MG/15ML		A085863	001	
ROXANE	80MG/15ML		A084739	001	
TARO	80MG/15ML		A089626	001	Oct 28, 1988
WOCKHARDT	80MG/15ML		A086748	001	
INJECTABLE; INJECTION					
THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	40MG/100ML		N019083	001	Nov 07, 1984
THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	80MG/100ML		N019083	002	Nov 07, 1984
THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	160MG/100ML		N019083	003	Nov 07, 1984
THEOPHYLLINE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	200MG/100ML		N019212	001	Nov 07, 1984
	200MG/100ML		N019826	004	Aug 14, 1992
THEOPHYLLINE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER					
B BRAUN	4MG/ML		N019212	003	Nov 07, 1984
	400MG/100ML		N019212	002	Nov 07, 1984
	400MG/100ML		N019826	005	Aug 14, 1992
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 321 (of 346)

THEOPHYLLINE

SYRUP; ORAL					
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	
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	
THEO-DUR					
SCHERING	100MG		A085328	001	
	200MG		A086998	001	
	300MG		A085328	002	
	450MG		A089131	001	Jun 25, 1986
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
T-PHYL					
PHARM RES ASSOC	200MG		A088253	001	Aug 17, 1983
UNI-DUR					
SCHERING	400MG		A089822	001	Jan 04, 1995
	600MG		A089823	001	Jan 04, 1995
UNIPHYL					
PURDUE PHARM PRODS	400MG		A087571	001	Sep 01, 1982
	600MG		A040086	001	Apr 15, 1996

THEOPHYLLINE SODIUM GLYCINATE

ELIXIR; ORAL					
SYNOPHYLATE					
CENT PHARMS	EQ 165MG BASE/15ML		N006333	008	

DISCONTINUED DRUG PRODUCT LIST

6 - 322 (of 346)

THEOPHYLLINE SODIUM GLYCINATE

TABLET; ORAL ASBRON NOVARTIS	EQ 150MG BASE	A085148 001
------------------------------------	---------------	-------------

THIABENDAZOLE

SUSPENSION; ORAL MINTEZOL MERCK	500MG/5ML	N016097 001
TABLET, CHEWABLE; ORAL MINTEZOL MERCK	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
BAXTER HLTHCARE	100MG/ML	A080575 001
BEL MAR	100MG/ML	A080718 001
	200MG/ML	A080712 001
DELL LABS	100MG/ML	A083775 001
HOSPIRA	100MG/ML	A040079 001
LUITPOLD	100MG/ML	A080667 001
PARKE DAVIS	100MG/ML	A080770 001
WATSON LABS	100MG/ML	A083534 001
	200MG/ML	A083534 002
WYETH AYERST	100MG/ML	A080553 001

THIAMYLAL SODIUM

INJECTABLE; INJECTION SURITAL PARKADEALE	1GM/VIAL	N007600 003
	5GM/VIAL	N007600 005
	10GM/VIAL	N007600 009

THIETHYLPERAZINE MALATE

INJECTABLE; INJECTION TORECAN NOVARTIS	5MG/ML	N012754 002
--	--------	-------------

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

DISCONTINUED DRUG PRODUCT LIST

6 - 323 (of 346)

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	
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	
TEVA PHARMS	30MG/ML	A089602	001	Nov 09, 1987	
	100MG/ML	A089603	001	Nov 09, 1987	
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					
IVAX PHARMS	10MG	A088270	001	Apr 14, 1983	
	15MG	A088271	001	Apr 14, 1983	
	25MG	A088272	001	Apr 14, 1983	
	50MG	A088194	001	Apr 14, 1983	
	100MG	A088273	001	Oct 03, 1983	
MUTUAL PHARM	10MG	A088375	001	Nov 18, 1983	
	15MG	A088461	001	Nov 18, 1983	
	25MG	A087264	001	Nov 18, 1983	
	50MG	A088370	001	Nov 18, 1983	
	100MG	A088379	001	Nov 16, 1983	
	150MG	A088737	001	Sep 26, 1984	
	200MG	A088738	001	Oct 16, 1984	
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	
	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 324 (of 346)

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

SANDOZ	150MG	A088136	001	Sep 17, 1986
	200MG	A088137	001	Sep 17, 1986
SUPERPHARM	10MG	A089103	001	Jul 02, 1985
	25MG	A089104	001	Jul 02, 1985
	50MG	A089105	001	Jul 02, 1985
TEVA	10MG	A088493	001	May 17, 1985
	100MG	A088456	001	May 17, 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	A088477	001	Nov 08, 1983
	15MG	A088562	001	May 11, 1984
	25MG	A088296	001	Jul 28, 1983
	25MG	A088478	001	Nov 08, 1983
	25MG	A088567	001	May 11, 1984
	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
	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				
APP PHARMS	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	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
WATSON LABS	1MG	A070600	001	Jun 05, 1987
	2MG	A071626	001	Jun 25, 1987
	5MG	A071627	001	Jun 25, 1987
	10MG	A070603	001	Jun 05, 1987
	10MG	A071628	001	Jun 25, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 325 (of 346)

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				
ROXANE	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	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

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
MYLAN	250MG		A075316	001 Nov 02, 1999
SANDOZ	250MG		A075318	001 Aug 20, 1999
WATSON LABS	250MG		A075309	001 Apr 26, 2000

DISCONTINUED DRUG PRODUCT LIST

6 - 326 (of 346)

TIMOLOL MALEATESOLUTION/DROPS; OPHTHALMIC
Timolol Maleate

AKORN	EQ 0.25% BASE	A074465	001	Mar 25, 1997
FOUGERA	EQ 0.25% BASE	A074667	001	Mar 25, 1997
	EQ 0.5% BASE	A074668	001	Mar 25, 1997
TABLET; ORAL				
BLOCADREN				
MERCK	5MG	N018017	001	
	10MG	N018017	002	
	20MG	N018017	004	
Timolol Maleate				
QUANTUM PHARMICS	5MG	A072466	001	May 19, 1989
	10MG	A072467	001	May 19, 1989
	20MG	A072468	001	May 19, 1989
SANDOZ	5MG	A072550	001	Apr 13, 1989
	10MG	A072551	001	Apr 13, 1989
	20MG	A072552	001	Apr 13, 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

TIOCONAZOLECREAM; TOPICAL
TZ-3
PFIZER

	1%	N018682	001	Feb 18, 1983
--	----	---------	-----	--------------

TIROFIBAN HYDROCHLORIDEINJECTABLE; 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

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

Tizanidine Hydrochloride

ACTAVIS ELIZABETH	EQ 2MG BASE	A076283	001	Jul 12, 2002
	EQ 4MG BASE	A076283	002	Jul 12, 2002
ACTAVIS TOTOWA	EQ 2MG BASE	A076281	001	Oct 20, 2003
	EQ 4MG BASE	A076281	002	Oct 20, 2003
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
ZANAFLEX				
ACORDA	EQ 2MG BASE	N020397	002	Feb 04, 2000

TOBRAMYCINSOLUTION/DROPS; OPHTHALMIC
TOBRAMYCIN

ALCON UNIVERSAL	0.3%	A063176	001	May 25, 1994
-----------------	------	---------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 327 (of 346)

TOBRAMYCIN SULFATEINJECTABLE; 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 40MG BASE/ML	A064021	002	May 31, 1994
	EQ 40MG BASE/ML	A064026	001	May 31, 1994
ASTRAZENECA	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
	EQ 40MG BASE/ML	A063122	001	Oct 31, 1994
BAXTER HLTHCARE	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
HOSPIRA	EQ 10MG BASE/ML	A063080	001	Apr 30, 1991
	EQ 40MG BASE/ML	A063161	001	May 29, 1991
MARSAM PHARMS LLC	EQ 40MG BASE/ML	A062945	002	Aug 09, 1989

TOCAINIDE HYDROCHLORIDETABLET; ORAL
TONOCARD

ASTRAZENECA	400MG	N018257	001	Nov 09, 1984
	600MG	N018257	002	Nov 09, 1984

TOLAZAMIDETABLET; 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
INTERPHARM	250MG	A071270	001	Sep 23, 1986
	500MG	A071271	001	Sep 23, 1986
IVAX SUB TEVA PHARMS	100MG	N018894	001	Nov 02, 1984
	250MG	N018894	002	Nov 02, 1984
	500MG	N018894	003	Nov 02, 1984
MUTUAL PHARM	100MG	A071357	001	Jul 16, 1987
	250MG	A071358	001	Jul 16, 1987
	500MG	A071359	001	Jul 16, 1987
PAR PHARM	100MG	A070159	001	Jan 06, 1986
	250MG	A070160	001	Jan 06, 1986
	500MG	A070161	001	Jan 06, 1986
SANDOZ	100MG	A071633	001	Dec 09, 1987
	250MG	A070289	001	Mar 13, 1986
	500MG	A070290	001	Mar 13, 1986
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
	500MG	A070244	001	Aug 01, 1986
TOLINASE				
PHARMACIA AND UPJOHN	100MG	N015500	002	

DISCONTINUED DRUG PRODUCT LIST

6 - 328 (of 346)

TOLAZAMIDE

TABLET; ORAL		
TOLINASE		
PHARMACIA AND UPJOHN	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	N010670 002
	500MG	N010670 001
TOLBUTAMIDE		
ALRA	500MG	A086141 001
ASCOT	500MG	A087541 001 Mar 01, 1983
BARR	500MG	A087121 001
CLONMEL HLTHCARE	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	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 PHARM INTL	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
MUTUAL PHARM	EQ 400MG BASE	A073311 001 Nov 27, 1991
SANDOZ	EQ 400MG BASE	A073462 001 Apr 30, 1992
TEVA	EQ 400MG BASE	A073519 001 May 29, 1992
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
IVAX SUB TEVA PHARMS	EQ 600MG BASE	A074399 001 Mar 28, 1996
SANDOZ	EQ 200MG BASE	A073588 001 Jul 31, 1992
	EQ 600MG BASE	A074002 001 Sep 27, 1993
TEVA	EQ 600MG BASE	A074729 001 Feb 27, 1997

DISCONTINUED DRUG PRODUCT LIST

6 - 329 (of 346)

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
TABLET; ORAL TOPAMAX				
JANSSEN PHARMS	300MG	N020505	003	Dec 24, 1996
	400MG	N020505	006	Dec 24, 1996
TOPIRAMATE BARR	25MG	A076315	001	Mar 27, 2009
	100MG	A076315	002	Mar 27, 2009
	200MG	A076315	003	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

TOPOTECAN HYDROCHLORIDE

SOLUTION; INTRAVENOUS
 TOPOTECAN
 SANDOZ INC EQ 1MG BASE/ML (EQ 1MG BASE/ML) N200199 001 Feb 25, 2011
 EQ 3MG BASE/3ML (EQ 1MG BASE/ML) N200199 002 Feb 25, 2011
 EQ 4MG BASE/4ML (EQ 1MG BASE/ML) N200199 003 Feb 25, 2011

TORSEMIDE

INJECTABLE; INJECTION
 DEMADEX
 ROCHE 50MG/5ML (10MG/ML) N020137 002 Aug 23, 1993
 20MG/2ML (10MG/ML) N020137 001 Aug 23, 1993

TRAMADOL HYDROCHLORIDE

TABLET; ORAL
 TRAMADOL HYDROCHLORIDE
 ACTAVIS ELIZABETH 50MG A075960 001 Jun 19, 2002
 ASTA 50MG A075974 001 Jul 12, 2002
 IVAX SUB TEVA PHARMS 50MG A075963 001 Jul 03, 2002
 ULTRAM
 JANSSEN PHARMS 100MG N020281 001 Mar 03, 1995

TRANEXAMIC ACID

TABLET; ORAL
 CYKLOKAPRON
 PHARMACIA AND UPJOHN 500MG N019280 001 Dec 30, 1986

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC
 TRAVATAN
 ALCON PHARMS LTD 0.004% N021257 001 Mar 16, 2001

DISCONTINUED DRUG PRODUCT LIST

6 - 330 (of 346)

TRAZODONE HYDROCHLORIDE

TABLET; ORAL DESYREL				
APOTHECON	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
MYLAN	50MG	A071405	001	Feb 27, 1991
	100MG	A071406	001	Feb 27, 1991
QUANTUM PHARMICS	100MG	A070921	001	Dec 01, 1986
SANDOZ	50MG	A072484	001	Apr 30, 1990
	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	A071112	001	Nov 17, 1986
	100MG	A071113	001	Nov 17, 1986
TRIALODINE				
QUANTUM PHARMICS	50MG	A070942	001	Dec 01, 1986

TRETINOIN

CAPSULE; ORAL VESANOID				
ROCHE	10MG	N020438	001	Nov 22, 1995
SOLUTION; TOPICAL TRETINOIN				
TEVA PHARMS	0.05%	A074873	001	Jun 19, 1998
SWAB; TOPICAL RETIN-A				
ORTHO JANSSEN	0.05%	N016921	002	

TRIAMCINOLONE

TABLET; ORAL ARISTOCORT				
ASTELLAS	1MG	N011161	009	
	2MG	N011161	004	
	4MG	N011161	007	
	8MG	N011161	011	
	16MG	N011161	010	
KENACORT				
BRISTOL MYERS SQUIBB	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	

DISCONTINUED DRUG PRODUCT LIST

6 - 331 (of 346)

TRIAMCINOLONE

TABLET; ORAL
TRIAMCINOLONE

TEVA	4MG	A084775	001
WATSON LABS	4MG	A084270	001
	4MG	A085834	001

TRIAMCINOLONE ACETONIDE

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

APOTHECON	0.5%	A083943	001
-----------	------	---------	-----

KENALOG-H

APOTHECON	0.1%	A086240	001
-----------	------	---------	-----

TRIACET

TEVA	0.1%	A084908	002
------	------	---------	-----

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

AMBIX	0.025%	A087932	001	May 09, 1983
-------	--------	---------	-----	--------------

G AND W LABS	0.1%	A089798	001	May 31, 1991
--------------	------	---------	-----	--------------

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
------	--------	---------	-----	--------------

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

DISCONTINUED DRUG PRODUCT LIST

6 - 332 (of 346)

TRIAMCINOLONE ACETONIDE

GEL; TOPICAL				
ARISTOGEL				
ASTELLAS	0.1%		A083380	001
INJECTABLE; INJECTION				
TRIAMCINOLONE ACETONIDE				
PARNELL	3MG/ML		N019503	001 Oct 16, 1987
WATSON LABS	40MG/ML		A085825	001
LOTION; TOPICAL				
KENALOG				
APOTHECON	0.025%		A084343	001
	0.1%		A084343	002
BRISTOL MYERS SQUIBB	0.025%		N011602	003
	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				
APOTHECON	0.5%		A083944	001
TRIAMCINOLONE ACETONIDE				
ACTAVIS MID ATLANTIC	0.1%		A087799	001 Jun 07, 1982
ALPHARMA US PHARMS	0.5%		A089913	001 Dec 23, 1988
G AND W LABS	0.025%		A089795	001 Dec 23, 1988
	0.1%		A089796	001 Dec 23, 1988
MORTON GROVE	0.025%		A088090	001 Sep 01, 1983
	0.1%		A088091	001 Sep 01, 1983
	0.5%		A088092	001 Sep 01, 1983
PHARMADERM	0.025%		A088692	001 Aug 02, 1984
	0.1%		A088690	001 Aug 02, 1984
TRYMEX				
SAVAGE LABS	0.025%		A088693	001 Aug 02, 1984
	0.1%		A088691	001 Aug 02, 1984
PASTE; DENTAL				
KENALOG IN ORABASE				
APOTHECON	0.1%		N012097	001
ORALONE				
TARO	0.1%		A071383	001 Jul 06, 1987
SPRAY, METERED; NASAL				
ALLERNAZE				
LUPIN ATLANTIS	0.05MG/SPRAY		N020120	001 Feb 04, 2000
NASCORT HFA				
SANOFI AVENTIS US	0.055MG/SPRAY		N020784	001 Apr 07, 2004

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION				
ARISTOCORT				
SANDOZ	25MG/ML		N011685	003
	40MG/ML		N012802	001
TRIAMCINOLONE DIACETATE				
AKORN	25MG/ML		A085122	001
	40MG/ML		A086394	001

DISCONTINUED DRUG PRODUCT LIST

6 - 333 (of 346)

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

TRIAMCINOLONE DIACETATE

WATSON LABS	40MG/ML	A084072	001
	40MG/ML	A085529	001
SYRUP; ORAL			
ARISTOCORT			
ASTELLAS	2MG / 5ML	N011960	004
KENACORT			
BRISTOL MYERS SQUIBB	EQ 4MG BASE/5ML	N012515	001

TRIAZOLAM

TABLET; ORAL

HALCION

PHARMACIA AND UPJOHN 0.5MG N017892 002 Nov 15, 1982

TRICHLORMETHIAZIDE

TABLET; ORAL

METAHYDRIN

SANOFI AVENTIS US	2MG	N012594	001	Jun 16, 1988
	4MG	N012594	002	Jun 16, 1988
NAQUA				
SCHERING	2MG	N012265	001	
	4MG	N012265	002	
TRICHOLOREX				
LANNETT	4MG	A083436	001	
	4MG	A085630	001	
TRICHLORMAS				
MAST MM	4MG	A086259	001	
TRICHLORMETHIAZIDE				
IMPAX LABS	4MG	A083967	001	
PAR PHARM	2MG	A087007	001	
	4MG	A087005	001	
SANDOZ	4MG	A086171	001	
TG UNITED LABS	4MG	A085568	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

TRIDIHEXYL 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

DISCONTINUED DRUG PRODUCT LIST

6 - 334 (of 346)

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL					
TRIFLUOPERAZINE HYDROCHLORIDE					
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	

TRIFLUPROMAZINE 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	

DISCONTINUED DRUG PRODUCT LIST

6 - 335 (of 346)

TRIHEXYPHENIDYL HYDROCHLORIDE

TABLET; ORAL				
TREMIN				
SCHERING	5MG		A080381	003
TRIHEXYPHENIDYL HYDROCHLORIDE				
NYLOS	5MG		A085622	001
VANGARD	2MG		A088035	001
WATSON LABS	2MG		A085117	001
	5MG		A085105	001

TRILOSTANE

CAPSULE; ORAL				
MODRASTANE				
BIOENVISION	30MG		N018719	002
	60MG		N018719	001
				Dec 31, 1984
				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
MORTON GROVE	EQ 2.5MG BASE/5ML		A088285	001
TABLET; ORAL				
TEMARIL				
ALLERGAN HERBERT	EQ 2.5MG BASE		N011316	001

TRIMETHADIONE

CAPSULE; ORAL				
TRIDIONE				
ABBOTT	300MG		N005856	005
SOLUTION; ORAL				
TRIDIONE				
ABBOTT	200MG/5ML		N005856	002

TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION				
ARFONAD				
ROCHE	50MG/ML		N008983	001

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION				
TRIMETHOBENZAMIDE HYDROCHLORIDE				
SMITH AND NEPHEW	100MG/ML		A088960	001
	100MG/ML		A089043	001
SOLOPAK	100MG/ML		A089094	001
WATSON LABS	100MG/ML		A086577	001
	100MG/ML		A087939	001
				Apr 04, 1986
				Apr 04, 1986
				Apr 04, 1986
				Oct 19, 1982
				Dec 28, 1982

TRIMETHOPRIM

TABLET; ORAL				
PROLOPRIM				
MONARCH PHARMS	100MG		N017943	001
	200MG		N017943	003
TRIMETHOPRIM				Jul 14, 1982
MUTUAL PHARM	100MG		A070494	001
	200MG		A070495	001
TRIMPEX				Jan 22, 1986
ROCHE	100MG		N017952	001
				Sep 24, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 336 (of 346)

TRIMETHOPRIM

TABLET; ORAL
 TRIMPEX 200
 ROCHE 200MG N017952 002 Nov 09, 1982

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL
 PRIMSOL
 FSC EQ 25MG BASE/5ML A074374 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
 50MG A085188 001
 TABLET, EXTENDED RELEASE; ORAL
 PBZ-SR
 NOVARTIS 50MG N010533 002
 100MG N010533 001

TRIPLE SULFA (SULFABENZAMIDE; SULFACETAMIDE; SULFATHIAZOLE)

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

DISCONTINUED DRUG PRODUCT LIST

6 - 337 (of 346)

TRIPLE SULFA (SULFABENZAMIDE; SULFACETAMIDE; SULFATHIAZOLE)

CREAM; VAGINAL					
TRIPLE SULFA					
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					
TEVA	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	
SULFALOID					
FOREST PHARMS	167MG;167MG;167MG		A080099	001	
SULFA-TRIPLE #2					
IMPAX LABS	167MG;167MG;167MG		A080079	001	
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	

DISCONTINUED DRUG PRODUCT LIST

6 - 338 (of 346)

TRISULFAPYRIMIDINES (SULFADIAZINE; SULFAMERAZINE; SULFAMETHAZINE)

TABLET; ORAL TRIPLE SULFOID PAL PAK	167MG;167MG;167MG	A080094 001
---	-------------------	-------------

TROGLITAZONE

TABLET; ORAL PRELAY SANKYO	200MG 300MG 400MG	N020719 001 Jan 29, 1997 N020719 003 Aug 04, 1997 N020719 002 Jan 29, 1997
REZULIN PFIZER PHARMS	200MG 300MG 400MG	N020720 001 Jan 29, 1997 N020720 003 Aug 04, 1997 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% 1%	N012111 002 N012111 004
MYDRIAFAIR PHARMAFAIR	0.5% 1%	A088274 001 Sep 16, 1983 A088230 001 Sep 16, 1983
TROPICAMIDE AKORN ALCON UNIVERSAL MIZA PHARMS USA WATSON LABS	1% 1% 0.5% 1% 0.5%	A088447 001 Aug 28, 1985 A089172 001 Dec 28, 1990 A087636 001 Jul 30, 1982 A087637 001 Aug 09, 1982 A089171 001 Dec 28, 1990

TROVAFLOXACIN MESYLATE

TABLET; ORAL TROVAN PFIZER	EQ 100MG BASE EQ 200MG BASE	N020759 001 Dec 18, 1997 N020759 002 Dec 18, 1997
----------------------------------	--------------------------------	--

TUBOCURARINE CHLORIDE

INJECTABLE; INJECTION TUBOCURARINE CHLORIDE BRISTOL MYERS SQUIBB HOSPIRA LILLY	3MG/ML 3MG/ML 3MG/ML	N005657 001 N006095 001 N006325 001
--	----------------------------	---

TYROPOANOATE SODIUM

CAPSULE; ORAL BILOPAQUE GE HEALTHCARE	750MG	N013731 001
---	-------	-------------

DISCONTINUED DRUG PRODUCT LIST

6 - 339 (of 346)

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC RESCULA SUCAMPO PHARMS	0.15%	N021214 001 Aug 03, 2000
---	-------	--------------------------

URACIL MUSTARD

CAPSULE; ORAL URACIL MUSTARD SHIRE	1MG	N012892 001
--	-----	-------------

UREA

INJECTABLE; INJECTION STERILE UREA HOSPIRA UREAPHIL HOSPIRA	40GM/VIAL	N017698 001
	40GM/VIAL	N012154 001

UREA C-13

FOR SOLUTION; ORAL HELICOSOL METABOLIC SOLUTIONS MERETEK UBT KIT (W/ PRANACTIN) OTSUKA AMERICA PYLORI-CHEK BREATH TEST DXS DEVICES	125MG/VIAL	N021092 001 Dec 17, 1999
	125MG/VIAL	N020586 001 Sep 17, 1996
	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 WATSON PHARMS	150MG	N019594 001 Dec 31, 1987
--	-------	--------------------------

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 SCHERER RP USL PHARMA	250MG	A070431 001 Feb 28, 1986
	250MG	A070195 001 Jul 02, 1987
	250MG	A070631 001 Jun 11, 1987
SYRUP; ORAL VALPROIC ACID APOTEX INC	250MG/5ML	A077105 001 Jul 29, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 340 (of 346)

VALSARTAN

CAPSULE; ORAL DIOVAN			
NOVARTIS	80MG	N020665	001 Dec 23, 1996
	160MG	N020665	002 Dec 23, 1996

VANCOMYCIN HYDROCHLORIDE

FOR SOLUTION; ORAL VANCOCIN HYDROCHLORIDE			
VIROPHARMA	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			
VIROPHARMA	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			
BAXTER HLTHCARE	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			
BAXTER HLTHCARE	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

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			
BAXTER HLTHCARE	10MG/VIAL	A075218	001 Aug 23, 1999
	20MG/VIAL	A075218	002 Aug 23, 1999
HOSPIRA	4MG/VIAL	A075558	001 Sep 11, 2001

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

DISCONTINUED DRUG PRODUCT LIST

6 - 341 (of 346)

VENLAFAXINE HYDROCHLORIDETABLET; 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
<u>VENLAFAXINE HYDROCHLORIDE</u>				
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 HYDROCHLORIDEINJECTABLE; INJECTION
CALAN

GD SEARLE LLC	2.5MG/ML	N018925	001	Mar 30, 1984
	2.5MG/ML	N019038	001	Mar 30, 1984
ISOPTIN				
FSC	2.5MG/ML	N018485	001	
<u>VERAPAMIL HYDROCHLORIDE</u>				
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
LUITPOLD	2.5MG/ML	A070225	001	Nov 12, 1985
	2.5MG/ML	A070617	001	Nov 12, 1985
MARSAM PHARMS LLC	2.5MG/ML	A072233	001	Feb 26, 1993
	2.5MG/ML	A073485	001	Sep 27, 1993
SMITH AND NEPHEW	2.5MG/ML	A070696	001	Jul 31, 1987
	2.5MG/ML	A070697	001	Jul 31, 1987
SOLOPAK	2.5MG/ML	A070695	001	Jul 31, 1987

TABLET; ORAL
CALAN

GD SEARLE LLC	160MG	N018817	004	Feb 23, 1988
ISOPTIN				
FSC	40MG	N018593	003	Nov 23, 1987
	80MG	N018593	001	Mar 08, 1982

VERAPAMIL HYDROCHLORIDE				
ACTAVIS ELIZABETH	80MG	A071019	001	Sep 24, 1986
	120MG	A070468	001	Sep 24, 1986
HERITAGE PHARMS INC	80MG	A071880	001	Apr 05, 1988
	120MG	A071881	001	Apr 05, 1988
MUTUAL PHARM	80MG	A070482	001	Sep 24, 1986
	80MG	A071489	002	Jan 13, 1988
	120MG	A070483	001	Sep 24, 1986
	120MG	A071489	001	Jan 13, 1988
PLIVA	40MG	A072751	001	Feb 23, 1996
	80MG	A072124	001	Jan 26, 1989
	120MG	A072125	001	Jan 26, 1989
SANDOZ	40MG	A073168	001	Jul 31, 1992
	80MG	A071423	001	May 24, 1988
	120MG	A071424	001	May 25, 1988

DISCONTINUED DRUG PRODUCT LIST

6 - 342 (of 346)

VERAPAMIL HYDROCHLORIDE

TABLET; ORAL					
VERAPAMIL HYDROCHLORIDE					
WARNER CHILCOTT	80MG		A070340	001	Sep 24, 1986
	120MG		A070341	001	Sep 24, 1986
WATSON LABS	40MG		A072799	001	Apr 28, 1989
TABLET, EXTENDED RELEASE; ORAL					
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					
PARKADEALE	EQ 187.4MG BASE/ML		N050523	001	
OINTMENT; OPHTHALMIC					
VIRA-A					
PARKADEALE	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

VINCRISTINE SULFATE

INJECTABLE; INJECTION					
ONCOVIN					
LILLY	1MG/VIAL		N014103	001	
	1MG/ML		N014103	003	Mar 07, 1984
	5MG/VIAL		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
APP PHARMS	1MG/ML		A076296	001	Dec 20, 2002
	1MG/ML		A076401	001	Oct 28, 2003
HOSPIRA	1MG/VIAL		A071559	001	Apr 11, 1988
	2MG/VIAL		A071560	001	Apr 11, 1988
	5MG/VIAL		A071561	001	Apr 11, 1988

VIOMYCIN SULFATE

INJECTABLE; INJECTION					
VIOCIN SULFATE					
PFIZER	EQ 1GM BASE/VIAL		A061086	001	
	EQ 5GM BASE/VIAL		A061086	002	

VITAMIN A

CAPSULE; ORAL					
AQUASOL A					
ASTRAZENECA	25,000USP UNITS		A083080	002	
	50,000USP UNITS		A083080	001	
VITAMIN A					
BANNER PHARMACAPS	50,000USP UNITS		A083973	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 343 (of 346)

VITAMIN A

CAPSULE; ORAL			
VITAMIN A			
CHASE CHEM	50,000 IU	A083351	001
EVERYLIFE	50,000 IU	A083134	001
IMPAKX 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
IMPAKX 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
	EQ 50,000 UNITS BASE	A083321	001
BANNER PHARMACAPS	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

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	50MG/VIAL	N009218	020
	75MG/VIAL	N009218	012
TABLET; ORAL			
ATHROMBIN			
PHARM RES ASSOC	5MG	N011771	003
	10MG	N011771	002
	25MG	N011771	001

DISCONTINUED DRUG PRODUCT LIST

6 - 344 (of 346)

WARFARIN SODIUM

TABLET; ORAL PANWARFIN ABBOTT	2MG 2.5MG 5MG 7.5MG 10MG	N017020 001 N017020 002 N017020 003 N017020 004 N017020 005
WARFARIN SODIUM SANDOZ	1MG 2MG 2.5MG 3MG 4MG 5MG 6MG 7.5MG 10MG	A040196 001 Sep 30, 1997 A040196 002 Sep 30, 1997 A040196 003 Sep 30, 1997 A040196 008 Jul 26, 2000 A040196 004 Sep 30, 1997 A040196 005 Sep 30, 1997 A040196 009 Jul 26, 2000 A040196 006 Sep 30, 1997 A040196 007 Sep 30, 1997
USL PHARMA	2MG 2.5MG 5MG	A088719 001 Jun 27, 1985 A088720 001 Aug 06, 1985 A088721 001 Jul 02, 1985
WATSON LABS	2MG 2.5MG 5MG 7.5MG 10MG	A086123 001 Aug 17, 1982 A086120 001 Aug 17, 1982 A086119 001 Aug 17, 1982 A086118 001 Aug 17, 1982 A086122 001 Aug 17, 1982

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	N017550 001
	0.25-5 CI/AMP	N017550 003
MALLINCKRODT	10mCi/VIAL 20mCi/VIAL	N018327 001 Mar 09, 1982 N018327 002 Mar 09, 1982
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	N017256 001
LANTHEUS MEDCL	6.3mCi/ML	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

DISCONTINUED DRUG PRODUCT LIST

6 - 345 (of 346)

ZALCITABINE

TABLET; ORAL HIVID			
ROCHE	0.375MG	N020199	001 Jun 19, 1992
	0.75MG	N020199	002 Jun 19, 1992

ZALEPLON

CAPSULE; ORAL ZALEPLON			
SANDOZ	5MG	A078095	001 Jun 06, 2008
	10MG	A078095	002 Jun 06, 2008

ZICONOTIDE

INJECTABLE; INTRATHECAL PRIALT			
AZUR PHARMA II	200MCG/2ML (100MCG/ML)	N021060	003 Dec 28, 2004

ZIDOVUDINE

TABLET; ORAL RETROVIR			
VIVI HLTHCARE	200MG	N020518	001 Dec 19, 1995
ZIDOVUDINE			
AUROBINDO PHARMA	60MG	N022294	001 Jul 23, 2009
MATRIX LABS LTD	100MG	N200732	001 Feb 23, 2011

ZILEUTON

TABLET; ORAL ZYFLO			
CORNERSTONE THERAP	300MG	N020471	001 Dec 09, 1996

ZINC SULFATE

INJECTABLE; INJECTION ZINC SULFATE			
ABRAXIS PHARM	EQ 1MG ZINC/ML	N019229	002 May 05, 1987

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION) ZOMETA			
NOVARTIS	EQ 4MG BASE/VIAL	N021223	001 Aug 20, 2001

ZOLPIDEM TARTRATE

TABLET; ORAL ZOLPIDEM TARTRATE			
MUTUAL PHARMA	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
WORLD GEN	5MG	A076062	001 Apr 23, 2007
	10MG	A076062	002 Apr 23, 2007
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			
MUTUAL PHARM	25MG	A077635	001 Dec 22, 2005
	50MG	A077635	002 Dec 22, 2005
	100MG	A077635	003 Dec 22, 2005
ROXANE	25MG	A077648	001 Dec 22, 2005
	50MG	A077648	002 Dec 22, 2005
	100MG	A077648	003 Dec 22, 2005

DISCONTINUED DRUG PRODUCT LIST

6 - 346 (of 346)

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

TEVA PHARMS

25MG

A077641 003 Dec 22, 2005

50MG

A077641 002 Dec 22, 2005

100MG

A077641 001 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 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	ASPIRIN; CAFFEINE; CARISOPRODOL TABLET; ORAL 160MG; 32MG; 200MG
ACETAMINOPHEN; ASPIRIN; BUTALBITAL CAPSULE OR TABLET; ORAL 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	ASPIRIN; CARISOPRODOL TABLET; ORAL 325MG; 200MG
ACETAMINOPHEN; BUTALBITAL CAPSULE OR TABLET; ORAL 325MG; 50MG 650MG; 50MG	ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE TABLET; ORAL 325MG; 200MG; 16MG
ACETAMINOPHEN; BUTALBITAL; CAFFEINE CAPSULE OR TABLET; ORAL 325MG; 50MG; 40MG 650MG; 50MG; 40MG	ASPIRIN; MEPROBAMATE TABLET; ORAL 325MG; 200MG
AMINOPHYLLINE TABLET; ORAL 100MG; 200MG	CHLOROTHIAZIDE TABLET; ORAL 250MG
ASPIRIN; BUTALBITAL CAPSULE OR TABLET; ORAL 325MG; 50MG 650MG; 50MG	HYDROXYZINE HYDROCHLORIDE TABLET; ORAL 10MG; 25MG; 50MG; 100MG
ASPIRIN; BUTALBITAL; CAFFEINE CAPSULE OR TABLET; ORAL 325MG; 50MG; 40MG 650MG; 50MG; 40MG	PREDNISONE TABLET; ORAL 1MG; 2.5MG; 5MG; 10MG; 20MG; 25MG; 50MG

APPENDIX A - PRODUCT NAME INDEX

A - 1

** * *

OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN

** 8 **

8-MOP, METHOXSALEN

** A **

ABELCET, AMPHOTERICIN B
ABILIFY, ARIPIPRAZOLE
ABLAVAR, GADOFOSVESET TRISODIUM
ABRAXANE, PACLITAXEL
ABREVA, DOCOSANOL (OTC)
ABSTRAL, FENTANYL CITRATE
ACANYA, BENZOYL PEROXIDE
ACARBOSE, ACARBOSE
ACCOLATE, ZAFIRLUKAST
ACCUNEB, ALBUTEROL SULFATE
ACCUPRIL, QUINAPRIL HYDROCHLORIDE
ACCURETIC, HYDROCHLOROTHIAZIDE
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACEON, PERINDOPRIL ERBUMINE
ACEPHEN, ACETAMINOPHEN (OTC)
ACETADOTE, ACETYL CYSTEINE
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE, ACETAMINOPHEN
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
ACETASOL HC, ACETIC ACID, GLACIAL
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETAZOLAMIDE, ACETAZOLAMIDE
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE, ACETIC ACID, GLACIAL
ACETIC ACID, ACETIC ACID, GLACIAL
ACETOHEXAMIDE, ACETOHEXAMIDE
ACETYLCYSTEINE, ACETYLCYSTEINE
ACIPHEX, RABEPRAZOLE SODIUM
ACLOVATE, ALCLOMETASONE DIPROPIONATE
ACTHREL, CORTICORELIN OVINE TRIFLUTATE
ACTIGALL, URSDIOL
ACTIQ, FENTANYL CITRATE
ACTIVELLA, ESTRADIOL
ACTONEL, RISEDRONATE SODIUM
ACTOPLUS MET XR, METFORMIN HYDROCHLORIDE
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
ACTOS, PIOGLITAZONE HYDROCHLORIDE
ACULAR LS, KETOROLAC TROMETHAMINE
ACULAR PRESERVATIVE FREE, KETOROLAC TROMETHAMINE
ACULAR, KETOROLAC TROMETHAMINE
ACUVAIL, KETOROLAC TROMETHAMINE
ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE, ACYCLOVIR SODIUM
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACYCLOVIR, ACYCLOVIR
ACZONE, DAPSONE
ADAGEN, PEGADEMASE BOVINE
ADALAT CC, NIFEDIPIINE
ADAPALENE, ADAPALENE
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 - 2

** A **

ADDERALL XR 5, AMPHETAMINE ASPARTATE
ADENOCARD, ADENOSINE
ADENOSCAN, ADENOSINE
ADENOSINE, ADENOSINE
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ADRENAClick, 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
ADVICOR, LOVASTATIN
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 PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADVIL, IBUPROFEN (OTC)
AEROBID, FLUNISOLIDE
AEROSPAN HFA, FLUNISOLIDE
AFEDITAB CR, NIFEDIPINE
AFINITOR, EVEROLIMUS
AFRINOL, PSEUDOEPHEDRINE SULFATE (OTC)
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
AGGRENOX, ASPIRIN
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
A-HYDROCORT, HYDROCORTISONE SODIUM SUCCINATE
AKBETA, LEVOBUNOLOL HYDROCHLORIDE
AK-FLUOR 10%, FLUORESCEIN SODIUM
AKINETON, BIPERIDEN HYDROCHLORIDE
AKNE-MYCIN, ERYTHROMYCIN
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKPRO, DIPIVEFRIN HYDROCHLORIDE
AKTEN, LIDOCAINE HYDROCHLORIDE
AKTOB, TOBRAMYCIN
ALA-CORT, HYDROCORTISONE
ALAMAST, PEMIROLAST POTASSIUM
ALA-SCALP, HYDROCORTISONE
ALAVERT, LORATADINE (OTC)
ALAWAY, KETOTIFEN FUMARATE (OTC)
ALBALON, NAPHAZOLINE HYDROCHLORIDE
ALBENZA, ALBENDAZOLE
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALCAINE, PROPARACAINA HYDROCHLORIDE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ALCOHOL 10% AND DEXTROSE 5%, ALCOHOL
ALCOHOL 5% AND DEXTROSE 5%, ALCOHOL
ALDACTAZIDE, HYDROCHLOROTHIAZIDE
ALDACTONE, SPIRONOLACTONE
ALDARA, IMIQUIMOD
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALEVE, NAPROXEN SODIUM (OTC)
ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)
ALFENTA, ALFENTANIL HYDROCHLORIDE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALIMTA, PEMETREXED DISODIUM
ALINIA, NITAZOXANIDE
ALKERAN, MELPHALAN

APPENDIX A - PRODUCT NAME INDEX

A - 3

** A **

ALKERAN, MELPHALAN HYDROCHLORIDE
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)
ALLI, ORLISTAT (OTC)
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
ALLOPURINOL, ALLOPURINOL
ALOCRIL, NEDOCROMIL SODIUM
ALOMIDE, LODOXAMIDE TROMETHAMINE
ALOPRIM, ALLOPURINOL SODIUM
ALORA, ESTRADIOL
ALOXI, PALONOSETRON HYDROCHLORIDE
ALPHAGAN P, BRIMONIDINE TARTRATE
ALPRAZOLAM, ALPRAZOLAM
ALPROSTADIL, ALPROSTADIL
ALREX, LOTEPREDNOL ETABONATE
ALSUMA, SUMATRIPTAN SUCCINATE
ALTABAX, RETAPAMULIN
ALTACE, RAMIPRIL
ALTAVERA, ETHINYL ESTRADIOL
ALTOPREV, LOVASTATIN
ALVESCO, CICLESONIDE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMARYL, GLIMEPIRIDE
AMBRIEN CR, ZOLPIDEM TARTRATE
AMBRIEN, ZOLPIDEM TARTRATE
AMBISOME, AMPHOTERICIN B
AMCINONIDE, AMCINONIDE
AMERGE, NARATRIPTAN HYDROCHLORIDE
A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE
AMICAR, AMINOCAPROIC ACID
AMIDATE, ETOMIDATE
AMIFOSTINE, AMIFOSTINE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOCAPROIC, AMINOCAPROIC ACID
AMINOHIPPURATE SODIUM, AMINOHIPPURATE SODIUM
AMINOPHYLLINE, AMINOPHYLLINE
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

APPENDIX A - PRODUCT NAME INDEX

A - 4

**** A ****

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 BESYLATE AND BENAZEPHIL HYDROCHLORIDE, AMLODIPINE BESYLATE
AMLODIPINE BESYLATE, 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 AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
AMOXIL, AMOXICILLIN
AMPHADASE, HYALURONIDASE
AMPHOTEC, AMPHOTERICIN B
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
AMTURNIDE, ALISKIREN HEMIFUMARATE
ANADROL-50, OXYMETHOLONE
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANAPROX DS, NAPROXEN SODIUM
ANAPROX, NAPROXEN SODIUM
ANASTROZOLE, ANASTROZOLE
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
ANCOBON, FLUCYTOSINE
ANDRODERM, TESTOSTERONE
ANDROGEL, TESTOSTERONE
ANDROID 10, METHYLTESTOSTERONE
ANDROID 25, METHYLTESTOSTERONE
AN-DTPA, TECHNETIUM TC-99M PENTETATE KIT
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ANESTACON, LIDOCAINE HYDROCHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANEXSIA 7.5/650, ACETAMINOPHEN
ANEXSIA, ACETAMINOPHEN
ANGELIQ, DROSPIRENONONE
ANGIOMAX, BIVALIRUDIN
ANSAID, FLURBIPROFEN
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
ANTABUSE, DISULFIRAM
ANTARA (MICRONIZED), FENOFLIBRATE
ANTHELIOS 20, AVOBENZONE (OTC)
ANTHELIOS 40, AVOBENZONE (OTC)
ANTHELIOS SX, AVOBENZONE (OTC)
ANTIVERT, MECLIZINE HYDROCHLORIDE
ANTIZOL, FOMEPIZOLE
ANTUROL, OXYBUTYNIN
ANUSOL HC, HYDROCORTISONE
ANZEMET, DOLASETRON MESYLATE
APHTHASOL, AMLEXANOX

APPENDIX A - PRODUCT NAME INDEX

A - 5

** A **

APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
 APIDRA, INSULIN GLULISINE RECOMBINANT
 APLENZIN, BUPROPION HYDROBROMIDE
 APOKYN, APOMORPHINE HYDROCHLORIDE
 APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
 APRISO, MESALAMINE
 APTIVUS, TIPRANAVIR
 AQUASOL A, VITAMIN A PALMITATE
 ARALEN, CHLOROQUINE PHOSPHATE
 ARANELLE, ETHINYL ESTRADIOL
 ARAVA, LEFLUNOMIDE
 ARCAPTA NEOHALER, INDACATEROL MALEATE
 AREDIA, PAMIDRONATE DISODIUM
 ARESTIN, MINOCYCLINE HYDROCHLORIDE
 ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
 ARGATROBAN, ARGATROBAN
 ARICEPT ODT, DONEPEZIL HYDROCHLORIDE
 ARICEPT, DONEPEZIL HYDROCHLORIDE
 ARIDOL KIT, MANNITOL
 ARIMIDEX, ANASTROZOLE
 ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
 ARIXTRA, FONDAPARINUX SODIUM
 AROMASIN, EXEMESTANE
 ARRANON, NELARABINE
 ARTHROTEC, DICLOFENAC SODIUM
 ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, ARTICAINE HYDROCHLORIDE
 ARTICAINE HYDROCHLORIDE WITH EPINEPHRINE, ARTICAINE HYDROCHLORIDE
 ASACOL HD, MESALAMINE
 ASACOL, MESALAMINE
 ASCLERA, POLIDOCANOL
 ASMANEX TWISTHALER, MOMETASONE FUROATE
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 ASTELIN, AZELASTINE HYDROCHLORIDE
 ASTEPRO, AZELASTINE HYDROCHLORIDE
 ASTRAMORPH PF, MORPHINE SULFATE
 ATACAND HCT, CANDESARTAN CILEXETIL
 ATACAND, CANDESARTAN CILEXETIL
 ATELVIA, RISEDRONATE SODIUM
 ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 ATENOLOL, ATENOLOL
 ATIVAN, LORAZEPAM
 ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 ATRALIN, TRETINOIN
 ATRIDOX, DOXYCYCLINE HYCLATE
 ATRIPLA, EFAVIRENZ
 ATROOPEN, ATROPINE
 ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE
 ATROVENT HFA, IPRATROPIUM BROMIDE
 ATROVENT, IPRATROPIUM BROMIDE
 AUGMENTIN '125', AMOXICILLIN
 AUGMENTIN '200', AMOXICILLIN
 AUGMENTIN '250', AMOXICILLIN
 AUGMENTIN '400', AMOXICILLIN
 AUGMENTIN '500', AMOXICILLIN
 AUGMENTIN '875', AMOXICILLIN
 AUGMENTIN ES-600, AMOXICILLIN
 AUGMENTIN XR, AMOXICILLIN
 AVAGARD, ALCOHOL (OTC)
 AVAGE, TAZAROTENE

APPENDIX A - PRODUCT NAME INDEX

A - 6

** A **

AVALIDE, HYDROCHLOROTHIAZIDE
 AVANDAMET, METFORMIN HYDROCHLORIDE
 AVANDARYL, GLIMEPIRIDE
 AVANDIA, ROSIGLITAZONE MALEATE
 AVAPRO, IRBESARTAN
 AVC, SULFANILAMIDE
 AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN HYDROCHLORIDE
 AVELOX, MOXIFLOXACIN HYDROCHLORIDE
 AVENTYL HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 AVIANE-28, ETHINYL ESTRADIOL
 AVINZA, MORPHINE SULFATE
 AVITA, TRETINOIN
 AVODART, DUTASTERIDE
 AXERT, ALMOTRIPTAN MALATE
 AXID AR, NIZATIDINE (OTC)
 AXID, NIZATIDINE
 AXIRON, TESTOSTERONE
 AYGESTIN, NORETHINDRONE ACETATE
 AZACTAM IN PLASTIC CONTAINER, AZTREONAM
 AZACTAM, AZTREONAM
 AZASAN, AZATHIOPRINE
 AZASITE, AZITHROMYCIN
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZATHIOPRINE, AZATHIOPRINE
 AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 AZELEX, AZELAIC ACID
 AZILECT, RASAGILINE MESYLATE
 AZITHROMYCIN, AZITHROMYCIN
 AZMACORT, TRIAMCINOLONE ACETONIDE
 AZOPT, BRINZOLAMIDE
 AZOR, AMLODIPINE BESYLATE
 AZTREONAM, AZTREONAM
 AZULFIDINE EN-TABS, SULFASALAZINE
 AZULFIDINE, SULFASALAZINE

** B **

BACIIM, BACITRACIN
 BACI-RX, BACITRACIN
 BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 BACITRACIN, BACITRACIN
 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
 BACTRIM DS, SULFAMETHOXAZOLE
 BACTRIM, SULFAMETHOXAZOLE
 BACTROBAN, MUPIROCIN
 BACTROBAN, MUPIROCIN CALCIUM
 BAL, DIMERCAPROL
 BALANCED SALT, CALCIUM CHLORIDE
 BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
 BALZIVA-28, ETHINYL ESTRADIOL
 BANZEL, RUFINAMIDE
 BARACLUDE, ENTECAVIR
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 BENADRYL PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 BENADRYL, DIPHENHYDRAMINE HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BENICAR HCT, HYDROCHLOROTHIAZIDE
 BENICAR, OLMESARTAN MEDOXOMIL
 BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 7

** B **

BENTYL, DICYCLOMINE HYDROCHLORIDE
 BENZACLIN, BENZOYL PEROXIDE
 BENZAMYCIN PAK, BENZOYL PEROXIDE
 BENZAMYCIN, BENZOYL PEROXIDE
 BENZONATATE, BENZONATATE
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 BEPREVE, BEPOTASTINE BESILATE
 BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
 BETADINE, POVIDONE-IODINE
 BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
 BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE
 BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 BETAPACE AF, SOTALOL HYDROCHLORIDE
 BETAPACE, SOTALOL HYDROCHLORIDE
 BETA-VAL, BETAMETHASONE VALERATE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 BETIMOL, TIMOLOL
 BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 BETOPTIC, BETAXOLOL HYDROCHLORIDE
 BEYAZ, DROSPIRENONE
 BIAXIN XL, CLARITHROMYCIN
 BIAXIN, CLARITHROMYCIN
 BICALUTAMIDE, BICALUTAMIDE
 BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
 BICILLIN C-R, PENICILLIN G BENZATHINE
 BICILLIN L-A, PENICILLIN G BENZATHINE
 BICNU, CARMUSTINE
 BIDIL, HYDRALAZINE HYDROCHLORIDE
 BILTRICIDE, PRAZIQUANTEL
 BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)
 BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BLEPH-10, SULFACETAMIDE SODIUM
 BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
 BLEPHAMIDE, PREDNISOLONE ACETATE
 BONIVA, IBANDRONATE SODIUM
 BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 BONTRIL, PHENDIMETRAZINE TARTRATE
 BRANCHAMIN 4% IN PLASTIC CONTAINER, AMINO ACIDS
 BRAVELLE, UROFOLLITROPIN
 BREATHTEK UBT FOR H-PYLORI, UREA C-13
 BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER, BRETYLIUM TOSYLATE
 BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 BREVIBLOC, ESMOLOL HYDROCHLORIDE
 BREVICON 28-DAY, ETHINYL ESTRADIOL
 BREVITAL SODIUM, METHOHEXITAL SODIUM
 BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)
 BRIELLYN, ETHINYL ESTRADIOL
 BRILINTA, TICAGRELOR
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 BROMDAY, BROMFENAC SODIUM
 BROMFED-DM, BROMPHENIRAMINE MALEATE
 BROMFENAC SODIUM, BROMFENAC SODIUM
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 BRONCHO SALINE, SODIUM CHLORIDE (OTC)
 BROVANA, ARFORMOTEROL TARTRATE
 BSS PLUS, CALCIUM CHLORIDE
 BSS, CALCIUM CHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 8

** B **

BUDESONIDE, BUDESONIDE
 BUMETANIDE, BUMETANIDE
 BUPHENYL, SODIUM PHENYLBUTYRATE
 BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
 BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 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
 BUTAPAP, ACETAMINOPHEN
 BUTISOL SODIUM, BUTABARBITAL SODIUM
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 BUTTRANS, BUPRENORPHINE
 BYETTA, EXENATIDE SYNTHETIC
 BYSTOLIC, NEBIVOLOL HYDROCHLORIDE

** C **

CABERGOLINE, CABERGOLINE
 CADUET, AMLODIPINE BESYLATE
 CAFCIT, CAFFEINE CITRATE
 CAFERGOT, CAFFEINE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALAN, VERAPAMIL HYDROCHLORIDE
 CALCIJEX, CALCITRIOL
 CALCIPOTRIENE, CALCIPOTRIENE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CALCITRIOL, CALCITRIOL
 CALCIUM ACETATE, CALCIUM ACETATE
 CALCIUM CARBONATE, FAMOTIDINE AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
 CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
 CALDOLOR, IBUPROFEN
 CAMBIA, DICLOFENAC POTASSIUM
 CAMILA, NORETHINDRONE
 CAMPRAL, ACAMPROSATE CALCIUM
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 CANASA, MESALAMINE
 CANCIDAS, CASPOFUNGIN ACETATE
 CANTIL, MEPENZOLETATE BROMIDE
 CAPASTAT SULFATE, CAPREOMYCIN SULFATE
 CAPEX, FLUOCINOLONE ACETONIDE
 CAPITAL AND CODEINE, ACETAMINOPHEN
 CAPITAL SOLEIL 15, AVOBENZONE (OTC)
 CAPOTEN, CAPTOPRIL
 CAPOZIDE 25/15, CAPTOPRIL
 CAPOZIDE 25/25, CAPTOPRIL
 CAPOZIDE 50/15, CAPTOPRIL
 CAPOZIDE 50/25, CAPTOPRIL
 CAPTOPRIL AND HYDROCHLORTHIAZIDE, CAPTOPRIL
 CAPTOPRIL, CAPTOPRIL
 CARAC, FLUOROURACIL
 CARAFATE, SUCRALFATE
 CARBAGLU, CARGLUMIC ACID
 CARBAMAZEPINE, CARBAMAZEPINE

APPENDIX A - PRODUCT NAME INDEX

A - 9

** C **

CARBATROL, CARBAMAZEPINE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
CARBOPLATIN, CARBOPLATIN
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 SR, NICARDIPINE HYDROCHLORIDE
CARDENE, NICARDIPINE HYDROCHLORIDE
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
CARDIZEM, DILTIAZEM HYDROCHLORIDE
CARDURA XL, DOXAZOSIN MESYLATE
CARDURA, DOXAZOSIN MESYLATE
CARISOPRODOL AND ASPIRIN, ASPIRIN
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CARISOPRODOL, CARISOPRODOL
CARMOL HC, HYDROCORTISONE ACETATE
CARNITOR SF, LEVOCARNITINE
CARNITOR, LEVOCARNITINE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CARTIA XT, DILTIAZEM HYDROCHLORIDE
CARVEDILOL, CARVEDILOL
CASQDEX, BICALUTAMIDE
CATAFLAM, DICLOFENAC POTASSIUM
CATAPRES, CLONIDINE HYDROCHLORIDE
CATAPRES-TTS-1, CLONIDINE
CATAPRES-TTS-2, CLONIDINE
CATAPRES-TTS-3, CLONIDINE
CAVERJECT IMPULSE, ALPROSTADIL
CAVERJECT, ALPROSTADIL
CAYSTON, AZTREONAM
CEDAX, CEFTIBUTEN DIHYDRATE
CEENU, LOMUSTINE
CEFACLOR, CEFACLOR
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFAZOLIN AND DEXTROSE, 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
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
CEFOTAXIME, CEFOTAXIME SODIUM
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, CEFOTETAN DISODIUM
CEFOTETAN, CEFOTETAN DISODIUM
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM
CEFOXITIN, CEFOXITIN
CEFOXITIN, CEFOXITIN SODIUM
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME
CEFTAZIDIME, CEFTAZIDIME
CEFTIN, CEFUROXIME AXETIL
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM

APPENDIX A - PRODUCT NAME INDEX

A - 10

**** C ****

CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME SODIUM IN PLASTIC CONTAINER, CEFUROXIME SODIUM
CEFUROXIME SODIUM, CEFUROXIME SODIUM
CELEBREX, CELECOXIB
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELESTONE, BETAMETHASONE
CELEXA, CITALOPRAM HYDROBROMIDE
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CELONTIN, MENTSUXIMIDE
CENESTIN, ESTROGENS, CONJUGATED SYNTHETIC A
CENTANY, MUPIROCIN
CEPHALEXIN, CEPHALEXIN
CEREDASE, ALGLUCERASE
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CEREZYME, IMIGLUCERASE
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
CERVIDIL, DINOPROSTONE
CESAMET, NABILONE
CETAMIDE, SULFACETAMIDE SODIUM
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE
CETROTIDE, CETRORELIX
CEVIMELINE, CEVIMELINE HYDROCHLORIDE
CHANTIX, VARENICLINE TARTRATE
CHEMET, SUCCIMER
CHENODIOL, CHENODIOL
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
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)
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 COLD, IBUPROFEN (OTC)
CHILDREN'S MOTRIN, IBUPROFEN (OTC)
CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
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)
CHLORASCRUB MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWAB, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE

APPENDIX A - PRODUCT NAME INDEX

A - 11

**** C ****

CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
CHLOROTHIAZIDE, CHLOROTHIAZIDE
CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORTHALIDONE, CHLORTHALIDONE
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)
CHLORZOXAZONE, CHLORZOXAZONE
CHOLAC, LACTULOSE
CHOLEDYL SA, OXTRIPHYLLINE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
CHOLOGRAFIN MEGLUMINE, IODIPAMIDE MEGLUMINE
CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
CHROMITOPE SODIUM, SODIUM CHROMATE CR-51
CIALIS, TADALAFIL
CICLOPIROX, CICLOPIROX
CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
CILOSTAZOL, CILOSTAZOL
CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIMETIDINE, CIMETIDINE
CIMETIDINE, CIMETIDINE (OTC)
CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPRO XR, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN HYDROCHLORIDE
CIPRODEX, CIPROFLOXACIN
CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPROFLOXACIN, CIPROFLOXACIN
CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
CISPLATIN, CISPLATIN
CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CITANEST FORTE DENTAL, EPINEPHRINE BITARTRATE
CITANEST PLAIN DENTAL, PRILOCAINE HYDROCHLORIDE
CLADRIBINE, CLADRIBINE
CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CLAFORAN, CEFOTAXIME SODIUM
CLARAVIS, ISOTRETINOIN
CLARINEX D 24 HOUR, DESLORATADINE
CLARINEX, DESLORATADINE
CLARINEX-D 12 HOUR, DESLORATADINE
CLARITHROMYCIN, CLARITHROMYCIN
CLARITIN HIVES RELIEF REDTAB, LORATADINE (OTC)
CLARITIN HIVES RELIEF, LORATADINE (OTC)
CLARITIN REDTABS, LORATADINE (OTC)
CLARITIN, LORATADINE (OTC)
CLARITIN-D 24 HOUR, LORATADINE (OTC)
CLARITIN-D, LORATADINE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
CLENZ-LYTE, POLYETHYLENE GLYCOL 3350
CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE

APPENDIX A - PRODUCT NAME INDEX

A - 12

** C **

CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLEOCIN T, CLINDAMYCIN PHOSPHATE
CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
CLEOCIN, CLINDAMYCIN PHOSPHATE
CLEVIPREX, CLEVIDIPIINE BUTYRATE
CLIMARA PRO, ESTRADIOL
CLIMARA, ESTRADIOL
CLINDA-DERM, CLINDAMYCIN PHOSPHATE
CLINDAGEL, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLINDESSE, CLINDAMYCIN PHOSPHATE
CLINDETS, CLINDAMYCIN PHOSPHATE
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, AMINO ACIDS
CLINIMIX E 2.75/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 2.75/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/20 SULFITE-FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/15 SULFITE-FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/20 SULFITE-FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/35 SULFITE-FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
CLINORIL, SULINDAC
CLOBETASOL PROPIONATE (EMOLlient), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOBEX, CLOBETASOL PROPIONATE
CLODERM, CLOCORTOLONE PIVALATE
CLOLAR, CLOFARABINE
CLOMID, CLOMIPHENE CITRATE
CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLONIDINE, CLONIDINE

APPENDIX A - PRODUCT NAME INDEX

A - 13

**** C ****

CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CLORPRES, CHLORTHALIDONE
CLOTTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOTTRIMAZOLE, CLOTTRIMAZOLE
CLOTTRIMAZOLE, CLOTTRIMAZOLE (OTC)
CLOZAPINE, CLOZAPINE
CLOZARIL, CLOZAPINE
COARTEM, ARTEMETHER
CODEINE SULFATE, CODEINE SULFATE
COGENTIN, BENZTROPIINE MESYLATE
CO-GESIC, ACETAMINOPHEN
COLAZAL, BALSALAZIDE DISODIUM
COLCRYS, COLCHICINE
COLESTID, COLESTIPOL HYDROCHLORIDE
COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
COLGATE TOTAL, SODIUM FLUORIDE (OTC)
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
COL-PROBENECID, COLCHICINE
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
COLYTE, POLYETHYLENE GLYCOL 3350
COLYTE-FLAVORED, POLYETHYLENE GLYCOL 3350
COMBIGAN, BRIMONIDINE TARTRATE
COMBIPATCH, ESTRADIOL
COMBIVENT RESPIMAT, ALBUTEROL SULFATE
COMBIVENT, ALBUTEROL SULFATE
COMBIVIR, LAMIVUDINE
COMMIT, NICOTINE POLACRILEX (OTC)
COMPLERA, EMTRICITABINE
COMPRO, PROCHLORPERAZINE
COMTAN, ENTACAPONE
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
CONDYLOX, PODOFILOX
CONRAY 30, IOTHALAMATE MEGLUMINE
CONRAY 43, IOTHALAMATE MEGLUMINE
CONRAY, IOTHALAMATE MEGLUMINE
CONSTILAC, LACTULOSE
CONSTULOSE, LACTULOSE
CONZIP, TRAMADOL HYDROCHLORIDE
COPAXONE, GLATIRAMER ACETATE
COPEGUS, RIBAVIRIN
CORDARONE, AMIODARONE HYDROCHLORIDE
CORDRAN SP, FLURANDRENOLIDE
CORDRAN, FLURANDRENOLIDE
COREG CR, CARVEDILOL PHOSPHATE
COREG, CARVEDILOL
CORGARD, NADOLOL
CORLOPAM, FENOLDOPAM MESYLATE
CORMAX, CLOBETASOL PROPIONATE
CORTEF, HYDROCORTISONE
CORTENEMA, HYDROCORTISONE
CORTIFOAM, HYDROCORTISONE ACETATE
CORTISONE ACETATE, CORTISONE ACETATE
CORTISPORIN, BACITRACIN ZINC
CORTISPORIN, HYDROCORTISONE
CORTISPORIN, HYDROCORTISONE ACETATE
CORTROSYN, COSYNTROPIN
COVERT, IBUTILIDE FUMARATE
CORZIDE, BENDROFLUMETHIAZIDE
COSMEGEN, DACTINOMYCYIN
COSOPT, DORZOLAMIDE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 14

**** C ****

COSYNTROPIN, COSYNTROPIN
COUMADIN, WARFARIN SODIUM
COVERA-HS, VERAPAMIL HYDROCHLORIDE
COZAAR, LOSARTAN POTASSIUM
CREON, LIPASE
CRESTOR, ROSUVASTATIN CALCIUM
CRINONE, PROGESTERONE
CRIXIVAN, INDINAVIR SULFATE
CROLOM, CROMOLYN SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
CROTAN, CROTAMITON
CRYSELLE, ETHINYLMESTRADIOL
CUBICIN, DAPTOMYCIN
CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
CUPRIMINE, PENICILLAMINE
CUROSURF, PORACTANT ALFA
CUTIVATE, FLUTICASONE PROPIONATE
CUVPOSA, GLYCOPYRROLATE
CYANOCOBALAMIN, CYANOCOBALAMIN
CYANOKIT, HYDROXOCOBALAMIN
CYCLAFEM 1/35, ETHINYLMESTRADIOL
CYCLAFEM 7/7/7, ETHINYLMESTRADIOL
CYCLESSA, DESOGESTREL
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
CYCLOSET, BROMOCRIPTINE MESYLATE
CYCLOSPORINE, CYCLOSPORINE
CYKLOKAPRON, TRANEXAMIC ACID
CYMBALTA, DULOXETINE HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
CYSTADANE, BETAINE HYDROCHLORIDE
CYSTAGON, CYSTEAMINE BITARTRATE
CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
CYTARABINE, CYTARABINE
CYTOMEL, LIOTHYRONINE SODIUM
CYTOSAR-U, CYTARABINE
CYTOTEC, MISOPROSTOL
CYTOVENE, GANCICLOVIR SODIUM
CYTOXAN, CYCLOPHOSPHAMIDE

**** D ****

D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
DACARBAZINE, DACARBAZINE
DACOGEN, DECITABINE
DACTINOMYCIN, DACTINOMYCIN
DALIRESP, ROFLUMILAST
DANAZOL, DANAZOL
DANTRIUM, DANTROLENE SODIUM
DANTROLENE SODIUM, DANTROLENE SODIUM
DAPSONE, DAPSONE
DARAPRIM, PYRIMETHAMINE
DASETTA 1/35, ETHINYLMESTRADIOL
DASETTA 7/7/7, ETHINYLMESTRADIOL
DATSCAN, IOFLUPANE I-123
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DAUNOXOME, DAUNORUBICIN CITRATE
DAYPRO, OXaprozin

APPENDIX A - PRODUCT NAME INDEX

A - 15

** D **

DAYTRANA, METHYLPHENIDATE
DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
DDAVP, DESMOPRESSIN ACETATE
DECLOMYCIN, DEMECLOCYCLINE HYDROCHLORIDE
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
DEFINITY, PERFLUTREN
DELATESTRYL, 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 3.5% 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
DELFLLEX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLLEX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLLEX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELFLLEX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
DEMADEX, TORSEMIDE
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
DEMEROL, MEPERIDINE HYDROCHLORIDE
DEMSER, METYROSINE
DENAVIR, PENCICLOVIR SODIUM
DENDRID, IDOXURIDINE
DEPACON, VALPROATE SODIUM
DEPAKENE, VALPROIC ACID
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
DEPEN, PENICILLAMINE
DEPOCYT, CYTARABINE
DEPODUR, MORPHINE SULFATE
DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
DERMABET, BETAMETHASONE VALERATE
DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
DERMATOP E EMOLLIENT, PREDNICARBATE
DERMATOP, PREDNICARBATE
DERMOTIC, FLUOCINOLONE ACETONIDE
DESFERAL, DEFEROXAMINE MESYLATE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DESLOTRADATINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLOTRADATINE
DESLOTRADATINE, DESLOTRADATINE
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DESOGEN, DESOGESTREL
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DESONATE, DESONIDE
DESONIDE, DESONIDE
DESOWEN, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
DETROL LA, TOLTERODINE TARTRATE
DETROL, TOLTERODINE TARTRATE
DEXAMETHASONE INTENSOL, DEXAMETHASONE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE

APPENDIX A - PRODUCT NAME INDEX

A - 16

** D **

DEXAMETHASONE, DEXAMETHASONE
DEXASPORIN, DEXAMETHASONE
DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
DEXEDRINE, DEXTROAMPHETAMINE SULFATE
DEXFERRUM, IRON DEXTRAN
DEXILANT, DEXLANSOPRAZOLE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
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 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 75 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 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
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 ACETATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
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 (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
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, DEXTROSE

APPENDIX A - PRODUCT NAME INDEX

A - 17

** D **

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER,
DEXTROSE
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 60% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
DIABETA, GLYBURIDE
DIABINESE, CHLORPROPAMIDE
DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIAMOX, ACETAZOLAMIDE
DIAMOX, ACETAZOLAMIDE SODIUM
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
DIASTAT ACUDIAL, DIAZEPAM
DIASTAT, DIAZEPAM
DIAZEPAM INTENSOL, DIAZEPAM
DIAZEPAM, DIAZEPAM
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICLOxacillin SODIUM, DICLOxacillin SODIUM
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE-FREE), DICYCLOMINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIDANOSINE, DIDANOSINE
DIDREX, BENZPHETAMINE HYDROCHLORIDE
DIDRONEL, ETIDRONATE DISODIUM
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
DIFFERIN, ADAPALENE
DIFCID, FIDAXOMICIN
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
DIFLUCAN IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
DIFLUCAN, FLUCONAZOLE
DIFLUNISAL, DIFLUNISAL
DIGOXIN, DIGOXIN
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DILACOR XR, DILTIAZEM HYDROCHLORIDE
DILANTIN, PHENYTOIN
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
DILATRATE-SR, ISOSORBIDE DINITRATE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILAUDID-HP, HYDROMORPHONE HYDROCHLORIDE
DILT-CD, DILTIAZEM HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 18

**** D ****

DILTZAC, DILTIAZEM HYDROCHLORIDE
DIMENHYDRINATE, DIMENHYDRINATE
DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
DIOVAN HCT, HYDROCHLOROTHIAZIDE
DIOVAN, VALSARTAN
DIPENTUM, OLSALAZINE SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIPIVEFRIN HYDROCHLORIDE, DIPIVEFRIN HYDROCHLORIDE
DIPRIVAN, PROPOFOL
DIPROLENE AF, BETAMETHASONE DIPROPIONATE
DIPROLENE, BETAMETHASONE DIPROPIONATE
DIPYRIDAMOLE, DIPYRIDAMOLE
DISOPHROL, DEXBROMPHENIRAMINE MALEATE (OTC)
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
DISPERMOX, AMOXICILLIN
DISULFIRAM, DISULFIRAM
DITROPAN XL, OXYBUTYNIN CHLORIDE
DITROPAN, OXYBUTYNIN CHLORIDE
DIURIL, CHLOROTHIAZIDE
DIURIL, CHLOROTHIAZIDE SODIUM
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DIVIGEL, ESTRADIOL
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOCEFREZ, DOCETAXEL
DOCETAXEL, DOCETAXEL
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
DORAL, QUAZEPAM
DORIBAX, DORIPENEM
DORYX, DOXYCYCLINE HYCLATE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DOVONEX, CALCIPOTRIENE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXE PIN HYDROCHLORIDE, DOXE PIN HYDROCHLORIDE
DOXERCALCI FEROL, DOXERCALCI FEROL
DOXIL, DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXY 100, DOXYCYCLINE HYCLATE
DOXY 200, DOXYCYCLINE HYCLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DOXYCYCLINE, DOXYCYCLINE HYCLATE
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
DRAIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
DRISDOL, ERGOCALCI FEROL
DRIXORAL PLUS, ACETAMINOPHEN (OTC)
DRIXORAL, DEXBROMPHENIRAMINE MALEATE (OTC)
DRONABINOL, DRONABINOL
DROPERIDOL, DROPERIDOL
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
DROXIA, HYDROXYUREA
DTIC-DOME, DACARBAZINE

APPENDIX A - PRODUCT NAME INDEX

A - 19

**** D ****

DTPA, TECHNETIUM TC-99M PENTETATE KIT
DUAC, BENZOYL PEROXIDE
DUETACT, GLIMEPIRIDE
DUEXIS, FAMOTIDINE
DULERA, FORMOTEROL FUMARATE
DUODOTE, ATROPINE
DUONEB, ALBUTEROL SULFATE
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
DUTASTERIDE, DUTASTERIDE
DUVOID, BETHANECHOL CHLORIDE
DYAZIDE, HYDROCHLORTIAZIDE
DYNACIN, MINOCYCLINE HYDROCHLORIDE
DYNACIRC CR, ISRADIPINE
DYN-HEX, CHLORHEXIDINE GLUCONATE (OTC)
DYRENIUM, TRIAMTERENE

**** E ****

E.E.S. 200, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
EC-NAPROSYN, NAPROXEN
ECONAZOLE NITRATE, ECONAZOLE NITRATE
EDARBI, AZILSARTAN KAMEDOXOMIL
EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
EDECRIN, ETHACRYNATE SODIUM
EDECRIN, ETHACRYNIC ACID
EDEX, ALPROSTADIL
EDLUAR, ZOLPIDEM TARTRATE
EDURANT, RILPIVIRINE HYDROCHLORIDE
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
EFFIENT, PRASUGREL HYDROCHLORIDE
EFUDEX, FLUOROURACIL
E-GLADES, ERYTHROMYCIN
EGRIFTA, TESAMORELIN ACETATE
ELDEPRYL, SELEGILINE HYDROCHLORIDE
ELESTAT, EPINASTINE HYDROCHLORIDE
ELESTRIN, ESTRADIOL
ELIDEL, PIMECROLIMUS
ELIGARD, LEUPROLIDE ACETATE
ELIMIT, PERMETHRIN
ELIPHOS, CALCIUM ACETATE
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 E, CLOBETASOL PROPIONATE
EMBELINE, CLOBETASOL PROPIONATE
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
EMEND, APREPITANT
EMEND, FOSAPREPITANT DIMEGLUMINE

APPENDIX A - PRODUCT NAME INDEX

A - 20

** E **

EMLA, LIDOCAINE
EMOQUETTE, DESOGESTREL
EMSAM, SELEGILINE
EMTRIVA, EMTRICITABINE
ENABLEX, DARIFENACIN HYDROBROMIDE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ENALAPRILAT, ENALAPRILAT
ENDOMETRIN, PROGESTERONE
ENDOSOL EXTRA, CALCIUM CHLORIDE
ENDURON, METHYCLOTHIAZIDE
ENFLURANE, ENFLURANE
ENJUVIA, ESTROGENS, CONJUGATED SYNTHETIC B
ENLON, EDROPHONIUM CHLORIDE
ENLON-PLUS, ATROPINE SULFATE
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
ENPRESSE-28, ETHINYLMESTRADIOL
ENTEREG, ALVIMOPAN
ENTOCORT EC, BUDESONIDE
ENULOSE, LACTULOSE
EOVIST, GADOXETATE DISODIUM
EPIDUO, ADAPALENE
EPIFOAM, HYDROCORTISONE ACETATE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
EPINEPHRINE, EPINEPHRINE (OTC)
EPIPEN JR., EPINEPHRINE
EPIPEN, EPINEPHRINE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
EPITOL, CARBAMAZEPINE
EPIVIR, LAMIVUDINE
EPIVIR-HBV, LAMIVUDINE
EPLERENONE, EPLERENONE
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
EPZICOM, ABACAVIR SULFATE
EQUETRO, CARBAMAZEPINE
ERAXIS, ANIDULAFUNGIN
ERGOCALCIFEROL, ERGOCALCIFEROL
ERGOLOID MESYLATES, ERGOLOID MESYLATES
ERGOMAR, ERGOTAMINE TARTRATE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ERRIN, NORETHINDRONE
ERTACZO, SERTACONAZOLE NITRATE
ERYC, ERYTHROMYCIN
ERYGEL, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERY-TAB, ERYTHROMYCIN
ERYTHRA-DERM, ERYTHROMYCIN
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN, ERYTHROMYCIN
ERYTHRO-STATIN, ERYTHROMYCIN
ESGIC-PLUS, ACETAMINOPHEN
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
ESTAZOLAM, ESTAZOLAM
ESTRACE, ESTRADIOL
ESTRADERM, ESTRADIOL
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
ESTRADIOL AND NORGESTIMATE, ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

A - 21

** E **

ESTRADIOL CYPIONATE, ESTRADIOL CYPIONATE
 ESTRADIOL VALERATE, ESTRADIOL VALERATE
 ESTRADIOL, ESTRADIOL
 ESTRASORB, ESTRADIOL HEMIHYDRATE
 ESTRING, ESTRADIOL
 ESTROGEL, ESTRADIOL
 ESTRONE, ESTRONE
 ESTROPIPATE, ESTROPIPATE
 ESTROSTEP FE, ETHINYL ESTRADIOL
 ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 ETHAMOLIN, ETHANOLAMINE OLEATE
 ETHOSUXIMIDE, ETHOSUXIMIDE
 ETHRANE, ENFLURANE
 ETHYOL, AMIFOSTINE
 ETIDRONATE DISODIUM, ETIDRONATE DISODIUM
 ETODOLAC, ETODOLAC
 ETOMIDATE, ETOMIDATE
 ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 ETOPOSIDE, ETOPOSIDE
 EURAX, CROTAMITON
 EVAMIST, ESTRADIOL
 EVISTA, RALOXIFENE HYDROCHLORIDE
 EVOCLIN, CLINDAMYCIN PHOSPHATE
 EVOXAC, CEVIMELINE HYDROCHLORIDE
 EXALGO, HYDROMORPHONE HYDROCHLORIDE
 EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
 EXELDERM, SULCONAZOLE NITRATE
 EXELON, RIVASTIGMINE
 EXELON, RIVASTIGMINE TARTRATE
 EXEMESTANE, EXEMESTANE
 EXFORGE HCT, AMLODIPINE BESYLATE
 EXFORGE, AMLODIPINE BESYLATE
 EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
 EXJADE, DEFERASIROX
 EXPAREL, BUPIVACAINE
 EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 EXTINA, KETOCONAZOLE
 EXTRANEAL, ICODEXTRIN
 E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)

** F **

FACTIVE, GEMIFLOXACIN MESYLATE
 FAMCICLOVIR, FAMCICLOVIR
 FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK), FAMOTIDINE
 FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FAMVIR, FAMCICLOVIR
 FANAPT, ILOPERIDONE
 FARESTON, TOREMIFENE CITRATE
 FASLODEX, FULVESTRANT
 FAZACLO ODT, CLOZAPINE
 FELBAMATE, FELBAMATE
 FELBATOL, FELBAMATE
 FELDENE, PIROXICAM
 FELODIPINE, FELODIPINE
 FEMARA, LETROZOLE
 FEMCON FE, ETHINYL ESTRADIOL
 FEMHRT, ETHINYL ESTRADIOL
 FEMRING, ESTRADIOL ACETATE
 FEMSTAT 3, BUTOCONAZOLE NITRATE (OTC)

APPENDIX A - PRODUCT NAME INDEX

A - 22

** F **

FEMTRACE, ESTRADIOL ACETATE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
FENOGLIDE, FENOFIBRATE
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
FENTANYL CITRATE, FENTANYL CITRATE
FENTANYL-100, FENTANYL
FENTANYL-12, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL
FENTORA, FENTANYL CITRATE
FERAHEME, FERUMOXYTOL
FERIDEX I.V., FERUMOXIDES
FERRIPROX, DEFERIPRONE
FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FIBRICOR, FENOFIBRIC ACID
FINACEA, AZELAIC ACID
FINASTERIDE, FINASTERIDE
FIORICET W/ CODEINE, ACETAMINOPHEN
FIORICET, ACETAMINOPHEN
FIORINAL W/CODEINE, ASPIRIN
FIORINAL, ASPIRIN
FIRAZYR, ICATIBANT ACETATE
FIRMAGON, DEGARELIX ACETATE
FLAGYL ER, METRONIDAZOLE
FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
FLAGYL, METRONIDAZOLE
FLAREX, FLUOROMETHOLONE ACETATE
FLAVORED COLESTID, COLESTIOL HYDROCHLORIDE
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLECTOR, DICLOFENAC EPOLAMINE
FLEXERIL, CYCLOBENZAPRINE HYDROCHLORIDE
FLOLAN, EPOPROSTENOL SODIUM
FLOMAX, TAMSULOSIN HYDROCHLORIDE
FLONASE, FLUTICASONE PROPIONATE
FLO-PRED, PREDNISOLONE ACETATE
FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
FLOVENT HFA, FLUTICASONE PROPIONATE
FLOXIN OTIC, OFLOXACIN
FLOXURIDINE, FLOXURIDINE
FLUCANAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUCONAZOLE, FLUCONAZOLE
FLUCYTOSINE, FLUCYTOSINE
FLUDARA, FLUDARABINE PHOSPHATE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FLUDEOXYGLUCOSE F 18, FLUDEOXYGLUCOSE F-18
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
FLUMADINE, RIMANTADINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 23

** F **

FLUMAZENIL, FLUMAZENIL
 FLUNISOLIDE, FLUNISOLIDE
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUORESCITE, FLUORESCIN SODIUM
 FLUOROPLEX, FLUOROURACIL
 FLUOROURACIL, FLUOROURACIL
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUOXYMESTERONE, FLUOXYMESTERONE
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
 FLURBIPROFEN, FLURBIPROFEN
 FLUTAMIDE, FLUTAMIDE
 FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FML FORTE, FLUOROMETHOLONE
 FML, FLUOROMETHOLONE
 FOAMCOAT, ALUMINUM HYDROXIDE (OTC)
 FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
 FLOTYN, PRALATREXATE
 FOMEPIZOLE, FOMEPIZOLE
 FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 FORADIL, FORMOTEROL FUMARATE
 FORANE, ISOFLURANE
 FORFIVO XL, BUPROPION HYDROCHLORIDE
 FORTAMET, METFORMIN HYDROCHLORIDE
 FORTAZ IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
 FORTAZ, CEFTAZIDIME
 FORTEO, TERIPARATIDE RECOMBINANT HUMAN
 FORTESTA, TESTOSTERONE
 FORTICAL, CALCITONIN SALMON RECOMBINANT
 FOSAMAX PLUS D, ALENDRONATE SODIUM
 FOSAMAX, ALENDRONATE SODIUM
 FOSCARNET SODIUM, FOSCARNET SODIUM
 FOSCAVIR, FOSCARNET SODIUM
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM, 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% W/ ELECTROLYTES, AMINO ACIDS
 FREAMINE III 8.5%, AMINO ACIDS
 FROVA, FROVATRIPTAN SUCCINATE
 FULCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 FURADANTIN, NITROFURANTOIN
 FUROSEMIDE, FUROSEMIDE
 FUSILEV, LEVOLEUCOVORIN CALCIUM
 FUZEON, ENFUVIRTIDE

** G **

GABAPENTIN, GABAPENTIN
 GABITRIL, TIAGABINE HYDROCHLORIDE
 GABLOFEN, BACLOFEN

APPENDIX A - PRODUCT NAME INDEX

A - 24

**** G ****

GADAVIST, GADOBUTROL
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
GALZIN, ZINC ACETATE
GANCICLOVIR, GANCICLOVIR
GANCICLOVIR, GANCICLOVIR SODIUM
GANIRELIX ACETATE INJECTION, GANIRELIX ACETATE
GANITE, GALLIUM NITRATE
GASTROCROM, CROMOLYN SODIUM
GASTROGRAFIN, DIATRIZOATE MEGLUMINE
GASTROMARK, FERUMOXSIL
GATIFLOXACIN, GATIFLOXACIN
GAVISCON, ALUMINUM HYDROXIDE (OTC)
GELNIQUE, OXYBUTYNIN CHLORIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GEMCITABINE, GEMCITABINE HYDROCHLORIDE
GEMFIBROZIL, GEMFIBROZIL
GEMZAR, GEMCITABINE HYDROCHLORIDE
GENERLAC, LACTULOSE
GENGRAF, CYCLOSPORINE
GENOPTIC, GENTAMICIN SULFATE
GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT
GENOTROPIN, SOMATROPIN RECOMBINANT
GENTAK, GENTAMICIN SULFATE
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
GEN-XENE, CLORAZEPATE DIPOTASSIUM
GEDON, ZIPRASIDONE HYDROCHLORIDE
GEDON, ZIPRASIDONE MESYLATE
GILDESS FE 1.5/30, ETHINYL ESTRADIOL
GILDESS FE 1/20, ETHINYL ESTRADIOL
GILENYA, FINGOLIMOD
GLEEVEC, IMATINIB MESYLATE
GLIADEL, CARMUSTINE
GLIMEPIRIDE, GLIMEPIRIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLIPIZIDE, GLIPIZIDE
GLUCAGEN, GLUCAGON HYDROCHLORIDE RECOMBINANT
GLUCAGON, GLUCAGON RECOMBINANT
GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
GLUCOPHAGE, METFORMIN HYDROCHLORIDE
GLUCOTROL XL, GLIPIZIDE
GLUCOTROL, GLIPIZIDE
GLUCOVANCE, GLYBURIDE
GLUMETZA, METFORMIN HYDROCHLORIDE
GLYBURIDE (MICRONIZED), GLYBURIDE
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYBURIDE, GLYBURIDE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
GLYCOLAX, POLYETHYLENE GLYCOL 3350
GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
GLCOPYRROLATE, GLCOPYRROLATE
GLYNASE, GLYBURIDE
GLYSET, MIGLITOL
GOLYTELY, POLYETHYLENE GLYCOL 3350
GONAL-F RFF PEN, FOLLITROPIN ALFA/BETA
GONAL-F RFF, FOLLITROPIN ALFA/BETA
GONAL-F, FOLLITROPIN ALFA/BETA
GRALISE, GABAPENTIN
GRANisetron HYDROCHLORIDE PRESERVATIVE FREE, GRANisetron HYDROCHLORIDE
GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
GRANISTERON HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
GRIFULVIN V, GRISEOFULVIN, MICROCRYSTALLINE

APPENDIX A - PRODUCT NAME INDEX

A - 25

** G **

GRISEOFULVIN, GRISEOFULVIN
GRISEOFULVIN, GRISEOFULVIN, MICROCRYSTALLINE
GRIS-PEG, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
GUAIFENESIN, GUAIFENESIN (OTC)
GUANABENZ ACETATE, GUANABENZ ACETATE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
GYNAZOLE-1, BUTOCONAZOLE NITRATE
GYNE-LOTRIMIN 3 COMBINATION PACK, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN 3, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN COMBINATION PACK, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN, CLOTRIMAZOLE (OTC)

** H **

H.P. ACTHAR GEL, CORTICOTROPIN
HABITROL, NICOTINE (OTC)
HALAVEN, ERIBULIN MESYLATE
HALCION, TRIAZOLAM
HALDOL, HALOPERIDOL DECANOATE
HALDOL, HALOPERIDOL LACTATE
HALFLYTELY, BISACODYL
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HALOG, HALCINONIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL
HALOPERIDOL, HALOPERIDOL LACTATE
HEATHER, NORETHINDRONE
HECTOROL, DOXERCALCIFEROL
HELIDAC, BISMUTH SUBSALICYLATE
HEMABATE, CARBOPROST TROMETHAMINE
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 AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS AND 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 SODIUM
HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
HEPATAMINE 8%, AMINO ACIDS
HEPATASOL 8%, AMINO ACIDS
HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
HEPSERA, ADEFOVIR DIPIVOXIL
HERPLEX, IDOXURIDINE
HEXBRIX, IOXAGLATE MEGLUMINE
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 KWIKPEN, INSULIN LISPRO RECOMBINANT
HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 50/50 PEN, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT

APPENDIX A - PRODUCT NAME INDEX

A - 26

** H **

HUMALOG MIX 75/25 PEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
 HUMALOG PEN, INSULIN LISPRO 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 PEN, INSULIN RECOMBINANT HUMAN (OTC)
 HUMULIN R, INSULIN RECOMBINANT HUMAN
 HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
 HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 HYDASE, HYALURONIDASE
 HYDERGINE, ERGOLOID MESYLATES
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDREA, HYDROXYUREA
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 HYDROCORTISONE ACETATE, HYDROCORTISONE ACETATE
 HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
 HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 HYDROCORTISONE, HYDROCORTISONE
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 HYDRO-RX, HYDROCORTISONE
 HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYUREA, HYDROXYUREA
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN
 HYTRIN, TERAZOSIN HYDROCHLORIDE
 HYZAAR, HYDROCHLOROTHIAZIDE

** I **

IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 IBUPROFEN AND DIPHENHYDRAMINE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
 IBUPROHM, IBUPROFEN (OTC)
 IBU-TAB 200, IBUPROFEN (OTC)
 IBU-TAB, IBUPROFEN
 IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 IC-GREEN, INDOCYANINE GREEN
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IDKIT:HP, UREA C-13
 IFEX, IFOSFAMIDE
 IFOSFAMIDE, IFOSFAMIDE
 IFOSFAMIDE/MESNA KIT, IFOSFAMIDE
 IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 IMIQUIMOD, IMIQUIMOD
 IMITREX STATDOSE, SUMATRIPTAN SUCCINATE

APPENDIX A - PRODUCT NAME INDEX

A - 27

** I **

IMITREX, SUMATRIPTAN
IMITREX, SUMATRIPTAN SUCCINATE
IMODIUM A-D EZ CHEWS, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM MULTI-SYMPOTM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM, LOPERAMIDE HYDROCHLORIDE
IMPLANON, ETONOGESTREL
IMURAN, AZATHIOPRINE
INAPSINE, DROPERIDOL
INCIVEK, TELAPREVIR
INCRELEX, MECASERMIN RECOMBINANT
INDAPAMIDE, INDAPAMIDE
INDERAL LA, PROPRANOLOL HYDROCHLORIDE
INDERAL, PROPRANOLOL HYDROCHLORIDE
INDERIDE-40/25, HYDROCHLOROTHIAZIDE
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 SODIUM, INDOMETHACIN SODIUM
INDOMETHACIN, INDOMETHACIN
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
INFASURF PRESERVATIVE FREE, CALFACTANT
INFED, IRON DEXTRAN
INFUMORPH, MORPHINE SULFATE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INFUVITE PEDIATRIC, ASCORBIC ACID
INNOHEP, TINZAPARIN SODIUM
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
INOMAX, NITRIC OXIDE
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INSPRA, EPLERENONE
INTAL, CROMOLYN SODIUM
INTEGRILIN, EPTIFIBATIDE
INTELENCE, ETRAVIRINE
INTERMEZZO, ZOLPIDEM TARTRATE
INTRALIPID 10%, SOYBEAN OIL
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
INTROVALE, ETHINYLY ESTRADIOL
INTUNIV, GUANFACINE HYDROCHLORIDE
INVAGESIC FORTE, ASPIRIN
INVAGESIC, ASPIRIN
INVANZ, ERTAPENEM SODIUM
INVEGA SUSTENNA, PALIPERIDONE PALMITATE
INVEGA, PALIPERIDONE
INVIRASE, SAQUINAVIR MESYLATE
IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IOPAMIDOL-250, IOPAMIDOL
IOPAMIDOL-300, IOPAMIDOL
IOPAMIDOL-370, IOPAMIDOL
IOPIDINE, APRACLONIDINE HYDROCHLORIDE
IOSAT, POTASSIUM IODIDE (OTC)
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IPRIVASK, DESIRUDIN RECOMBINANT
IQUIX, LEVOFLOXACIN

APPENDIX A - PRODUCT NAME INDEX

A - 28

**** I ****

IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 ISENTRESS, RALTEGRAVIR POTASSIUM
 ISOCaine HYDROCHLORIDE W/ LEVONORDEFRIN, LEVONORDEFRIN
 ISOCaine HYDROCHLORIDE, Mepivacaine HYDROCHLORIDE
 ISOFLURANE, ISOFLURANE
 ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE E IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE S 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
 ISOPTIN SR, VERAPAMIL HYDROCHLORIDE
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 ISORDIL, ISOSORBIDE DINITRATE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 ISOSULFAN BLUE, ISOSULFAN BLUE
 ISOTONIC GENTAMICIN SULFATE IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 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
 IVY BLOCK, BENTOQUATAM (OTC)
 IXEMpra KIT, IXABEPILONE

**** J ****

JAKAFI, RUXOLITINIB PHOSPHATE
 JALYN, DUTASTERIDE
 JANTOVEN, WARFARIN SODIUM
 JANUMET, METFORMIN HYDROCHLORIDE
 JANUVIA, SITAGLIPTIN PHOSPHATE
 JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 JEVTANA KIT, CABAZITAXEL
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL
 JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 JUNEL FE 1/20, ETHINYL ESTRADIOL
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 JUVISYNC, SIMVASTATIN

**** K ****

K+10, POTASSIUM CHLORIDE
 KADIAN, MORPHINE SULFATE
 KALETRA, LOPINAVIR
 KALEXATE, SODIUM POLYSTYRENE SULFONATE
 KANAMYCIN SULFATE, KANAMYCIN SULFATE
 KAON CL-10, POTASSIUM CHLORIDE
 KAPVAY, CLONIDINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 29

**** K ****

KARIVA, DESOGESTREL
 KAYEXALATE, SODIUM POLYSTYRENE SULFONATE
 KEFLEX, CEPHALEXIN
 KEFZOL, CEFAZOLIN SODIUM
 KELNOR, ETHINYL ESTRADIOL
 KENALOG, TRIAMCINOLONE ACETONIDE
 KENALOG-10, TRIAMCINOLONE ACETONIDE
 KENALOG-40, TRIAMCINOLONE ACETONIDE
 KEPTRA XR, LEVETIRACETAM
 KEPTRA, LEVETIRACETAM
 KETALAR, KETAMINE HYDROCHLORIDE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETEK, TELITHROMYCIN
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 KETOZOLO, KETOCONAZOLE
 KINEVAC, SINCALIDE
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 KLARON, SULFACETAMIDE SODIUM
 KLOPIN, CLONAZEPAM
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE
 KLOR-CON M20, POTASSIUM CHLORIDE
 KLOR-CON, POTASSIUM CHLORIDE
 KLOTRIX, POTASSIUM CHLORIDE
 KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
 K-TAB, POTASSIUM CHLORIDE
 KUVAN, SAPROPTERIN DIHYDROCHLORIDE

**** L ****

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LABETALOL HYDROCLORIDE, LABETALOL HYDROCHLORIDE
 LAC-HYDRIN, AMMONIUM LACTATE
 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 CD, LAMOTRIGINE
 LAMICTAL ODT, LAMOTRIGINE
 LAMICTAL XR, LAMOTRIGINE
 LAMICTAL, LAMOTRIGINE
 LAMISIL AT, TERBINAFINE (OTC)
 LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMISIL, TERBINAFINE HYDROCHLORIDE
 LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
 LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
 LAMIVUDINE, LAMIVUDINE
 LAMOTRIGINE, LAMOTRIGINE
 LAMPRENE, CLOFAZIMINE
 LANORINAL, ASPIRIN
 LANOXIN PEDIATRIC, DIGOXIN
 LANOXIN, DIGOXIN
 LANSOPRAZOLE, LANSOPRAZOLE
 LANTUS, INSULIN GLARGINE RECOMBINANT
 LAROTID, AMOXICILLIN
 LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LASIX, FUROSEMIDE
 LASTACAFT, ALCAFTADINE
 LATANOPROST, LATANOPROST
 LATISSE, BIMATOPROST
 LATUDA, LURASIDONE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 30

** L **

LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
LAZANDA, FENTANYL CITRATE
LEFLUNOMIDE, LEFLUNOMIDE
LESCOL XL, FLUVASTATIN SODIUM
LESCOL, FLUVASTATIN SODIUM
LESSINA-28, ETHINYLMESTRADIOL
LETAIRIS, AMBRISENTAN
LETROZOLE, LETROZOLE
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEUKERAN, CHLORAMBUCYL
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
LEUSTATIN, CLADRIBINE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVAQUIN, LEVOFLOXACIN
LEVATOL, PENBUTOLOL SULFATE
LEVEMIR, INSULIN DETEMIR RECOMBINANT
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
LEVETIRACETAM, LEVETIRACETAM
LEVITRA, VARDENAFIL HYDROCHLORIDE
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
LEVOCARNITINE, LEVOCARNITINE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVOFLOXACIN, LEVOFLOXACIN
LEVONEST, ETHINYLMESTRADIOL
LEVONORGESTREL AND ETHINYLMESTRADIOL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL
LEVONORGESTREL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL
LEVONORGESTREL, LEVONORGESTREL
LEVONORGESTREL, LEVONORGESTREL (OTC)
LEVOPHED, NOREPINEPHRINE BITARTRATE
LEVORA 0.15/30-28, ETHINYLMESTRADIOL
LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
LEVO-T, LEVOTHYROXINE SODIUM
LEVOTHROID, LEVOTHYROXINE SODIUM
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
LEVOXYL, LEVOTHYROXINE SODIUM
LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
LEXAPRO, ESCITALOPRAM OXALATE
LEXISCAN, REGADENOSON
LEXIVA, FOSAMPRENAVIR CALCIUM
LIALDA, MESALAMINE
LIBRUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
LIDEX, FLUOCINONIDE
LIDEX-E, FLUOCINONIDE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
LIDOCAINE AND TETRACAINE, LIDOCAINE
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 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.8% IN 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 IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
LIDOCAINE, LIDOCAINE
LIDODERM, LIDOCAINE

APPENDIX A - PRODUCT NAME INDEX

A - 31

** L **

LIDOPEN, LIDOCAINE HYDROCHLORIDE
LIGNOSPIN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPIN STANDARD, EPINEPHRINE BITARTRATE
LIMBITROL DS, AMITRIPTYLINE HYDROCHLORIDE
LIMBITROL, AMITRIPTYLINE HYDROCHLORIDE
LINCOCIN, LINCOMYCIN HYDROCHLORIDE
LINDANE, LINDANE
LIORESAL, BACLOFEN
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
LIPITOR, ATORVASTATIN CALCIUM
LIPOFEN, FENOFLIBRATE
LIPOSYN II 10%, SAFFLOWER OIL
LIPOSYN II 20%, SAFFLOWER OIL
LIPOSYN III 10%, SOYBEAN OIL
LIPOSYN III 20%, SOYBEAN OIL
LIPOSYN III 30%, SOYBEAN OIL
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LITHIUM CARBONATE, LITHIUM CARBONATE
LITHIUM CITRATE, LITHIUM CITRATE
LITHOBID, LITHIUM CARBONATE
LITHOSTAT, ACETOHYDROXAMIC ACID
LIVALO, PITAVASTATIN CALCIUM
LO LOESTRIN FE, ETHINYLMESTRADIOL
LO/OVRAL-28, ETHINYLMESTRADIOL
LOCHOLEST LIGHT, CHOLESTYRAMINE
LOCHOLEST, CHOLESTYRAMINE
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
LOCOID, HYDROCORTISONE BUTYRATE
LODOSYN, CARBIDOPA
LOESTRIN 21 1.5/30, ETHINYLMESTRADIOL
LOESTRIN 21 1/20, ETHINYLMESTRADIOL
LOESTRIN 24 FE, ETHINYLMESTRADIOL
LOESTRIN FE 1.5/30, ETHINYLMESTRADIOL
LOESTRIN FE 1/20, ETHINYLMESTRADIOL
LOMOTIL, ATROPINE SULFATE
LONOX, ATROPINE SULFATE
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOOPERAMIDE HYDROCHLORIDE (OTC)
LOPERAMIDE HYDROCHLORIDE, LOOPERAMIDE HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOOPERAMIDE HYDROCHLORIDE (OTC)
LOPID, GEMFIBROZIL
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPRESSOR, METOPROLOL TARTRATE
LOPROX, CICLOPIROX
LOPURIN, ALLOPURINOL
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE REDIDOSE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
LORAZEPAM INTENSOL, LORAZEPAM
LORAZEPAM PRESERVATIVE FREE, LORAZEPAM
LORAZEPAM, LORAZEPAM
LORTAB, ACETAMINOPHEN
LORYNA, DROSPIRENONE
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOSEASONIQUE, ETHINYLMESTRADIOL
LOTEMAX, LOTEPREDNOL ETABONATE
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
LOTENSIN, BENAZEPRIL HYDROCHLORIDE
LOTREL, AMLODIPINE BESYLATE
LOTTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
LOTRISONE, BETAMETHASONE DIPROPIONATE
LOTRONEX, ALOSETRON HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 32

** L **

LOVASTATIN, LOVASTATIN
 LOVAZA, OMEGA-3-ACID ETHYL ESTERS
 LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 LOVENOX, ENOXAPARIN SODIUM
 LOW-OGESTREL-21, ETHINYL ESTRADIOL
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 LTA II KIT, LIDOCAINE HYDROCHLORIDE
 LUFYLLIN, DYPHYLLINE
 LUMIGAN, BIMATOPROST
 LUNESTA, ESZOPICLONE
 LUPRON DEPOT, LEUPROLIDE ACETATE
 LUPRON DEPOT-PED, LEUPROLIDE ACETATE
 LUSEDRA, FOSPROPOFOL DISODIUM
 LUVERIS, LUTROPIN ALFA
 LUVOX CR, FLUVOXAMINE MALEATE
 LUVOX, FLUVOXAMINE MALEATE
 LUXIQ, BETAMETHASONE VALERATE
 LYBREL, ETHINYL ESTRADIOL
 LYMPHAZURIN, ISOSULFAN BLUE
 LYRICA, PREGABALIN
 LYSODREN, MITOTANE
 LYSTEDA, TRANEXAMIC ACID

** M **

M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
 M.V.I. ADULT, ASCORBIC ACID
 M.V.I. PEDIATRIC, ASCORBIC ACID
 M.V.I.-12 (WITHOUT VITAMIN K), ASCORBIC ACID
 MACROBID, NITROFURANTOIN
 MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
 MACUGEN, PEGAPTANIB SODIUM
 MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE, MAGNESIUM HYDROXIDE
 MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MAGNEVIST, GADOPENTETATE DIMEGLUMINE
 MAKENA, HYDROXYPROGESTERONE CAPROATE
 MALARONE PEDIATRIC, ATOVAQUONE
 MALARONE, ATOVAQUONE
 MALATHION, MALATHION
 MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
 MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER, MANNITOL
 MANNITOL 10%, MANNITOL
 MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%, MANNITOL
 MANNITOL 15%, MANNITOL
 MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 20%, MANNITOL
 MANNITOL 25%, MANNITOL
 MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%, MANNITOL
 MANNITOL 5%, MANNITOL
 MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
 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
 MARINOL, DRONABINOL
 MARPLAN, ISOCARBOXAZID
 MATULANE, PROCARBAZINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 33

** M **

MAVIK, TRANDOLAPRIL
MAXAIR, PIRBUTEROL ACETATE
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MAXIDEX, DEXAMETHASONE
MAXIPIME, CEFEPIME HYDROCHLORIDE
MAXITROL, DEXAMETHASONE
MAXZIDE, HYDROCHLORTIAZIDE
MAXZIDE-25, HYDROCHLORTIAZIDE
MD-76R, DIATRIZOATE MEGLUMINE
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
MDP-BRACCO, TECHNETIUM TC-99M MEDRONATE KIT
MEBENDAZOLE, MEDEBENDAZOLE
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
MEGACE, MEGESTROL ACETATE
MEGATOPE, ALBUMIN IODINATED I-131 SERUM
MEGESTROL ACETATE, MEGESTROL ACETATE
MELOXICAM, MELOXICAM
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
MEMBRANEBLUE, TRYPLAN BLUE
MENEST, ESTROGENS, ESTERIFIED
MENOPUR, LUTEINIZING HORMONE
MENOSTAR, ESTRADIOL
MEN'S ROGAINE, MINOXIDIL (OTC)
MENTAX, BUTENAFINE HYDROCHLORIDE
MENTAX-TC, BUTENAFINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPHYTON, PHYTONADIONE
MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
MEPRON, ATOVAQUONE
MERCAPTOPURINE, MERCAPTOPURINE
MEROPENEM, 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 INTENSOL, METHADONE HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHADOSE, METHADONE HYDROCHLORIDE
METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
METHAZOLAMIDE, METHAZOLAMIDE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
METHERGINE, METHYLERGONOVINE MALEATE
METHIMAZOLE, METHIMAZOLE
METHOCARBAMOL AND ASPIRIN, ASPIRIN
METHOCARBAMOL, METHOCARBAMOL

APPENDIX A - PRODUCT NAME INDEX

A - 34

** M **

METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
METHYCLOTHIAZIDE, METHYCLOTHIAZIDE
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPA, METHYLDOPA
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METHYLTESTOSTERONE, METHYLTESTOSTERONE
METIPRANOL, METIPRANOL HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOLAZONE, METOLAZONE
METOPIRONE, METYRAPONE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METOZOLV ODT, METOCLOPRAMIDE HYDROCHLORIDE
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
METROCREAM, METRONIDAZOLE
METROGEL, METRONIDAZOLE
METROGEL-VAGINAL, METRONIDAZOLE
METROLOTION, METRONIDAZOLE
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
METRONIDAZOLE, METRONIDAZOLE
METVIXIA, METHYL AMINOLEVULINATE HYDROCHLORIDE
MEVACOR, LOVASTATIN
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
MIACALCIN, CALCITONIN SALMON
MICARDIS HCT, HYDROCHLOROTHIAZIDE
MICARDIS, TELMISARTAN
MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
MICONAZOLE 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
MICORT-HC, HYDROCORTISONE ACETATE
MICROGESTIN 1.5/30, ETHINYL ESTRADIOL
MICROGESTIN 1/20, ETHINYL ESTRADIOL
MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
MICRO-K 10, POTASSIUM CHLORIDE
MICRO-K, POTASSIUM CHLORIDE
MICRONOR, NORETHINDRONE
MICROZIDE, HYDROCHLOROTHIAZIDE
MIDAMOR, AMILORIDE HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIDOL LIQUID GELS, IBUPROFEN (OTC)
MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIFEPREX, MIFEPRISTONE
MIGERGOT, CAFFEINE
MIGRALAN, DIHYDROERGOTAMINE MESYLATE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE

APPENDIX A - PRODUCT NAME INDEX

A - 35

**** M ****

MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE, MILRINONE LACTATE
MINIPRESS, PRAZOSIN HYDROCHLORIDE
MINIRIN, DESMOPRESSIN ACETATE
MINITRAN, NITROGLYCERIN
MINOCIN, MINOCYCLINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL, MINOXIDIL
MINOXIDIL, MINOXIDIL (OTC)
MIOCHOL-E, ACETYLCHOLINE CHLORIDE
MIOSTAT, CARBACHOL
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
MIRCETTE, DESOGESTREL
MIRENA, LEVONORGESTREL
MIRTAZAPINE, MIRTAZAPINE
MISOPROSTOL, MISOPROSTOL
MITOMYCIN, MITOMYCIN
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
MOBIC, MELOXICAM
MODICON 28, ETHINYL ESTRADIOL
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
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)
MONISTAT-3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
MONODOX, DOXYCYCLINE
MONOKET, ISOSORBIDE MONONITRATE
MONOPRIL, FOSINOPRIL SODIUM
MONUROL, FOSFOMYCIN TROMETHAMINE
MORPHINE SULFATE, MORPHINE SULFATE
MOTOFEN, ATROPINE SULFATE
MOTRIN IB, IBUPROFEN (OTC)
MOTRIN MIGRAINE PAIN, IBUPROFEN (OTC)
MOVIPREP, ASCORBIC ACID
MOXATAG, AMOXICILLIN
MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
MOZOBIL, PLERIXAFOR
MPI DMSA KIDNEY REAGENT, TECHNETIUM TC-99M SUCCIMER KIT
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
MS CONTIN, MORPHINE SULFATE
MUCINEX D, GUAIFENESIN (OTC)
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
MUCINEX, GUAIFENESIN (OTC)
MULTAQ, DRONEDARONE HYDROCHLORIDE
MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
MULTIHANCE, GADOBENATE DIMEGLUMINE
MUPIROCIN, MUPIROCIN
MUSE, ALPROSTADIL
MUSTARGEN, MECHLORETHAMINE HYDROCHLORIDE
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
MYCAMINE, MICAFUNGIN SODIUM

APPENDIX A - PRODUCT NAME INDEX

A - 36

**** M ****

MYCELEX, CLOTRIMAZOLE
 MYCELEX-7 COMBINATION PACK, CLOTRIMAZOLE (OTC)
 MYCELEX-7, CLOTRIMAZOLE (OTC)
 MYCOBUTIN, RIFABUTIN
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 MYDRIACYL, TROPICAMIDE
 MYFORTIC, MYCOPHENOLIC ACID
 MYLERAN, BUSULFAN
 MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYOVIEW, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYSOLINE, PRIMIDONE
 MYTELASE, AMBENONIUM CHLORIDE
 M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)

**** N ****

NABUMETONE, NABUMETONE
 NADOLOL AND BENDROFLUMETHAZIDE, BENDROFLUMETHIAZIDE
 NADOLOL AND BENDROFLUMETHAZIDE, BENDROFLUMETHIAZIDE
 NADOLOL, NADOLOL
 NAFCILLIN SODIUM, NAFCILLIN SODIUM
 NAFTIN, NAFTIFINE HYDROCHLORIDE
 NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
 NALFON, FENOPROFEN CALCIUM
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM
 NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 NAMENDA XR, MEMANTINE HYDROCHLORIDE
 NAMENDA, MEMANTINE HYDROCHLORIDE
 NANDROLONE DECANOATE, NANDROLONE DECANOATE
 NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPHAZOLINE HYDROCHLORIDE, NAPHAZOLINE HYDROCHLORIDE
 NAPHCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NAPRELAN, NAPROXEN SODIUM
 NAPROSYN, NAPROXEN
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 NAPROXEN, NAPROXEN
 NARATRIPTAN, NARATRIPTAN
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NARDIL, PHENELZINE SULFATE
 NAROPIN, ROPIVACAINE HYDROCHLORIDE MONOHYDRATE
 NASACORT AQ, TRIAMCINOLONE ACETONIDE
 NASCOBAL, CYANOCOBALAMIN
 NASONEX, MOMETASONE FUROATE MONOHYDRATE
 NATACYN, NATAMYCIN
 NATAZIA, ESTRADIOL VALERATE
 NATEGLINIDE, NATEGLINIDE
 NATRECOR, NESIRITIDE RECOMBINANT
 NATROBA, SPINOSAD
 NAVANE, THIOTHIXENE
 NAVELBINE, VINORELBINE TARTRATE
 NAVSTEL, CALCIUM CHLORIDE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
 NEO-FRADIN, NEOMYCIN SULFATE
 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

APPENDIX A - PRODUCT NAME INDEX

A - 37

** N **

NEOMYCIN SULFATE, NEOMYCIN SULFATE
NEOPAP, ACETAMINOPHEN (OTC)
NEOPROFEN, IBUPROFEN LYSINE
NEORAL, CYCLOSPORINE
NEO-RX, NEOMYCIN SULFATE
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
NEOSPORIN, GRAMICIDIN
NEPHRAMINE 5.4%, AMINO ACIDS
NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
NEURONTIN, GABAPENTIN
NEVANAC, NEPAFENAC
NEXAVAR, SORAFENIB TOSYLATE
NEXCEDE, KETOPROFEN (OTC)
NEXIUM IV, ESOMEPRAZOLE SODIUM
NEXIUM, ESOMEPRAZOLE MAGNESIUM
NEXPLANON, ETONOGESTREL
NEXTERONE, AMIODARONE HYDROCHLORIDE
NIACIN, NIACIN
NIACOR, NIACIN
NIASPAN, NIACIN
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NICODERM CQ, NICOTINE (OTC)
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
NICOTINE, NICOTINE (OTC)
NICOTROL, NICOTINE
NIFEDIPINE, NIFEDIPINE
NILANDRON, NILUTAMIDE
NILSTAT, NYSTATIN
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMBEX, CISATRACURIUM BESYLATE
NIMODIPINE, NIMODIPINE
NIPENT, PENTOSTATIN
NIRAVAM, ALPRAZOLAM
NISOLDIPINE, NISOLDIPINE
NITHIODOTE, SODIUM NITRITE
NITRO-DUR, NITROGLYCERIN
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
NITROGLYCERIN, NITROGLYCERIN
NITROLINGUAL PUMPSPRAY, NITROGLYCERIN
NITROMIST, NITROGLYCERIN
NITROPRESS, SODIUM NITROPRUSSIDE
NITROSTAT, NITROGLYCERIN
NIX, PERMETHRIN (OTC)
NIZATIDINE, NIZATIDINE
NIZORAL A-D, KETOCONAZOLE (OTC)
NIZORAL, KETOCONAZOLE
NORCO, ACETAMINOPHEN
NORDETTE-28, ETHINYL ESTRADIOL
NORDITROPIN FLEXPRO, SOMATROPIN RECOMBINANT
NORDITROPIN NORDIFLEX, SOMATROPIN RECOMBINANT
NORDITROPIN, SOMATROPIN RECOMBINANT
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
NORETHIDRONE, NORETHINDRONE
NORETHIN 1/35E-21, ETHINYL ESTRADIOL
NORETHIN 1/35E-28, ETHINYL ESTRADIOL
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

A - 38

** N **

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL (7/14), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE, NORETHINDRONE
 NORFLEX, ORPHENADRINE CITRATE
 NORGESIC FORTE, ASPIRIN
 NORGESIC, ASPIRIN
 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
 NOROXIN, NORFLOXACIN
 NORPACE CR, DISOPYRAMIDE PHOSPHATE
 NORPACE, DISOPYRAMIDE PHOSPHATE
 NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
 NOR-QD, NORETHINDRONE
 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 BEZYLATE
 NORVIR, RITONAVIR
 NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
 NOVOLOG, INSULIN ASPART RECOMBINANT
 NOXAFILE, POSACONAZOLE
 NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
 NUCYNTA, TAPENTADOL HYDROCHLORIDE
 NUDEXTA, DEXTROMETHORPHAN HYDROBROMIDE
 NULYTLY, POLYETHYLENE GLYCOL 3350
 NULYTLY-FLAVORED, POLYETHYLENE GLYCOL 3350
 NUTRACORT, HYDROCORTISONE
 NUTRESTORE, GLUTAMINE
 NUTRILIPID 10%, SOYBEAN OIL
 NUTRILIPID 20%, SOYBEAN OIL
 NUTROPIN AQ PEN, SOMATROPIN RECOMBINANT
 NUTROPIN AQ, SOMATROPIN RECOMBINANT
 NUTROPIN, SOMATROPIN RECOMBINANT
 NUVARING, ETHINYL ESTRADIOL
 NUVIGIL, ARMODAFINIL
 NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
 NYSTATIN, NYSTATIN
 NYSTOP, NYSTATIN

** O **

OCTOCAINE, EPINEPHRINE
 OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OCUCLEAR, OXYMETAZOLINE HYDROCHLORIDE (OTC)
 OCUFEN, FLURBIPROFEN SODIUM

APPENDIX A - PRODUCT NAME INDEX

A - 39

** O **

OCUFLOX, OFLOXACIN
OCUPRESS, CARTEOLOL HYDROCHLORIDE
OFIRMEV, ACETAMINOPHEN
OFLOXACIN, OFLOXACIN
OFORTA, FLUDARABINE PHOSPHATE
OGEN .625, ESTROPIPATE
OGEN 1.25, ESTROPIPATE
OGEN 2.5, ESTROPIPATE
OGEN 5, ESTROPIPATE
OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
OLANZAPINE, OLANZAPINE
OLEPTRO, TRAZODONE HYDROCHLORIDE
OLUX E, CLOBETASOL PROPIONATE
OLUX, CLOBETASOL PROPIONATE
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
OMEPRAZOLE, OMEPRAZOLE
OMEPRAZOLE, OMEPRAZOLE (OTC)
OMNARIS, CICLESONIDE
OMNIPAQ 140, IOHEXOL
OMNIPAQ 180, IOHEXOL
OMNIPAQ 240, IOHEXOL
OMNIPAQ 300, IOHEXOL
OMNIPAQ 350, IOHEXOL
OMNIPRED, PREDNISOLONE ACETATE
OMNISCAN, GADODIAMIDE
OMNITROPE, SOMATROPIN RECOMBINANT
ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE AND SODIUM CHLORIDE IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON
ONFI, CLOBAZAM
ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
ONMEL, ITRACONAZOLE
ONSOLIS, FENTANYL CITRATE
OPANA ER, OXYMORPHONE HYDROCHLORIDE
OPANA, OXYMORPHONE HYDROCHLORIDE
OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
OPHTHAINE, PROPARACAIN HYDROCHLORIDE
OPHTHETIC, PROPARACAIN HYDROCHLORIDE
OPTICROM, CROMOLYN SODIUM
OPTIMARK IN PLASTIC CONTAINER, GADOVERSETAMIDE
OPTIMARK, GADOVERSETAMIDE
OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
OPTIRAY 240, IOVERSOL
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL
OPTIRAY 350, IOVERSOL
OPTISON, ALBUMIN HUMAN
OPTIVAR, AZELASTINE HYDROCHLORIDE
ORABASE HCA, HYDROCORTISONE ACETATE
ORACEA, DOXYCYCLINE
ORAMORPH SR, MORPHINE SULFATE
ORAP, PIMOZIDE
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
ORAPRED, PREDNISOLONE SODIUM PHOSPHATE
ORAQIX, LIDOCAINE
ORAVESE, PHENTOLAMINE MESYLATE
ORAVIG, MICONAZOLE
ORETIC, HYDROCHLOROTHIAZIDE
ORFADIN, NITISINONE
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN

APPENDIX A - PRODUCT NAME INDEX

A - 40

** O **

ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 ORSYTHIA, ETHINYL ESTRADIOL
 ORTHO CYCLEN-28, ETHINYL ESTRADIOL
 ORTHO EVRA, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
 ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
 ORTHO-CEPT, DESOGESTREL
 ORTHO-EST, ESTROPIPATE
 ORTHO-NOVUM 1/35-28, ETHINYL ESTRADIOL
 ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
 ORVATEN, MIDODRINE HYDROCHLORIDE
 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
 OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 OTICAIR, HYDROCORTISONE
 OVCON-35, ETHINYL ESTRADIOL
 OVCON-50, ETHINYL ESTRADIOL
 OVIDE, MALATHION
 OVIDREL, CHORIOGONADOTROPIN ALFA
 OXACILLIN SODIUM, OXACILLIN SODIUM
 OXALIPLATIN, OXALIPLATIN
 OXANDRIN, OXANDROLONE
 OXANDROLONE, OXANDROLONE
 OXaprozin, OXaprozin
 OXAZEPAM, OXAZEPAM
 OXCARBAZEPINE, OXCARBAZEPINE
 OXECTA, OXYCODONE HYDROCHLORIDE
 OXILAN-300, IOXILAN
 OXILAN-350, IOXILAN
 OXISTAT, OXICONAZOLE NITRATE
 OXSORALEN, METHOXSALEN
 OXSORALEN-ULTRA, METHOXSALEN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCET, ACETAMINOPHEN
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 OXYCONTIN, OXYCODONE HYDROCHLORIDE
 OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 OXYTOCIN, OXYTOCIN
 OXYTROL, OXYBUTYNIN
 OZURDEX, DEXAMETHASONE

** P **

PACERONE, AMIODARONE HYDROCHLORIDE
 PACLITAXEL, PACLITAXEL
 PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PAMINE FORTE, METHSCOPOLAMINE BROMIDE
 PAMINE, METHSCOPOLAMINE BROMIDE
 PANCREAZE, LIPASE
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PANDEL, HYDROCORTISONE PROBUTATE
 PANRETIN, ALITRETNINOIN
 PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 PARAFON FORTE DSC, CHLORZOXAZONE

APPENDIX A - PRODUCT NAME INDEX

A - 41

** P **

PARAGARD T 380A, COPPER
PARCOPA, CARBIDOPA
PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
PARICALCITOL, PARICALCITOL
PARLODEL, BROMOCRIPTINE MESYLATE
PARNATE, TRANYLCYPROMINE SULFATE
PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PASER, AMINOSALICYLIC ACID
PATADAY, OLOPATADINE HYDROCHLORIDE
PATANASE, OLOPATADINE HYDROCHLORIDE
PATANOL, OLOPATADINE HYDROCHLORIDE
PAXIL CR, PAROXETINE HYDROCHLORIDE
PAXIL, PAROXETINE HYDROCHLORIDE
PCE, ERYTHROMYCIN
PEDIAMYCIN 400, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAMYCIN, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
PEDIATRIC ADVIL, IBUPROFEN (OTC)
PEDIOTIC, HYDROCORTISONE
PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
PEG-3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE, POLYETHYLENE GLYCOL 3350
PEGANONE, ETHOTOIN
PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
PENICILLIN G POTASSIUM, 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
PENNNSAID, DICLOFENAC SODIUM
PENTAM, PENTAMIDINE ISETHIONATE
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
PENTASA, MESALAMINE
PENTAZOCINE AND NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
PENTAZOCINE AND NALOXONE HYDROCHLORIDES, NALOXONE HYDROCHLORIDE
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PENTOSTATIN, PENTOSTATIN
PENTOXIFYLLINE, PENTOXIFYLLINE
PENTOXIL, PENTOXIFYLLINE
PEPCID AC (GELTAB), FAMOTIDINE (OTC)
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
PEPCID, FAMOTIDINE
PERCO CET, ACETAMINOPHEN
PERCODAN, ASPIRIN
PERFOROMIST, FORMOTEROL FUMARATE
PERIDEX, CHLORHEXIDINE GLUCONATE
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
PERIOCHIP, CHLORHEXIDINE GLUCONATE
PERIOGARD, CHLORHEXIDINE GLUCONATE
PERIOSTAT, DOXYCYCLINE HYCLATE
PERMAPEN, PENICILLIN G BENZATHINE
PERMETHRIN, PERMETHRIN
PERMETHRIN, PERMETHRIN (OTC)
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PERPHENAZINE, PERPHENAZINE
PERSANTINE, DIPYRIDAMOLE
PEXEGA, PAROXETINE MESYLATE
PFIZERPEN, PENICILLIN G POTASSIUM
PHARMASEAL SCRUB CARE, CHLORHEXIDINE GLUCONATE (OTC)
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENELZINE SULFATE, PHENELZINE SULFATE

APPENDIX A - PRODUCT NAME INDEX

A - 42

** P **

PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PHENTERMINE RESIN COMPLEX, PHENTERMINE RESIN COMPLEX
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PHENYTEK, PHENYTOIN SODIUM
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHENYTOIN, PHENYTOIN
PHILITH, ETHINYL ESTRADIOL
PHISOHEX, HEXACHLOROPHENE
PHOSLO GELCAPS, CALCIUM ACETATE
PHOSLYRA, CALCIUM ACETATE
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
PHOTOFRIN, PORFIMER SODIUM
PHRENILIN FORTE, ACETAMINOPHEN
PHRENILIN WITH CAFFEINE AND CODEINE, ACETAMINOPHEN
PHRENILIN, ACETAMINOPHEN
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYTONADIONE, PHYTONADIONE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PILOPINE HS, PILOCARPINE HYDROCHLORIDE
PINDOLOL, PINDOLOL
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PIPERACILLIN, PIPERACILLIN SODIUM
PIROXICAM, PIROXICAM
PITOCIN, OXYTOCIN
PLAN B ONE-STEP, LEVONORGESTREL
PLAN B ONE-STEP, LEVONORGESTREL (OTC)
PLAN B, LEVONORGESTREL
PLAN B, LEVONORGESTREL (OTC)
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
PLASMA-LYTE 56 IN PLASTIC CONTAINER, MAGNESIUM ACETATE TETRAHYDRATE
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLASMA-LYTE R IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLAVIX, CLOPIDOGREL BISULFATE
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLENDIL, FELODIPIINE
PLETAL, CILOSTAZOL
PODOFILOX, PODOFILOX
POLOCAINE, MEPIVACAINE HYDROCHLORIDE
POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
POLYMYCIN B SULFATE, POLYMYXIN B SULFATE
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POLY-PRED, NEOMYCIN SULFATE
POLY-RX, POLYMYXIN B SULFATE
POLYTRIM, POLYMYXIN B SULFATE
PONSTEL, MEFENAMIC ACID
PORTIA-28, ETHINYL ESTRADIOL
POTASSIUM ACETATE IN PLASTIC CONTAINER, POTASSIUM ACETATE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX A - PRODUCT NAME INDEX

A - 43

*** P ***

APPENDIX A - PRODUCT NAME INDEX

A - 44

** P **

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
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, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
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, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
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 CHLORIDE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
POTASSIUM CITRATE, POTASSIUM CITRATE
POTIGA, EZOGABINE
POVIDONE IODINE, POVIDONE-IODINE (OTC)
PRADAXA, DABIGATRAN ETEXILATE MESYLATE
PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRAMOSONE, HYDROCORTISONE ACETATE
PRANDIMET, METFORMIN HYDROCHLORIDE
PRANDIN, REPAGLINIDE
PRAVACHOL, PRAVASTATIN SODIUM
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
PRECOSE, ACARBOSE
PRED FORTE, PREDNISOLONE ACETATE
PRED MILD, PREDNISOLONE ACETATE
PRED-G, GENTAMICIN SULFATE
PREDNICARBATE, PREDNICARBATE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
PREDNISONE INTENSOL, PREDNISONE

APPENDIX A - PRODUCT NAME INDEX

A - 45

** P **

PREDNISONE, PREDNISONE
PREFEST, ESTRADIOL
PREGNYL, GONADOTROPIN, CHORIONIC
PRELONE, PREDNISOLONE
PREMARIN, ESTROGENS, CONJUGATED
PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
PREMPHASE 14/14, ESTROGENS, CONJUGATED
PREMPRO, ESTROGENS, CONJUGATED
PRE-OP II, HEXACHLOROPHENE
PRE-OP, HEXACHLOROPHENE
PRE-PEN, BENZYL PENICILLOYL POLYLYSINE
PREPIDIL, DINOPROSTONE
PREVACID 24 HR, LANSOPRAZOLE (OTC)
PREVACID, LANSOPRAZOLE
PREVALITE, CHOLESTYRAMINE
PREVIFEM, ETHINYL ESTRADIOL
PREVPAC, AMOXICILLIN
PREZISTA, DARUNAVIR ETHANOLATE
PRIALT, ZICONOTIDE ACETATE
PRIFTIN, RIFAPENTINE
PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
PRILOSEC, OMEPRAZOLE
PRILOSEC, OMEPRAZOLE MAGNESIUM
PRIMAQUINE, PRIMAQUINE PHOSPHATE
PRIMAXIN, CILASTATIN SODIUM
PRIMIDONE, PRIMIDONE
PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE
PRINVIL, LISINOPRIL
PRINZIDE, HYDROCHLORTIAZIDE
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
PROAMATINE, MIDODRINE HYDROCHLORIDE
PROBALAN, PROBENECID
PROBENECID AND COLCHICINE, COLCHICINE
PROBENECID, PROBENECID
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCALAMINE, AMINO ACIDS
PROCARDIA XL, NIFEDIPINE
PROCARDIA, NIFEDIPINE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROCHLORPERAZINE, PROCHLORPERAZINE
PROCOMP, PROCHLORPERAZINE MALEATE
PROCTOFOAM HC, HYDROCORTISONE ACETATE
PROFEN, IBUPROFEN (OTC)
PROFERDEX, IRON DEXTRAN
PROGESTERONE, PROGESTERONE
PROGLYCEM, DIAZOXIDE
PROGRAF, TACROLIMUS
PROHANCE MULTIPACK, GADOTERIDOL

APPENDIX A - PRODUCT NAME INDEX

A - 46

** P **

PROHANCE, GADOTERIDOL
 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 AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE
 PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 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
 PROparacaine HYDROCHLORIDE, PROparacaine HYDROCHLORIDE
 PROpecia, FINASTERIDE
 PROPINE, DIPIVEFRIN HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 PROpranolol HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 PROpranolol HYDROCHLORIDE, PROpranolol HYDROCHLORIDE
 PROPYlTHIouracil, PROPYlTHIouracil
 PROSCAR, FINASTERIDE
 PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
 PROSTIN E2, DINOPROSTONE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PROTONIX IV, PANTOPRAZOLE SODIUM
 PROTONIX, PANTOPRAZOLE SODIUM
 PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
 PROtopic, TACROLIMUS
 PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 PROVENTIL-HFA, ALBUTEROL SULFATE
 PROVERA, MEDROXYPROGESTERONE ACETATE
 PROVIGIL, MODAFINIL
 PROVOCHOLINE, METHACHOLINE CHLORIDE
 PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
 PROZAC, FLUOXETINE HYDROCHLORIDE
 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 PULMICORT FLEXHALER, BUDESONIDE
 PULMICORT RESPULES, BUDESONIDE
 PULMICORT, BUDESONIDE
 PULMOLITE, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 PURINETHOL, MERCAPTOPURINE
 PUR-WASH, PURIFIED WATER (OTC)
 PYLERA, BISMUTH SUBCITRATE POTASSIUM
 PYRAZINAMIDE, PYRAZINAMIDE
 PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 PYTEST KIT, UREA, C-14
 PYTEST, UREA, C-14

** Q **

QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM
 QUALAQUIN, QUININE SULFATE
 QUASENSE, ETHINYL ESTRADIOL
 QUELICIN PRESERVATIVE FREE, SUCCINYLCHOLINE CHLORIDE
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 47

** Q **

QUINARETIC, HYDROCHLOROTHIAZIDE
 QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 QUINIDINE SULFATE, QUINIDINE SULFATE
 QUIXIN, LEVOFLOXACIN
 QUTENZA, CAPSAICIN
 QVAR 40, BECLOMETHASONE DIPROPIONATE
 QVAR 80, BECLOMETHASONE DIPROPIONATE

** R **

RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)
 RAMIPRIL, RAMIPRIL
 RANEXA, RANOLAZINE
 RANICLOR, CEFACLOR
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 RAPAFLO, SILODOSIN
 RAPAMUNE, SIROLIMUS
 RAZADYNE ER, GALANTAMINE HYDROBROMIDE
 RAZADYNE, GALANTAMINE HYDROBROMIDE
 REBETOL, RIBAVIRIN
 RECLAST, ZOLEDRONIC ACID
 RECTIV, NITROGLYCERIN
 REFLUDAN, LEPIRUDIN RECOMBINANT
 REGITINE, PHENTOLAMINE MESYLATE
 REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 REGIONOL, PYRIDOSTIGMINE BROMIDE
 RELENZA, ZANAMIVIR
 RELISTOR, METHYLNALTREXONE BROMIDE
 RELPAX, ELETRIPTAN HYDROBROMIDE
 REMERON SOLTAB, MIRTAZAPINE
 REMERON, MIRTAZAPINE
 REMODULIN, TREPROSTINIL SODIUM
 RENACIDIN, CITRIC ACID
 RENAGEL, SEVELAMER HYDROCHLORIDE
 RENAMIN W/O ELECTROLYTES, AMINO ACIDS
 RENOGRAFIN-76, DIATRIZOATE MEGLUMINE
 RENOVA, TRETINOIN
 RENVELA, SEVELAMER CARBONATE
 REPREXAIN, HYDROCODONE BITARTRATE
 REPRONEX, LUTEINIZING HORMONE
 REQUIP XL, ROPINIROLE HYDROCHLORIDE
 REQUIP, ROPINIROLE HYDROCHLORIDE
 DESCRIPTOR, DELAVIRDINE MESYLATE
 RESECTISOL IN PLASTIC CONTAINER, MANNITOL
 RESERPINE, RESERPINE
 RESTASIS, CYCLOSPORINE
 RESTORIL, TEMAZEPAM
 RETIN-A MICRO, TRETINOIN
 RETIN-A, TRETINOIN
 RETISERT, FLUOCINOLONE ACETONIDE
 RETROVIR, ZIDOVUDINE
 REVATIO, SILDENAFIL CITRATE
 REVIA, NALTREXONE HYDROCHLORIDE
 REVIMID, LENALIDOMIDE
 REYATAZ, ATAZANAVIR SULFATE
 REZIRA, HYDROCODONE BITARTRATE
 R-GENE 10, ARGININE HYDROCHLORIDE
 RHINOCORT, BUDESONIDE
 RIBASPHERE, RIBAVIRIN
 RIBAVARIN, RIBAVIRIN
 RIBAVIRIN, RIBAVIRIN
 RID MOUSSE, PIPERONYL BUTOXIDE (OTC)
 RIDAURA, AURANOFIN

APPENDIX A - PRODUCT NAME INDEX

A - 48

**** R ****

RIFADIN, RIFAMPIN
 RIFAMATE, ISONIAZID
 RIFAMPIN AND ISONIAZID, ISONIAZID
 RIFAMPIN, RIFAMPIN
 RIFATER, ISONIAZID
 RILUTEK, RILUZOLE
 RILUZOLE, RILUZOLE
 RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 RIMSO-50, DIMETHYL SULFOXIDE
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 RIOMET, METFORMIN HYDROCHLORIDE
 RISEDRONATE SODIUM, RISEDRONATE SODIUM
 RISPERDAL CONSTA, RISPERIDONE
 RISPERDAL, RISPERIDONE
 RISPERIDONE, RISPERIDONE
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN-SR, METHYLPHENIDATE HYDROCHLORIDE
 RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 ROBAXIN, METHOCARBAMOL
 ROBAXIN-750, METHOCARBAMOL
 ROBINUL FORTE, GLYCOPYRROLATE
 ROBINUL, GLYCOPYRROLATE
 ROCALTROL, CALCITRIOL
 ROCEPHIN, CEFTRIAXONE SODIUM
 ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 ROGAINE (FOR MEN), MINOXIDIL (OTC)
 ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
 ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 ROMAZICON, FLUMAZENIL
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 ROWASA, MESALAMINE
 ROXICET 5/500, ACETAMINOPHEN
 ROXICET, ACETAMINOPHEN
 ROXICODONE, OXYCODONE HYDROCHLORIDE
 ROXILOX, ACETAMINOPHEN
 ROZEREM, RAMELTEON
 RYTHMOL SR, PROPAFENONE HYDROCHLORIDE
 RYTHMOL, PROPAFENONE HYDROCHLORIDE
 RYZOLT, TRAMADOL HYDROCHLORIDE

**** S ****

SABRIL, VIGABATRIN
 SAFYRAL, DROSPIRENONE
 SAIZEN, SOMATROPIN RECOMBINANT
 SALAGEN, PILOCARPINE HYDROCHLORIDE
 SALONPAS, MENTHOL (OTC)
 SALURON, HYDROFLUMETHIAZIDE
 SAMSCA, TOLVAPTAN
 SANCTURA XR, TROSPiUM CHLORIDE
 SANCTURA, TROSPiUM CHLORIDE
 SANCUSO, GRANisetron
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 SAPHRIS, ASENAPINE MALEATE
 SARAFEM, FLUOXETINE HYDROCHLORIDE
 SAVELLA, MILNACIPRAN HYDROCHLORIDE
 SCANDONEST L, LEVONORDEFRIN
 SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
 SCANLUX-300, IOPAMIDOL
 SCANLUX-370, IOPAMIDOL
 SCLEROSOL, TALC

APPENDIX A - PRODUCT NAME INDEX

A - 49

** S **

SEASONALE, ETHINYL ESTRADIOL
SEASONIQUE, ETHINYL ESTRADIOL
SECONAL SODIUM, SECOBARBITAL SODIUM
SECTRAL, ACEBUTOLOL HYDROCHLORIDE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SELENIUM SULFIDE, SELENIUM SULFIDE
SELSUN, SELENIUM SULFIDE
SELZENTRY, MARAVIROC
SEMPREX-D, ACRIVASTINE
SENSIPAR, CINACALCET HYDROCHLORIDE
SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
SEPTOCaine, ARTICAINE HYDROCHLORIDE
SEPTRA DS, SULFAMETHOXAZOLE
SEPTRA, SULFAMETHOXAZOLE
SEREVENT, SALMETEROL XINAFOATE
SEROMYCIN, CYCLOSERINE
SEROPHENE, CLOMIPHENE CITRATE
SEROQUEL XR, QUETIAPINE FUMARATE
SEROQUEL, QUETIAPINE FUMARATE
SEROSTIM, SOMATROPIN RECOMBINANT
SERPALAN, RESERPINE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SEVOFLURANE, SEVOFLURANE
SFROWASA, MESALAMINE
SHADE UVAGUARD, AVOBENZONE (OTC)
SILENOR, DOXEPEPIN HYDROCHLORIDE
SILVADENE, SILVER SULFADIAZINE
SIMCOR, NIACIN
SIMVASTATIN, SIMVASTATIN
SINE-AID IB, IBUPROFEN (OTC)
SINEMET CR, CARBIDOPA
SINEMET, CARBIDOPA
SINGULAIR, MONTELUKAST SODIUM
SINOGRAFIN, DIATRIZOATE MEGLUMINE
SKELAXIN, METAXALONE
SKELID, TILUDRONATE DISODIUM
SODIUM ACETATE IN PLASTIC CONTAINER, SODIUM ACETATE ANHYDROUS
SODIUM BICARBONATE, SODIUM BICARBONATE
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
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX
SODIUM IODIDE I-123, SODIUM IODIDE I-123
SODIUM IODIDE I-131, SODIUM IODIDE I-131
SODIUM IODIDE I-131, SODIUM IODIDE I-131
SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SOJOURN, SEVOFLURANE
SOLAGE, MEQUINOL
SOLARAZE, DICLOFENAC SODIUM
SOLODYN, MINOCYCLINE HYDROCHLORIDE
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
SOMA COMPOUND W/ CODEINE, ASPIRIN
SOMA COMPOUND, ASPIRIN
SOMA, CARISOPRODOL

APPENDIX A - PRODUCT NAME INDEX

A - 50

**** S ****

SOMATULINE DEPOT, LANREOTIDE ACETATE
SOMAVERT, PEGVISOMANT
SONATA, ZALEPLON
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
SOTRET, ISOTRETINOIN
SPECTRACEF, CEFEDITOREN PIVOXIL
SPIRIVA, TIOTROPIUM BROMIDE MONOHYDRATE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPIRONOLACTONE, SPIRONOLACTONE
SPORANOX, ITRACONAZOLE
SPRINTEC, ETHINYL ESTRADIOL
SPRIX, KETOROLAC TROMETHAMINE
SPRYCEL, DASATINIB
SPS, SODIUM POLYSTYRENE SULFONATE
SSD AF, SILVER SULFADIAZINE
SSD, SILVER SULFADIAZINE
STADOL PRESERVATIVE FREE, BUTORPHANOL TARTRATE
STADOL, BUTORPHANOL TARTRATE
STALEVO 100, CARBIDOPA
STALEVO 125, CARBIDOPA
STALEVO 150, CARBIDOPA
STALEVO 200, CARBIDOPA
STALEVO 50, CARBIDOPA
STALEVO 75, CARBIDOPA
STARLIX, NATEGLINIDE
STAVUDINE, STAVUDINE
STAVZOR, VALPROIC ACID
STAXYN, VARDENAFIL HYDROCHLORIDE
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
STIE-CORT, HYDROCORTISONE
STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
STRATTERA, ATOMOXETINE HYDROCHLORIDE
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
STRIANT, TESTOSTERONE
STROMECTOL, IVERMECTIN
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89
SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
SUBOXONE, BUPRENORPHINE
SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
SUBUTEX, BUPRENORPHINE HYDROCHLORIDE
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 AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
SULFADIAZINE, SULFADIAZINE
SULFAMETHOPRIM, SULFAMETHOXAZOLE
SULFAMETHOPRIM-DS, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE

APPENDIX A - PRODUCT NAME INDEX

A - 51

**** S ****

SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 SULFAMYRON, MAFENIDE ACETATE
 SULFASALAZINE, SULFASALAZINE
 SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
 SULINDAC, SULINDAC
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SUMAVEL DOSEPRO, SUMATRIPTAN SUCCINATE
 SUPPRELIN LA, HISTRELIN ACETATE
 SUPRANE, DESFLURANE
 SUPRAX, CEFIXIME
 SUPRENZA, PHENTERMINE HYDROCHLORIDE
 SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE ANHYDROUS
 SURMONTIL, TRIMIPRAMINE MALEATE
 SURVANTA, BERACTANT
 SUSTIVA, EFAVIRENZ
 SUTENT, SUNITINIB MALATE
 SYEDA, DROSPIRENONE
 SYMBICORT, BUDESONIDE
 SYMBYAX, FLUOXETINE HYDROCHLORIDE
 SYMLIN, PRAMLINTIDE ACETATE
 SYNACORT, HYDROCORTISONE
 SYNALAR, FLUOCINOLONE ACETONIDE
 SYNALGOS-DC, ASPIRIN
 SYNAREL, NAFARELIN ACETATE
 SYNERA, LIDOCAINE
 SYNERCID, DALFOPRISTIN
 SYNTHETIC CONJUGATED ESTROGENS A, ESTROGENS, CONJUGATED SYNTHETIC A
 SYNTHROID, LEVOTHYROXINE SODIUM
 SPRINE, TRIENTINE HYDROCHLORIDE

**** T ****

TAB-PROFEN, IBUPROFEN (OTC)
 TACLONEX SCALP, BETAMETHASONE DIPROPIONATE
 TACLONEX, BETAMETHASONE DIPROPIONATE
 TACROLIMUS, TACROLIMUS
 TAGAMET HB, CIMETIDINE (OTC)
 TAGAMET, CIMETIDINE
 TALC, TALC
 TALWIN, PENTAZOCINE LACTATE
 TAMBOCOR, FLECAINIDE ACETATE
 TAMIFLU, OSSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TAPAZOLE, METHIMAZOLE
 TARCEVA, ERLOTINIB HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARKA, TRANDOLAPRIL
 TASIGNA, NILOTINIB HYDROCHLORIDE MONOHYDRATE
 TASMAR, TOLCAPONE
 TAVIST ALLERGY/SINUS/HEADACHE, ACETAMINOPHEN (OTC)
 TAVIST-1, CLEMASTINE FUMARATE (OTC)
 TAXOTERE, DOCETAXEL
 TAZICEF, CEFTAZIDIME
 TAZORAC, TAZAROTENE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT
 TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99 SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT

APPENDIX A - PRODUCT NAME INDEX

A - 52

**** T ****

TEFLARO, CEFTAROLINE FOSAMIL
TEGRETOL, CARBAMAZEPINE
TEGRETOL-XR, CARBAMAZEPINE
TEKAMLO, ALISKIREN HEMIFUMARATE
TEKTURNA HCT, ALISKIREN HEMIFUMARATE
TEKTURNA, ALISKIREN HEMIFUMARATE
TEMAZEPAM, TEMAZEPAM
TEMODAR, TEMOZOLOMIDE
TEMOVATE E, CLOBETASOL PROPIONATE
TEMOVATE, CLOBETASOL PROPIONATE
TEMOZOLOMIDE, TEMOZOLOMIDE
TENEX, GUANFACINE HYDROCHLORIDE
TENORETIC 100, ATENOLOL
TENORETIC 50, ATENOLOL
TENORMIN, ATENOLOL
TENSILON PRESERVATIVE FREE, EDROPHONIUM CHLORIDE
TENSILON, EDROPHONIUM CHLORIDE
TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE
TENUATE, DIETHYLPROPION HYDROCHLORIDE
TERAZOL 3, TERCONAZOLE
TERAZOL 7, TERCONAZOLE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TERCONAZOLE, TERCONAZOLE
TERIL, CARBAMAZEPINE
TERRAMYCIN W/ POLYMYXIN B SULFATE, OXYTETRACYCLINE HYDROCHLORIDE
TESSALON, BENZONATATE
TESTIM, TESTOSTERONE
TESTOPEL, TESTOSTERONE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TESTRED, METHYLTESTOSTERONE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
TEVETEN HCT, EPROSARTAN MESYLATE
TEVETEN, EPROSARTAN MESYLATE
TEV-TROPIN, SOMATROPIN RECOMBINANT
TEXACORT, HYDROCORTISONE
THALITONE, CHLORTHALIDONE
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
THALOMID, THALIDOMIDE
THAM, TROMETHAMINE
THEO-24, THEOPHYLLINE
THEOCHRON, THEOPHYLLINE
THEOLAIR, 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 AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE, THEOPHYLLINE
THERMAZENE, SILVER SULFADIAZINE
THEROXIDIL, MINOXIDIL (OTC)
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
THIOGUANINE, THIOGUANINE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTEPA, THIOTEPA
THIOTHIXENE, THIOTHIXENE
THRIVE, NICOTINE POLACRILEX (OTC)
THYROGEN, THYROTROPIN ALFA
THYROLAR-0.25, LIOTHYRONINE SODIUM

APPENDIX A - PRODUCT NAME INDEX

A - 53

** T **

THYROLAR-0.5, LIOTHYRONINE SODIUM
THYROLAR-1, LIOTHYRONINE SODIUM
THYROLAR-2, LIOTHYRONINE SODIUM
THYROLAR-3, LIOTHYRONINE SODIUM
THYROSafe, POTASSIUM IODIDE (OTC)
THYROSHIELD, POTASSIUM IODIDE (OTC)
TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
TIAZAC, DILTIAZEM HYDROCHLORIDE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
TIKOSYN, DOFETILIDE
TIMENTIN IN PLASTIC CONTAINER, CLAVULANATE POTASSIUM
TIMENTIN, CLAVULANATE POTASSIUM
TIMOLOL MALEATE, TIMOLOL MALEATE
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
TIMOPTIC, TIMOLOL MALEATE
TIMOPTIC-XE, TIMOLOL MALEATE
TINDAMAX, TINIDAZOLE
TIOCONAZOLE, TIOCONAZOLE (OTC)
TIOPRONIN, TIOPRONIN
TIROSINT, LEVOTHYROXINE SODIUM
TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
TIS-U-SOL, MAGNESIUM SULFATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOBI, TOBRAMYCIN
TOBRADEX ST, DEXAMETHASONE
TOBRADEX, DEXAMETHASONE
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOBRAMYCIN, TOBRAMYCIN
TOBREX, TOBRAMYCIN
TODAY, NONOXYNOL-9 (OTC)
TOFRANIL, IMIPRAMINE HYDROCHLORIDE
TOFRANIL-PM, IMIPRAMINE PAMOATE
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TOPAMAX, TOPIRAMATE
TOPICORT LP, DESOXIMETASONE
TOPICORT, DESOXIMETASONE
TOPIRAMATE, TOPIRAMATE
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
TOPOTECAN, TOPOTECAN HYDROCHLORIDE
TOPROL-XL, METOPROLOL SUCCINATE
TORISEL, TEMSIROLIMUS
TORSEMIDE, TORSEMIDE
TOTECT, DEXRAZOXANE HYDROCHLORIDE
TOVIAZ, FESOTERODINE FUMARATE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TRACLEER, BOSENTAN
TRADJENTA, LINAGLIPTIN
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRANDATE, LABETALOL HYDROCHLORIDE
TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
TRANDOLAPRIL, TRANDOLAPRIL
TRANEXAMIC ACID, TRANEXAMIC ACID
TRANSDERM SCOP, SCOPOLAMINE
TRANXENE, CLORAZEPATE DIPOTASSIUM
TRANYLCYPROMINE SULFATE, TRANYLCYPROMINE SULFATE
TRASYLOL, APROTININ

APPENDIX A - PRODUCT NAME INDEX

A - 54

**** T ****

TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 10% W/O ELECTROLYTES, AMINO ACIDS
TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 3.5% W/ ELECTROLYTES, AMINO ACIDS
TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 5.5% W/ ELECTROLYTES, AMINO ACIDS
TRAVASOL 5.5% W/O ELECTROLYTES, AMINO ACIDS
TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 8.5% W/ ELECTROLYTES, AMINO ACIDS
TRAVASOL 8.5% W/O ELECTROLYTES, AMINO ACIDS
TRAVATAN Z, TRAVOPROST
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TREANDA, BENDAMUSTINE HYDROCHLORIDE
TRECATOR, ETHIONAMIDE
TRELSTAR, TRIPTORELIN PAMOATE
TRENTAL, PENTOXIFYLLINE
TRETINOIN, TRETINOIN
TREXALL, METHOTREXATE SODIUM
TREXIMET, NAPROXEN SODIUM
TRI LO SPRINTEC, ETHINYLMESTRADIOL
TRIACET, TRIAMCINOLONE ACETONIDE
TRIACIN-C, CODEINE PHOSPHATE
TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIAMTERENE AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE
TRIAZOLAM, TRIAZOLAM
TRIBENZOR, AMLODIPINE BESYLATE
TRICOR, FENOFIBRATE
TRIDERM, TRIAMCINOLONE ACETONIDE
TRIDIONE, TRIMETHADIONE
TRIESENCE, TRIAMCINOLONE ACETONIDE
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
TRIFLURIDINE, TRIFLURIDINE
TRIGLIDE, FENOFIBRATE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
TRI-LEGEST 21, ETHINYLMESTRADIOL
TRI-LEGEST FE, ETHINYLMESTRADIOL
TRILEPTAL, OXCARBAZEPINE
TRILIPIX, CHOLINE FENOFIBRATE
TRI-LUMA, FLUOCINOLONE ACETONIDE
TRILYTE, POLYETHYLENE GLYCOL 3350
TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE, TRIMETHOBENZAMIDE HYDROCHLORIDE
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
TRIMETHOPRIM, TRIMETHOPRIM
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE
TRI-NORINYL 28-DAY, ETHINYLMESTRADIOL
TRIOSTAT, LIOTHYRONINE SODIUM
TRI-PREVIFEM, ETHINYLMESTRADIOL
TRISENOX, ARSENIC TRIOXIDE
TRI-SPRINTEC, ETHINYLMESTRADIOL
TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
TRIVARIS, TRIAMCINOLONE ACETONIDE
TRIVORA-28, ETHINYLMESTRADIOL
TRIZIVIR, ABACAVIR SULFATE
TROPHAMINE 10%, AMINO ACIDS
TROPHAMINE, AMINO ACIDS
TROPICACYL, TROPICAMIDE
TROPICAMIDE, TROPICAMIDE
TROSPiUM CHLORIDE, TROSPiUM CHLORIDE
TRUSOPT, DORZOLAMIDE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 55

** T **

TRUVADA, EMTRICITABINE
 TUSSICAPS, CHLORPHENIRAMINE POLISTIREX
 TUSSIGON, HOMATROPINE METHYLBROMIDE
 TUSSIONEX PENN KINETIC, CHLORPHENIRAMINE POLISTIREX
 TWINJECT 0.15, EPINEPHRINE
 TWINJECT 0.3, EPINEPHRINE
 TWYNSTA, AMLODIPINE BESYLATE
 TYGACIL, TIGECYCLINE
 TYKERB, LAPATINIB DITOSYLATE
 TYLENOL (CAPLET), ACETAMINOPHEN (OTC)
 TYLENOL (GELTAB), ACETAMINOPHEN (OTC)
 TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
 TYLENOL W/ CODEINE NO. 4, ACETAMINOPHEN
 TYLOX, ACETAMINOPHEN
 TYVASO, TREPROSTINIL SODIUM
 TYZEKA, TELBIVUDINE
 TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

** U **

U-CORT, HYDROCORTISONE ACETATE
 ULESFIA, BENZYL ALCOHOL
 ULORIC, FEBUXOSTAT
 ULTANE, SEVOFLURANE
 ULTIVA, REMIFENTANIL HYDROCHLORIDE
 ULTRACET, ACETAMINOPHEN
 ULTRAM ER, TRAMADOL HYDROCHLORIDE
 ULTRAM, TRAMADOL HYDROCHLORIDE
 ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
 ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 ULTRAVATE, HALOBETASOL PROPIONATE
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 150, IOPROMIDE
 ULTRAVIST 240, IOPROMIDE
 ULTRAVIST 300 IN PLASTIC CONTAINER, IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNIRETIC, HYDROCHLOROTHIAZIDE
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM
 UNIVASC, MOXIPRIL HYDROCHLORIDE
 URECHOLINE, BETHANECHOL CHLORIDE
 UREX, METHENAMINE HIPPURATE
 UROCIT-K, POTASSIUM CITRATE
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSDIOL
 URSO FORTE, URSDIOL
 URSDIOL, URSDIOL
 UVADEX, METHOXSALEN

** V **

VAGIFEM, ESTRADIOL
 VAGISTAT-1, TIOCONAZOLE (OTC)
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
 VALIUM, DIAZEPAM
 VALNAC, BETAMETHASONE VALERATE
 VALPROATE SODIUM, VALPROATE SODIUM
 VALPROIC ACID, VALPROIC ACID
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 VALTROPIN, SOMATROPIN RECOMBINANT

APPENDIX A - PRODUCT NAME INDEX

A - 56

** V **

VALTURNA, ALISKIREN HEMIFUMARATE
VANCOGIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
VANCOGIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VANDAZOLE, METRONIDAZOLE
VANDETANIB, VANDETANIB
VANIQA, EFLORNITHINE HYDROCHLORIDE
VANOS, FLUOCINONIDE
VANTAS, HISTRELIN ACETATE
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
VASERETIC, ENALAPRIL MALEATE
VASOCIDIN, PREDNISOLONE SODIUM PHOSPHATE
VASOTEC, ENALAPRIL MALEATE
VECTICAL, CALCITRIOL
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VELCADE, BORTEZOMIB
VELETRI, EPOPROSTENOL SODIUM
VELIVET, DESOGESTREL
VELTIN, CLINDAMYCIN PHOSPHATE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VENOFER, IRON SUCROSE
VENTAVIS, ILOPROST
VENTOLIN HFA, ALBUTEROL SULFATE
VERAMYST, FLUTICASONE FUROATE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VERDESO, DESONIDE
VEREGEN, SINECATECHINS
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERELAN, VERAPAMIL HYDROCHLORIDE
VESICARE, SOLIFENACIN SUCCINATE
VEXOL, RIMEXOLONE
VFEND, VORICONAZOLE
VIAGRA, SILDENAFIL CITRATE
VIBATIV, TELAVANCIN HYDROCHLORIDE
VIBISONE, CYANOCOBALAMIN
VIBRAMYCIN, DOXYCYCLINE
VIBRAMYCIN, DOXYCYCLINE CALCIUM
VIBRAMYCIN, DOXYCYCLINE HYCLATE
VICODIN ES, ACETAMINOPHEN
VICODIN HP, ACETAMINOPHEN
VICODIN, ACETAMINOPHEN
VICOPROFEN, HYDROCODONE BITARTRATE
VICTOZA, LIRAGLUTIDE RECOMBINANT
VICTRELIS, BOCEPREVIR
VIDAZA, AZACITIDINE
VIDEX EC, DIDANOSINE
VIDEX, DIDANOSINE
VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
VIBRYD, VILAZDONE HYDROCHLORIDE
VIMOVO, ESOMEPRAZOLE MAGNESIUM
VIMPAT, LACOSAMIDE
VINBLASTINE SULFATE, VINBLASTINE SULFATE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
VIRACEPT, NELFINAVIR MESYLATE
VIRAMUNE XR, NEVIRAPINE
VIRAMUNE, NEVIRAPINE
VIRAZOLE, RIBAVIRIN
VIREAD, TENOFOVIR DISOPROXIL FUMARATE
VIROPTIC, TRIFLURIDINE
VISICOL, SODIUM PHOSPHATE, DIBASIC ANHYDROUS
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISINE-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

APPENDIX A - PRODUCT NAME INDEX

A - 57

** V **

VISIONBLUE, TRYPLAN BLUE
 VISIPAQUE 270, IODIXANOL
 VISIPAQUE 320, IODIXANOL
 VISTARIL, HYDROXYZINE PAMOATE
 VISTIDE, CIDOFOVIR
 VISUDYNE, VERTEPORFIN
 VITAMIN D, ERGOCALCIFEROL
 VITAMIN K1, PHYTONADIONE
 VITRASE, HYALURONIDASE
 VITRASERT, GANCICLOVIR
 VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE
 VIVELLE, ESTRADIOL
 VIVELLE-DOT, ESTRADIOL
 VIVITROL, NALTREXONE
 VOLTAREN, DICLOFENAC SODIUM
 VOLTAREN-XR, DICLOFENAC SODIUM
 VORICONAZOLE, VORICONAZOLE
 VOSOL HC, ACETIC ACID, GLACIAL
 VOSOL, ACETIC ACID, GLACIAL
 VOSPIRE ER, ALBUTEROL SULFATE
 VOTRIENT, PAZOPANIB HYDROCHLORIDE
 VPRIV, VELAGLUCERASE ALFA
 VUMON, TENIPOSIDE
 VUSION, MICONAZOLE NITRATE
 VYTORIN, EZETIMIBE
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE

** W **

WARFARIN SODIUM, WARFARIN SODIUM
 WELCHOL, COLESEVELAM HYDROCHLORIDE
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 WELLBUTRIN, BUPROPION HYDROCHLORIDE
 WESTCORT, HYDROCORTISONE VALERATE

** X **

XALATAN, LATANOPROST
 XALKORI, CRIZOTINIB
 XANAX XR, ALPRAZOLAM
 XANAX, ALPRAZOLAM
 XARELTO, RIVAROXABAN
 XELODA, CAPECITABINE
 XENAZINE, TETRABENAZINE
 XENICAL, ORLISTAT
 XENON XE 133, XENON XE-133
 XERESE, ACYCLOVIR
 XIFAXAN, RIFAXIMIN
 XOLEGEL, KETOCONAZOLE
 XOPENEX HFA, LEVALBUTEROL TARTRATE
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 XYLOCAINE 4% PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE DENTAL WITH EPINEPHRINE, EPINEPHRINE
 XYLOCAINE DENTAL, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYREM, SODIUM OXYBATE
 XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE

** Y **

YASMIN, DROSPIRENONONE

APPENDIX A - PRODUCT NAME INDEX

A - 58

**** Y ****

YAZ, DROSPIRENONE

**** Z ****

ZADITOR, KETOTIFEN FUMARATE (OTC)
ZAFIRLUKAST, ZAFIRLUKAST
ZALEPLON, ZALEPLON
ZANAFLEX, TIZANIDINE HYDROCHLORIDE
ZANOSAR, STREPTOZOCIN
ZANTAC 150, RANITIDINE HYDROCHLORIDE
ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
ZANTAC 25, RANITIDINE HYDROCHLORIDE
ZANTAC 300, RANITIDINE HYDROCHLORIDE
ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)
ZANTAC IN PLASTIC CONTAINER, RANITIDINE HYDROCHLORIDE
ZANTAC, RANITIDINE HYDROCHLORIDE
ZARONTIN, ETHOSUXIMIDE
ZAROXOLYN, METOLAZONE
ZAVESCA, MIGLUSTAT
ZEBETA, BISOPROLOL FUMARATE
ZEGERID OTC, OMEPRAZOLE (OTC)
ZEGERID, OMEPRAZOLE
ZELAPAR, SELEGILINE HYDROCHLORIDE
ZELBORAF, VEMURAFENIB
ZEMPLAR, PARICALCITOL
ZEMURON, ROCURONIUM BROMIDE
ZENPEP, LIPASE
ZERIT, STAVUDINE
ZESTORETIC, HYDROCHLOROTHIAZIDE
ZESTRIL, LISINOPRIL
ZETIA, EZETIMIBE
ZIAC, BISOPROLOL FUMARATE
ZIAGEN, ABACAVIR SULFATE
ZIANA, CLINDAMYCIN PHOSPHATE
ZIDOVUDINE, ZIDOVUDINE
ZINACEF IN PLASTIC CONTAINER, CEFUROXIME SODIUM
ZINACEF, CEFUROXIME SODIUM
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
ZINECARD, DEXRAZOXANE HYDROCHLORIDE
ZIPSOR, DICLOFENAC POTASSIUM
ZIRGAN, GANCICLOVIR
ZITHROMAX, AZITHROMYCIN
ZMAX, AZITHROMYCIN
ZOCOR, SIMVASTATIN
ZOFTRAN ODT, ONDANSETRON
ZOFTRAN PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ZOFTRAN, ONDANSETRON HYDROCHLORIDE
ZOLADEX, GOSERELIN ACETATE
ZOLINZA, VORINOSTAT
ZOLOFT, SERTRALINE HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOLPIMIST, ZOLPIDEM TARTRATE
ZOMETA, ZOLEDRONIC ACID
ZOMIG, ZOLMITRIPTAN
ZOMIG-ZMT, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZORBTIVE, SOMATROPIN RECOMBINANT
ZORTRESS, EVEROLIMUS
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOSYN, PIPERACILLIN SODIUM
ZOVIA 1/35E-28, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

A - 59

**** Z ****

ZOVIA 1/50E-28, ETHINYL ESTRADIOL
ZOVIRAX, ACYCLOVIR
ZUPLENZ, ONDANSETRON
ZUTRIPRO, CHLORPHENIRAMINE MALEATE
ZYBAN, BUPROPION HYDROCHLORIDE
ZYCLARA, IMIQUIMOD
ZYDONE, ACETAMINOPHEN
ZYFLO CR, ZILEUTON
ZYFLO, ZILEUTON
ZYLET, LOTEPEREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAR, GATIFLOXACIN
ZYMAXID, GATIFLOXACIN
ZYPREXA RELPREVV, OLANZAPINE PAMOATE
ZYPREXA ZYDIS, OLANZAPINE
ZYPREXA, OLANZAPINE
ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
ZYRTEC, CETIRIZINE HYDROCHLORIDE
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)
ZYTIGA, ABIRATERONE ACETATE
ZYVOX, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** 3 **

3M

- * 3M CO
PERIDEX, CHLORHEXIDINE GLUCONATE
- * 3M HEALTH CARE INC
AVAGARD, ALCOHOL (OTC)
DURAPREP, IODINE POVACRYLEX (OTC)
- * 3M PHARMACEUTICALS INC
PROVENTIL-HFA, ALBUTEROL SULFATE

3M INFECTION

- * 3M INFECTION PREVENTION DIV
CHLORASCRUB MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWAB, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

AAIPHARMA LLC

- * AAIPHARMA LLC
AZASAN, AZATHIOPRINE

ABBOTT

- * ABBOTT LABORATORIES
ADVICOR, LOVASTATIN
AKINETON, BIPERIDEN HYDROCHLORIDE
AZMACORT, TRIAMCINOLONE ACETONIDE
BIAXIN XL, CLARITHROMYCIN
BIAXIN, CLARITHROMYCIN
CYCLOSPORINE, CYCLOSPORINE
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
GENGRAF, CYCLOSPORINE
KALETRA, LOPINAVIR
NIASPAN, NIACIN
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMBEX, CISATRACURIUM BESYLATE
NORVIR, RITONAVIR
SIMCOR, NIACIN
SYNTHROID, LEVOTHYROXINE SODIUM
TEVETEN HCT, EPROSARTAN MESYLATE
TEVETEN, EPROSARTAN MESYLATE
ULTANE, SEVOFLURANE
ZEMPLAR, PARICALCITOL
- * ABBOTT LABORATORIES HOSP PRODUCTS DIV
CALCIJEX, CALCITRIOL
- * ABBOTT LABORATORIES PHARMACEUTICAL PRODUCTS DIV
BIAXIN, CLARITHROMYCIN
DEPACON, VALPROATE SODIUM
DEPAKENE, VALPROIC ACID
DEPAKOTE, DIVALPROEX SODIUM
ENDURON, METHYCLOTHIAZIDE
HYTRIN, TERAZOSIN HYDROCHLORIDE
K-TAB, POTASSIUM CHLORIDE
MAVIK, TRANDOLAPRIL
NORVIR, RITONAVIR
ORETIC, HYDROCHLOROTHIAZIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
TARKA, TRANDOLAPRIL
TRIDIONE, TRIMETHADIONE
VICODIN ES, ACETAMINOPHEN
VICODIN HP, ACETAMINOPHEN
VICODIN, ACETAMINOPHEN
VICOPROFEN, HYDROCODONE BITARTRATE
ZEMPLAR, PARICALCITOL

ABBOTT ENDOCRINE

- * ABBOTT ENDOCRINE INC
LUPRON DEPOT, LEUPROLIDE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ABBOTT ENDOCRINE INC
LUPRON DEPOT-PED, LEUPROLIDE ACETATE

ABBOTT LABS

- * ABBOTT ENDOCRINE INC SUB ABBOTT LABORATORIES
LUPRON DEPOT, LEUPROLIDE ACETATE
- * ABBOTT LABORATORIES
ANDROGEL, TESTOSTERONE
NORVIR, RITONAVIR
PROMETRIUM, PROGESTERONE
TRILIPPIX, CHOLINE FENOFIBRATE

ABBOTT LABS PHARM

- * ABBOTT LABORATORIES PHARMACEUTICAL PRODUCTS DIV
TRICOR, FENOFIBRATE

ABBOTT PRODS

- * ABBOTT PRODUCTS INC
ACEON, PERINDOPRIL ERBUMINE
CREON, LIPASE
MARINOL, DRONABINOL

ABRAXIS BIOSCIENCE

- * ABRAXIS BIOSCIENCE LLC
ABRAXANE, PACLITAXEL

ABRAXIS PHARM

- * ABRAXIS PHARMACEUTICAL PRODUCTS
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
MAGNESIUM SULFATE, MAGNESIUM SULFATE

ACADEMIC PHARMS

- * ACADEMIC PHARMACEUTICALS INC
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

ACCORD HLTH

- * ACCORD HEALTH CARE INC
METHYLDOPA, METHYLDOPA

ACCORD HLTHCARE

- * ACCORD HEALTHCARE INC
DOCETAXEL, DOCETAXEL
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GLIMEPIRIDE, GLIMEPIRIDE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LETROZOLE, LETROZOLE
LEVETIRACETAM, LEVETIRACETAM
MITOMYCIN, MITOMYCIN
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
SIMVASTATIN, SIMVASTATIN
TACROLIMUS, TACROLIMUS
TOPIRAMATE, TOPIRAMATE

ACCORD HLTHCARE INC

- * ACCORD HEALTHCARE INC USA
ANASTROZOLE, ANASTROZOLE
BICALUTAMIDE, BICALUTAMIDE
CLONAZEPAM, CLONAZEPAM
ETOPOSIDE, ETOPOSIDE
FINASTERIDE, FINASTERIDE
GLIPIZIDE, GLIPIZIDE
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
PACLITAXEL, PACLITAXEL

ACIC FINE CHEMS

- * ACIC FINE CHEMICALS INC
AMPICILLIN SODIUM, AMPICILLIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ACIC FINE CHEMICALS INC
NAFCILLIN SODIUM, NAFCILLIN SODIUM

ACORDA

* ACORDA THERAPEUTICS INC
AMPYRA, DALFAMPRIDINE
ZANAFLEX, TIZANIDINE HYDROCHLORIDE

ACS DOBFAR

* ACS DOBFAR SPA
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFEPIME 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
FLUCANAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN

ACTAVIS

* ACTAVIS SOUTH ATLANTIC LLC
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
* ACTAVIS SOUTHLANTLATIC LLC
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
FENTANYL-100, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL
NIFEDIPIINE, NIFEDIPIINE
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE

ACTAVIS ELIZABETH

* ACTAVIS ELIZABETH LLC
ALPRAZOLAM, ALPRAZOLAM
CARBIDOPA AND LEVODOPA, CARBIDOPA
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
GABAPENTIN, GABAPENTIN
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
INDAPAMIDE, INDAPAMIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
KADIAN, MORPHINE SULFATE
LEVETIRACETAM, LEVETIRACETAM
LORAZEPAM, LORAZEPAM
LOVASTATIN, LOVASTATIN
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NIFEDIPIINE, NIFEDIPIINE
OXAZEPAM, OXAZEPAM
OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
PROPYLTIOURACIL, PROPYLTIOURACIL
SPIRONOLACTONE, SPIRONOLACTONE
TEMAZEPM, TEMAZEPM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ACTAVIS INC**

- * ACTAVIS INC
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM

ACTAVIS MID ATLANTIC

- * ACTAVIS MID ATLANTIC LLC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETASOL HC, ACETIC ACID, GLACIAL
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE, ALBUTEROL SULFATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CICLOPIROX, CICLOPIROX
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
CONSTULOSE, LACTULOSE
ENULOSE, LACTULOSE
FLUOCINONIDE, FLUOCINONIDE
GRISEOFULVIN, GRISEOFULVIN, MICROCRYSTALLINE
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
HYDROCORTISONE, HYDROCORTISONE
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
LEVETIRACETAM, LEVETIRACETAM
MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
NYSTATIN, NYSTATIN
PERMETHRIN, PERMETHRIN
PERMETHRIN, PERMETHRIN (OTC)
PROMETH VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE
PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
PROMETH W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
VALNAC, BETAMETHASONE VALERATE

ACTAVIS PHARMA

- * ACTAVIS PHARMA MANUFACTURING PRIVATE LTD
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE

ACTAVIS S ATLANTIC

- * ACTAVIS SOUTH ATLANTIC LLC
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZOLPIDEM TATRATE, ZOLPIDEM TARTRATE

ACTAVIS TOTOWA

- * ACTAVIS TOTOWA LLC
BICALUTAMIDE, BICALUTAMIDE
DESIPIRAMINE HYDROCHLORIDE, DESIPIRAMINE HYDROCHLORIDE
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
FINASTERIDE, FINASTERIDE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LETROZOLE, LETROZOLE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ACTAVIS TOTOWA LLC
 - PACLITAXEL, PACLITAXEL
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - VINORELBINE TARTRATE, VINORELBINE TARTRATE

ACTELION

- * ACTELION LTD
 - TRACLEER, BOSENTAN
 - VELETRI, EPOPROSTENOL SODIUM

ACTELION PHARMS LTD

- * ACTELION PHARMACEUTICALS LTD
 - VENTAVIS, ILOPROST
 - ZAVESCA, MIGLUSTAT

ACTIENT PHARMS

- * ACTIENT PHARMACEUTICALS LLC
 - STRIANT, TESTOSTERONE

ACTON PHARMS

- * ACTON PHARMACEUTICALS INC
 - AEROSPAN HFA, FLUNISOLIDE

ADOLOR

- * ADOLOR CORP
 - ENTEREG, ALVIMOPAN

ADVENT PHARMS

- * ADVENT PHARMACEUTICALS INC
 - CARISOPRODOL, CARISOPRODOL

AEGIS PHARMS

- * AEGIS PHARMACEUTICALS INC
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE

AGOURON

- * AGOURON PHARMACEUTICALS INC
 - VIRACEPT, NELFINAVIR MESYLATE

AJANTA PHARMA

- * AJANTA PHARMA LTD
 - LEVETIRACETAM, LEVETIRACETAM

AJANTA PHARMA LTD

- * AJANTA PHARMA LTD
 - RISPERIDONE, RISPERIDONE

AKORN

- * AKORN INC
 - ADENOSINE, ADENOSINE
 - AKBETA, LEVOBUNOLOL HYDROCHLORIDE
 - AK-FLUOR 10%, FLUORESCIN SODIUM
 - AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
 - AKPRO, DIPIVEFRIN HYDROCHLORIDE
 - AKTEN, LIDOCAINE HYDROCHLORIDE
 - AKTOB, TOBRAMYCIN
 - ALFENTA, ALFENTANIL HYDROCHLORIDE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 - BAL, DIMERCAPROL
 - BALANCED SALT, CALCIUM CHLORIDE
 - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 - BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 - CALCITRIOL, CALCITRIOL
 - CAPASTAT SULFATE, CAPREOMYCIN SULFATE
 - CARBOPLATIN, CARBOPLATIN
 - CROMOLYN SODIUM, CROMOLYN SODIUM
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - ENDOSOL EXTRA, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AKORN INC
 - ERYTHROMYCIN, ERYTHROMYCIN
 - GENTAK, GENTAMICIN SULFATE
 - GENTAMICIN SULFATE, GENTAMICIN SULFATE
 - 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)
 - LEVOFLOXACIN, LEVOFLOXACIN
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
 - OFLOXACIN, OFLOXACIN
 - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 - PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
 - SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
 - SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
 - TERBUTALINE SULFATE, TERBUTALINE SULFATE
 - TIMOLOL MALEATE, TIMOLOL MALEATE
 - TROPICACYL, TROPICAMIDE

AKORN INC

- * AKORN INC
 - APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - HYDASE, HYALURONIDASE
 - INAPSINE, DROPERIDOL

AKORN STRIDES

- * AKORN STRIDES LLC
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

AKRIMAX PHARMS

- * AKRIMAX PHARMACEUTICALS LLC
 - INDERAL LA, PROPRANOLOL HYDROCHLORIDE
 - INDERAL, PROPRANOLOL HYDROCHLORIDE
 - INDERIDE-40/25, HYDROCHLOROTHIAZIDE

ALARA PHARM

- * ALARA PHARMACEUTICAL CORPORATION
 - LEVO-T, LEVOTHYROXINE SODIUM

ALCON

- * ALCON INC
 - TRIESENCE, TRIAMCINOLONE ACETONIDE
- * ALCON LABORATORIES INC
 - ALCAINE, PROPARACAINA HYDROCHLORIDE
 - ALOMIDE, LODOXAMIDE TROMETHAMINE
 - BETOPTIC S, BETAXOLOL HYDROCHLORIDE
 - BETOPTIC, BETAXOLOL HYDROCHLORIDE
 - BSS PLUS, CALCIUM CHLORIDE
 - BSS, CALCIUM CHLORIDE
 - CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 - CETAMIDE, SULFACETAMIDE SODIUM
 - CROMOLYN SODIUM, CROMOLYN SODIUM
 - CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
 - CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
 - DENDRID, IDOXURIDINE
 - EMADINE, EMEDASTINE DIFUMARATE
 - FLAREX, FLUOROMETHOLONE ACETATE
 - IOPIDINE, APRACLONIDINE HYDROCHLORIDE
 - MAXIDEX, DEXAMETHASONE
 - MAXITROL, DEXAMETHASONE
 - MIOSTAT, CARBACHOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* ALCON LABORATORIES INC
 MYDRIACYL, TROPICAMIDE
 NAPHCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 NATACYN, NATAMYCIN
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 OMNIPRED, PREDNISOLONE ACETATE
 PATANOL, OLOPATADINE HYDROCHLORIDE
 PILOPINE HS, PILOCARPINE HYDROCHLORIDE
 TOBRADEX, DEXAMETHASONE
 TOBREX, TOBRAMYCIN
 TRIFLURIDINE, TRIFLURIDINE
 VEXOL, RIMEXOLONE

ALCON PHARMA

* ALCON RESEARCH LTD
 ZADITOR, KETOTIFEN FUMARATE (OTC)

ALCON PHARMS LTD

* ALCON PHARMACEUTICALS LTD
 AZOPT, BRINZOLAMIDE
 BETADINE, POVIDONE-IODINE
 BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
 CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
 CIPRODEX, CIPROFLOXACIN
 DICLOFENAC SODIUM, DICLOFENAC SODIUM
 DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
 DUREZOL, DIFLUPREDNATE
 FLUORESCITE, FLUORESCEIN SODIUM
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
 MOXEZA, MOXIFLOXACIN HYDROCHLORIDE
 NAVSTEL, CALCIUM CHLORIDE
 NEVANAC, NEPAFENAC
 OFLOXACIN, OFLOXACIN
 PATADAY, OLOPATADINE HYDROCHLORIDE
 PATANASE, OLOPATADINE HYDROCHLORIDE
 SULFACTAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 TOBRADEX ST, DEXAMETHASONE
 TRAVATAN Z, TRAVOPROST
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE

ALCON RES

* ALCON RESEARCH LTD
 DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE
 LATANOPROST, LATANOPROST

ALCON UNIVERSAL

* ALCON UNIVERSAL LTD
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
 NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM

ALKOPHARMA USA

* ALKOPHARMA USA INC
 AMPHOTEC, AMPHOTERICIN B

ALEMBIC LTD

* ALEMBIC LTD
 LITHIUM CARBONATE, LITHIUM CARBONATE
 METRONIDAZOLE, METRONIDAZOLE
 PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
 ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 THEOPHYLLINE, THEOPHYLLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ALEMBIC PHARMS LTD**

- * ALEMBIC PHARMACEUTICALS LTD
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - FAMOTIDINE, FAMOTIDINE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LAMOTRIGINE, LAMOTRIGINE
 - LEFLUNOMIDE, LEFLUNOMIDE
 - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 - MEPROBAMATE, MEPROBAMATE
 - METRONIDAZOLE, METRONIDAZOLE
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

ALKEM

- * ALKEM LABORATORIES LTD
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - CEPHALEXIN, CEPHALEXIN
 - GABAPENTIN, GABAPENTIN
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

ALKEM LABS LTD

- * ALKEM LABORATORIES LTD
 - CEFUXIME AXETIL, CEFUXIME AXETIL
 - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

ALKERMES

- * ALKERMES INC
 - VIVITROL, NALTREXONE

ALKERMES GAINESVILLE

- * ALKERMES GAINESVILLE LLC
 - VERELAN, VERAPAMIL HYDROCHLORIDE

ALLEGIANCE HLTHCARE

- * ALLEGIANCE HEALTHCARE CORP
 - POVIDONE IODINE, POVIDONE-IODINE (OTC)

ALLERGAN

- * ALLERGAN
 - ACULAR LS, KETOROLAC TROMETHAMINE
 - ALPHAGAN P, BRIMONIDINE TARTRATE
 - BLEPH-10, SULFACETAMIDE SODIUM
 - GENOPTIC, GENTAMICIN SULFATE
 - ZYMAXID, GATIFLOXACIN
- * ALLERGAN INC
 - ACULAR PRESERVATIVE FREE, KETOROLAC TROMETHAMINE
 - ACULAR, KETOROLAC TROMETHAMINE
 - ACUVAIL, KETOROLAC TROMETHAMINE
 - ACZONE, DAPSONE
 - ALBALON, NAPHAZOLINE HYDROCHLORIDE
 - ALOCRIL, NEDOCROMIL SODIUM
 - ALPHAGAN P, BRIMONIDINE TARTRATE
 - AVAGE, TAZAROTENE
 - AZELEX, AZELAIC ACID
 - COMBIGAN, BRIMONIDINE TARTRATE
 - ELESTAT, EPINASTINE HYDROCHLORIDE
 - ELIMITE, PERMETHRIN
 - LASTACAFT, ALCAFTADINE
 - LATISSE, BIMATOPROST
 - LUMIGAN, BIMATOPROST
 - OCUFLOX, OFLOXACIN
 - OPTICROM, CROMOLYN SODIUM
 - OZURDEX, DEXAMETHASONE
 - POLYTRIM, POLYMYXIN B SULFATE
 - RESTASIS, CYCLOSPORINE
 - SANCTURA XR, TROSPiUM CHLORIDE
 - SANCTURA, TROSPiUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ALLERGAN INC
 - TAZORAC, TAZAROTENE
 - TRIVARIS, TRIAMCINOLONE ACETONIDE
 - ZYMAR, GATIFLOXACIN
- * ALLERGAN PHARMACEUTICAL
 - BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
 - BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
 - BLEPHAMIDE, PREDNISOLONE ACETATE
 - FML FORTE, FLUOROMETHOLONE
 - FML, FLUOROMETHOLONE
 - HERPLEX, IDOXURIDINE
 - OCUFEN, FLURBIPROFEN SODIUM
 - OPHTHETIC, PROPARACAINA HYDROCHLORIDE
 - POLY-PRED, NEOMYCIN SULFATE
 - PRED FORTE, PREDNISOLONE ACETATE
 - PRED MILD, PREDNISOLONE ACETATE
 - PRED-G, GENTAMICIN SULFATE
 - PROPINE, DIPIVEFRIN HYDROCHLORIDE

ALLERGAN HERBERT

- * ALLERGAN HERBERT SKIN CARE DIV ALLERGAN INC
 - FLUOROPLEX, FLUOROURACIL

ALLERQUEST

- * ALLERQUEST LLC
 - PRE-PEN, BENZYL PENICILLOYL POLYLYSINE

ALLOS

- * ALLOS THERAPEUTICS INC
 - FOLOTYN, PRALATREXATE

ALPHAPHARM

- * ALPHAPHARM PARTY LTD
 - ALPRAZOLAM, ALPRAZOLAM
 - CILOSTAZOL, CILOSTAZOL
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLONAZEPAM, CLONAZEPAM
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - INDAPAMIDE, INDAPAMIDE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - MIRTAZAPINE, MIRTAZAPINE
 - NAPROXEN, NAPROXEN
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRIAZOLAM, TRIAZOLAM
 - ZONISAMIDE, ZONISAMIDE
- * ALPHAPHARM PTY LTD
 - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

ALPHARMA

- * ALPHARMA USPD INC
 - PREDNISOLONE, PREDNISOLONE
 - VALPROIC ACID, VALPROIC ACID

ALPHARMA KING

- * ALPHARMA PHARMACEUTICALS LLC KING PHARMACEUTICALS
 - EMBEDA, MORPHINE SULFATE

ALRA

- * ALRA LABORATORIES INC
 - CHOLAC, LACTULOSE
 - CONSTILAC, LACTULOSE
 - GEN-XENE, CLORAZEPATE DIPOTASSIUM
 - IBU-TAB 200, IBUPROFEN (OTC)
 - IBU-TAB, IBUPROFEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******ALTAIRE PHARMS INC**

- * ALTAIRE PHARMACEUTICALS INC
NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)

ALTANA

- * ALTANA INC
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
AMCINONIDE, AMCINONIDE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE (EMOLlient), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CUTIVATE, FLUTICASONE PROPIONATE
DESONIDE, DESONIDE
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
HYDROCORTISONE, HYDROCORTISONE
KETOCONAZOLE, KETOCONAZOLE
METRONIDAZOLE, METRONIDAZOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NYSTATIN, NYSTATIN
OXISTAT, OXICONAZOLE NITRATE
PREDNICARBATE, PREDNICARBATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TERCONAZOLE, TERCONAZOLE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

ALTANA PHARMA

- * ALTANA PHARMA AG
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE

ALTERNA TCHP LLC

- * ALTERNA TCHP LLC
CHILDREN'S ELIXSURE, IBUPROFEN (OTC)

ALVOGEN

- * ALVOGEN INC
MACROBID, NITROFURANTOIN
MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

ALVOGEN INC

- * ALVOGEN INC
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

AM ANTIBIOTICS

- * AMERICAN ANTIBIOTICS LLC
AMOXICILLIN, AMOXICILLIN

AMAG PHARMS INC

- * AMAG PHARMACEUTICALS INC
FERAHeme, FERUMOXYTOL
FERIDEX I.V., FERUMOXIDES
GASTROMARK, FERUMOXSIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AMGEN**

- * AMGEN INC
SENSIPAR, CINACALCET HYDROCHLORIDE

AMNEAL PHARM

- * AMNEAL PHARMACEUTICAL
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLUCONAZOLE, FLUCONAZOLE
FOLIC ACID, FOLIC ACID
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
PRIMIDONE, PRIMIDONE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

AMNEAL PHARMS

- * AMNEAL PHARMACEUTICALS
ALBUTEROL SULFATE, ALBUTEROL SULFATE
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
FELBAMATE, FELBAMATE
IBUPROFEN, IBUPROFEN (OTC)
INDOMETHACIN, INDOMETHACIN
LEVETIRACETAM, LEVETIRACETAM
LORAZEPAM, LORAZEPAM
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
NITROFURANTOIN, NITROFURANTOIN
NIZATIDINE, NIZATIDINE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
SPIRONOLACTONE, SPIRONOLACTONE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- * AMNEAL PHARMACEUTICALS LLC
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AMNEAL PHARMACEUTICALS NY LLC
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
REPREXAIN, HYDROCODONE BITARTRATE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

AMPHASTAR PHARM

- * AMPHASTAR PHARMACEUTICAL INC
AMPHADASE, HYALURONIDASE
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM

AMPHASTAR PHARMS INC

- * AMPHASTAR PHARMACEUTICALS INC
CORTROSYN, COSYNTROPIN

AMPOLGEN

- * AMPOLGEN PHARMACEUTICALS LLC
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE

AMYLIN

- * AMYLIN PHARMACEUTICALS INC
BYETTA, EXENATIDE SYNTHETIC
SYMLIN, PRAMLINTIDE ACETATE

ANBEX

- * ANBEX INC
IOSAT, POTASSIUM IODIDE (OTC)

ANCHEN PHARMS

- * ANCHEN PHARMACEUTICALS INC
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * ANCHEN PHARMACEUTICALS TAIWAN INC
DIVALPROEX SODIUM, DIVALPROEX SODIUM
- * ANCHEN PHARMACEUTICALS, INC
ALPRAZOLAM, ALPRAZOLAM
CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN

ANDRX LABS LLC

- * ANDRX LABS LLC
ALTOPREV, LOVASTATIN
FORTAMET, METFORMIN HYDROCHLORIDE

ANESTA AG

- * ANESTA AG
AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE

ANGELINI LLC

- * ANGELINI LABOPHARM LLC
OLEPTRO, TRAZODONE HYDROCHLORIDE

ANI PHARMS

- * ANI PHARMACEUTICALS INC
CORTENEMA, HYDROCORTISONE
LACTULOSE, LACTULOSE
LUVOX, FLUVOXAMINE MALEATE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE

ANTARES PHARMA INC

- * ANTARES PHARMA INC
ANTUROL, OXYBUTYNIN

ANTIBIOTICOS BRASIL

- * ANTIBIOTICOS DO BRASIL LTDA
CEFOXITIN, CEFOXITIN

APOPHARMA INC

- * APOPHARMA INC
FERRIPROX, DEFERIPRONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******APOTEX**

- * APOTEX CORP
 - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
- * APOTEX INC
 - ALENDRONATE SODIUM, ALENDRONATE SODIUM
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - BUDESONIDE, BUDESONIDE
 - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 - CAPTOPRIL, CAPTOPRIL
 - CARBIDOPA AND LEVODOPA, CARBIDOPA
 - CEFUROXIME AXETIL, CEFUROXIME AXETIL
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - CIMETIDINE, CIMETIDINE
 - CIMETIDINE, CIMETIDINE (OTC)
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - CLONAZEPAM, CLONAZEPAM
 - CYCLOSPORINE, CYCLOSPORINE
 - DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 - DILT-CD, DILTIAZEM HYDROCHLORIDE
 - 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
 - KETOCONAZOLE, KETOCONAZOLE
 - LAMIVUDINE, LAMIVUDINE
 - LATANOPROST, LATANOPROST
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVOFLOXACIN, LEVOFLOXACIN
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 - NIZATIDINE, NIZATIDINE
 - OMEPRAZOLE, OMEPRAZOLE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 - PENTOXIFYLLINE, PENTOXIFYLLINE
 - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 - RAMIPRIL, RAMIPRIL
 - RANITIDINE HYDROCHLORIDE, RANITIDINE 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******APOTEX CORP**

* APOTEX CORP

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AZITHROMYCIN, AZITHROMYCIN
CICLOPIROX, CICLOPIROX
CLARITHROMYCIN, CLARITHROMYCIN
GATIFLOXACIN, GATIFLOXACIN
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NARatriptan, NARatriptan

APOTEX INC

* APOTEX INC

ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALPRAZOLAM, ALPRAZOLAM
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
CALCITONIN-SALMON, CALCITONIN SALMON
CARBAMAZEPINE, CARBAMAZEPINE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CEFPROZIL, CEFPROZIL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CEVIMELINE, CEVIMELINE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
LOVASTATIN, LOVASTATIN
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
NABUMETONE, NABUMETONE
OFLOXACIN, OFLOXACIN
OLANZAPINE, OLANZAPINE
OXCARBAZEPINE, OXCARBAZEPINE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

* APOTEX INC ETOBICOKE SITE

ACYCLOVIR, ACYCLOVIR
ALLOPURINOL, ALLOPURINOL
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
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
KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
LEFLUNOMIDE, LEFLUNOMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * APOTEX INC ETOBICOKE SITE
 - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LISINOPRIL, LISINOPRIL
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - LORATADINE, LORATADINE (OTC)
 - MELOXICAM, MELOXICAM
 - MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 - MIRTAZAPINE, MIRTAZAPINE
 - OXaprozin, OXaprozin
 - SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 - TOPIRAMATE, TOPIRAMATE
 - TORSEMIDE, TORSEMIDE
 - ZONISAMIDE, ZONISAMIDE
- * APOTEX INC RICHMOND HILL
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
 - DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 - FLUNISOLIDE, FLUNISOLIDE
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - LORATADINE, LORATADINE (OTC)
 - MEGESTROL ACETATE, MEGESTROL ACETATE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - PREDNISOLONE, PREDNISOLONE
 - RISPERIDONE, RISPERIDONE
- * APOTEX INC.
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

APOTHECON

- * APOTHECON INC DIV BRISTOL MYERS SQUIBB
 - CAPOZIDE 25/15, CAPTOPRIL
 - CAPOZIDE 25/25, CAPTOPRIL
 - CAPOZIDE 50/15, CAPTOPRIL
 - CAPOZIDE 50/25, CAPTOPRIL
 - KENALOG, TRIAMCINOLONE ACETONIDE
 - KENALOG-10, TRIAMCINOLONE ACETONIDE
 - KENALOG-40, TRIAMCINOLONE ACETONIDE
 - KLOTRIX, POTASSIUM CHLORIDE
 - OPHTHAINE, PROPARACAINA HYDROCHLORIDE
 - STADOL PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 - STADOL, BUTORPHANOL TARTRATE
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

APP PHARMS

- * APP PHARMACEUTICALS
 - GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
 - PROTAMINE SULFATE, PROTAMINE SULFATE
- * APP PHARMACEUTICALS LLC
 - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 - ADENOSINE, ADENOSINE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - ASTRAMORPH PF, MORPHINE SULFATE
 - AZITHROMYCIN, AZITHROMYCIN
 - AZTREONAM, AZTREONAM
 - BACITRACIN, BACITRACIN
 - BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 - BLEOMYCIN SULFATE, BLEOMYCIN SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * APP PHARMACEUTICALS LLC
- CAFFEINE CITRATE, CAFFEINE CITRATE
- CALCITRIOL, CALCITRIOL
- CARBOPLATIN, CARBOPLATIN
- CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
- CEFOTAXIME, CEFOTAXIME SODIUM
- CEFOXITIN, CEFOXITIN SODIUM
- CEFTRIAXONE, CEFTRIAXONE SODIUM
- CEFUROXIME SODIUM, CEFUROXIME SODIUM
- CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
- CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
- CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
- CISPLATIN, CISPLATIN
- CLADRIBINE, CLADRIBINE
- CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
- COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
- CYTARABINE, CYTARABINE
- DACARBAZINE, DACARBAZINE
- DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
- DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
- DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
- DIMENHYDRINATE, DIMENHYDRINATE
- DIPHENHYDRAMINE HYDROCHLORIDE, 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 HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
- FOLIC ACID, FOLIC ACID
- FUROSEMIDE, FUROSEMIDE
- GANCICLOVIR, GANCICLOVIR SODIUM
- GENTAMICIN SULFATE, GENTAMICIN SULFATE
- GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
- HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
- HALOPERIDOL, HALOPERIDOL LACTATE
- HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
- HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
- HEPARIN SODIUM, HEPARIN SODIUM
- HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
- HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
- IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
- IFOSFAMIDE, IFOSFAMIDE
- INDOMETHACIN, INDOMETHACIN
- IOPAMIDOL-250, IOPAMIDOL
- IOPAMIDOL-300, IOPAMIDOL
- IOPAMIDOL-370, IOPAMIDOL
- KANAMYCIN SULFATE, KANAMYCIN SULFATE
- KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
- LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
- LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
- LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
- MANNITOL 25%, MANNITOL
- MESNA, MESNA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * APP PHARMACEUTICALS LLC
 - METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
 - METHOTREXATE SODIUM, METHOTREXATE SODIUM
 - METHYLSPREDNISOLONE SODIUM SUCCINATE, METHYLSPREDNISOLONE SODIUM SUCCINATE
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - MILRINONE LACTATE, MILRINONE LACTATE
 - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 - NAROPIN, ROPIVACAINE HYDROCHLORIDE MONOHYDRATE
 - NEBUPENT, PENTAMIDINE ISETHIONATE
 - 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
 - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 - PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
 - PENTAM, PENTAMIDINE ISETHIONATE
 - 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
 - PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 - SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 - STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TERBUTALINE SULFATE, TERBUTALINE SULFATE
 - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 - TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 - 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 VISCOS, LIDOCAINE HYDROCHLORIDE
 - XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 - XYLOCAINE, LIDOCAINE HYDROCHLORIDE
- * APP PHARMACEUTICALS, LLC--A CO. OF THE FRESENIUS KABI GROUP
 - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 - CEFOTETAN, CEFOTETAN DISODIUM
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

APTALIS PHARMA US

- * APTALIS PHARMA US INC
 - BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
 - BENTYL, DICYCLOMINE HYDROCHLORIDE
 - CARAFATE, SUCRALFATE
 - PYLERA, BISMUTH SUBCITRATE POTASSIUM
 - URSO 250, URSODIOL
 - URSO FORTE, URSODIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

* APTALIS PHARMA US INC
ZENPEP, LIPASE

AQUA PHARMS

* AQUA PHARMACEUTICALS LLC
SOLAGE, MEQUINOL
XOLEGEL, KETOCONAZOLE

AR HOLDING CO INC

* AR HOLDING CO INC
COLCRYS, COLCHICINE
FIBRICOR, FENOFLIBRIC ACID
QUALAQUIN, QUININE SULFATE

ARBOR PHARMS INC

* ARBOR PHARMACEUTICALS INC
BIDIL, HYDRALAZINE HYDROCHLORIDE
E.E.S. 200, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERY-TAB, ERYTHROMYCIN
ERYTHROGIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN, ERYTHROMYCIN
PCE, ERYTHROMYCIN
PEDIAMYCIN 400, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAMYCIN, ERYTHROMYCIN ETHYLSUCCINATE

ARCHIMEDES

* ARCHIMEDES DEVELOPMENT LTD
LAZANDA, FENTANYL CITRATE

AREVA PHARMS

* AREVA PHARMACEUTICALS INC
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

ARMSTRONG PHARMS

* ARMSTRONG PHARMACEUTICALS INC
EPINEPHRINE, EPINEPHRINE (OTC)

ASCEND

* ASCEND THERAPEUTICS INC
ESTROGEL, ESTRADIOL

ASPEN GLOBAL

* ASPEN GLOBAL INC
MYLERAN, BUSULFAN

ASTELLAS

* ASTELLAS PHARMA US INC
ADENOCARD, ADENOSINE
ADENOSCAN, ADENOSINE
AMBISOME, AMPHOTERICIN B
LEXISCAN, REGADENOSON
MYCAMINE, MICAFUNGIN SODIUM
PROGRAF, TACROLIMUS
PROTOPIC, TACROLIMUS
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE
VESICARE, SOLIFENACIN SUCCINATE

ASTRAZENECA

* ASTRAZENECA LP
ENTOCORT EC, BUDESONIDE
NEXIUM IV, ESOMEPRAZOLE SODIUM
NEXIUM, ESOMEPRAZOLE MAGNESIUM
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
PRILOSEC, OMEPRAZOLE
PRILOSEC, OMEPRAZOLE MAGNESIUM
PULMICORT FLEXHALER, BUDESONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * ASTRAZENECA LP
 - PULMICORT RESPULES, BUDESONIDE
 - RHINOCORT, BUDESONIDE
 - SEROQUEL, QUETIAPINE FUMARATE
 - SYMBICORT, BUDESONIDE
 - TENORMIN, ATENOLOL
 - TOPROL-XL, METOPROLOL SUCCINATE
- * ASTRAZENECA PHARMACEUTICALS LP
 - ATACAND HCT, CANDESARTAN CILEXETIL
 - ATACAND, CANDESARTAN CILEXETIL
 - FASLODEX, FULVESTRANT
 - PLENDIL, FELODIPINE
 - PULMICORT, BUDESONIDE
 - SEROQUEL XR, QUETIAPINE FUMARATE
 - TENORETIC 100, ATENOLOL
 - TENORETIC 50, ATENOLOL
 - ZOMIG, ZOLMITRIPTAN
 - ZOMIG-ZMT, ZOLMITRIPTAN
- * ASTRAZENECA UK LTD
 - ACCOLATE, ZAFIRLUKAST
 - ARIMIDEX, ANASTROZOLE
 - CASODEX, BICALUTAMIDE
 - MERREM, MEROPENEM
 - ZESTORETIC, HYDROCHLOROTHIAZIDE
 - ZESTRIL, LISINOPRIL
 - ZOLADEX, GOSERELIN ACETATE

ASTRAZENECA LP

- * ASTRAZENECA LP
 - BRILINTA, TICAGRELOR
 - VIMOVO, ESOMEPRAZOLE MAGNESIUM

ATON

- * ATON PHARMA INC
 - CUPRIMINE, PENICILLAMINE
 - DEMSE, METYROSINE
 - EDECIN, ETHACRYNATE SODIUM
 - EDECIN, ETHACRYNIC ACID
 - LACRISERT, HYDROXYPROPYL CELLULOSE
 - LODOSYN, CARBIDOPA
 - SPRINE, TRIENTINE HYDROCHLORIDE
 - TIMOPIK IN OCUDOSE, TIMOLOL MALEATE
 - TIMOPIK, TIMOLOL MALEATE
 - TIMOPIK-XE, TIMOLOL MALEATE

AUROBINDO

- * AUROBINDO PHARMA LTD
 - AMOXICILLIN, AMOXICILLIN
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LISINOPRIL, LISINOPRIL
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - MIRTAZAPINE, MIRTAZAPINE
 - ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARM

- * AUROBINDO PHARMA USA INC
 - ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 - FLUCONAZOLE, FLUCONAZOLE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - GABAPENTIN, GABAPENTIN
 - LEVETIRACETAM, LEVETIRACETAM
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AUROBINDO PHARMA**

- * AUROBINDO PHARMA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
- * AUROBINDO PHARMA LTD
 ALENDRONATE SODIUM, ALENDRONATE SODIUM
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN, AMOXICILLIN
 AMPICILLIN SODIUM, AMPICILLIN SODIUM
 ATENOLOL, ATENOLOL
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
 CARISOPRODOL, CARISOPRODOL
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFDINIR, CEFDINIR
 CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFTAZIDIME, CEFTAZIDIME
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 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
 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ****

- * AUROBINDO PHARMA LTD
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
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
FAMCICLOVIR, FAMCICLOVIR
GABAPENTIN, GABAPENTIN
LAMIVUDINE, LAMIVUDINE
NAPROXEN SODIUM, NAPROXEN SODIUM
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
RAMIPRIL, RAMIPRIL
- * AUROBINDO PHARMA LTD INC
CEFPROZIL, CEFPROZIL
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
MIRTAZAPINE, MIRTAZAPINE
ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA USA

- * AUROBINDO PHARMA USA INC
ALPRAZOLAM, ALPRAZOLAM
NAPROXEN, NAPROXEN

AUROSAL PHARMS

- * AUROSAL PHARMACEUTICALS LLC
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE

AUSTARPHARMA LLC

- * AUSTARPHARMA LLC
ALENDRONATE SODIUM, ALENDRONATE SODIUM
METHOCARBAMOL, METHOCARBAMOL
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

AUXILIUM PHARMS

- * AUXILIUM PHARMACEUTICALS
TESTIM, TESTOSTERONE

AVACOR PRODS

- * AVACOR PRODUCTS LLC
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)

AVANIR PHARMS

- * AVANIR PHARMACEUTICALS INC
NUDEXTA, DEXTROMETHORPHAN HYDROBROMIDE

AVANTHI INC

- * AVANTHI INC
CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
INDOMETHACIN, INDOMETHACIN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

AVEMA PHARMA

- * AVEMA PHARMA SOLUTIONS
IBUPROFEN, IBUPROFEN (OTC)

AVENT

- * AVENT INC
PYTEST KIT, UREA, C-14
PYTEST, UREA, C-14

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** A ******AVEVA**

- * AVEVA DRUG DELIVERY SYSTEMS INC
CLONIDINE, CLONIDINE
NICOTINE, NICOTINE (OTC)

AXCAN

- * AXCAN PHARMA US INC
CANASA, MESALAMINE

AZTIQ PHARMA

- * AZTIQ PHARMA INC
TODAY, NONOXYNOL-9 (OTC)

AZUR PHARMA

- * AZUR PHARMA INTERNATIONAL LTD
AVC, SULFANILAMIDE
GASTROCROM, CROMOLYN SODIUM

AZUR PHARMA II

- * AZUR PHARMA INTERNATIONAL II LTD
ELESTRIN, ESTRADIOL
PRIALT, ZICONOTIDE ACETATE

AZUR PHARMA INTL

- * AZUR PHARMA INTERNATIONAL III LTD
FAZACLO ODT, CLOZAPINE

B BRAUN

- * B BRAUN MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ALCOHOL 10% AND DEXTROSE 5%, ALCOHOL
ALCOHOL 5% AND DEXTROSE 5%, ALCOHOL
BALANCED SALT, CALCIUM CHLORIDE
BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER, BRETYLIUM TOSYLATE
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 CHLORIDE
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 ACETATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
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
DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * B BRAUN MEDICAL INC
DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
FREAMINE HBC 6.9%, AMINO ACIDS
FREAMINE III 10%, AMINO ACIDS
FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5%, AMINO ACIDS
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
HEPARIN SODIUM 1,000 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
HEPATAMINE 8%, AMINO ACIDS
ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
ISOLYTE E IN PLASTIC CONTAINER, CALCIUM CHLORIDE
ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
ISOLYTE S 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
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
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER, MANNITOL
MANNITOL 10%, MANNITOL
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%, MANNITOL
MANNITOL 15%, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20%, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%, MANNITOL
MANNITOL 5%, MANNITOL
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
NEPHRAMINE 5.4%, AMINO ACIDS
NUTRILIPID 10%, SOYBEAN OIL
NUTRILIPID 20%, SOYBEAN OIL
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC
CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,
DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * B BRAUN MEDICAL INC
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * B BRAUN MEDICAL INC
 - POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PROCALAMINE, AMINO ACIDS
 - RESECTISOL IN PLASTIC CONTAINER, MANNITOL
 - 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
 - SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
 - 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

BANNER PHARMACAPS

- * BANNER PHARMACAPS INC
 - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 - BENZONATATE, BENZONATATE
 - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - ETHOSUXIMIDE, ETHOSUXIMIDE
 - IBUPROFEN AND DIPHENHYDRAMINE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - MIDOL LIQUID GELS, IBUPROFEN (OTC)
 - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 - NIMODIPINE, NIMODIPINE
 - STAVZOR, VALPROIC ACID
 - VALPROIC ACID, VALPROIC ACID
 - VITAMIN D, ERGOCALCIFEROL
 - ZONISAMIDE, ZONISAMIDE

BARR

- * BARR LABORATORIES INC
 - ALPRAZOLAM, ALPRAZOLAM
 - AMILORIDE HYDROCHLORIDE AND HYDROCHLORTIAZIDE, AMILORIDE HYDROCHLORIDE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 - ARANELLE, ETHINYLMESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BARR LABORATORIES INC
 - ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
 - BALZIVA-28, ETHINYL ESTRADIOL
 - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 - CAMILA, NORETHINDRONE
 - CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 - CHLORZOXAZONE, CHLORZOXAZONE
 - CLARAVIS, ISOTRETINOIN
 - CLONAZEPAM, CLONAZEPAM
 - CLONIDINE, CLONIDINE
 - DANAZOL, DANAZOL
 - DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
 - 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
 - ERRIN, NORETHINDRONE
 - ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL, ERYTHROMYCIN ETHYLSUCCINATE
 - ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
 - ESTRADIOL AND NORGESTIMATE, ESTRADIOL
 - ESTRADIOL, ESTRADIOL
 - ESTROPIPATE, ESTROPIPATE
 - ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
 - FENTANYL CITRATE, FENTANYL CITRATE
 - FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 - FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 - FLECAINIDE ACETATE, FLECAINIDE ACETATE
 - FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 - 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
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
 - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BARR LABORATORIES INC
 - 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
 - TRETINOIN, TRETINOIN
 - TREXALL, METHOTREXATE SODIUM
 - TRI LO SPRINTEC, ETHINYL ESTRADIOL
 - TRI-LEGEST 21, ETHINYL ESTRADIOL
 - TRI-LEGEST FE, ETHINYL ESTRADIOL
 - TRI-SPRINTEC, ETHINYL ESTRADIOL
 - WARFARIN SODIUM, WARFARIN SODIUM
 - ZONISAMIDE, ZONISAMIDE
- * BARR PHARMACEUTICALS
 - LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

BARR LABS INC

- * BARR LABORATORIES INC
 - NIMODIPINE, NIMODIPINE
 - OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN

BAUSCH AND LOMB

- * BAUSCH AND LOMB INC
 - ALAWAY, KETOTIFEN FUMARATE (OTC)
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - ALREX, LOTEPRENDNOL 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
 - LATANOPROST, LATANOPROST
 - LOTEMAX, LOTEPRENDNOL ETABONATE
 - MIOCHOL-E, ACETYLCHOLINE CHLORIDE
 - OFLOXACIN, OFLOXACIN
 - OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
 - RETISERT, FLUOCINOLONE ACETONIDE
 - TIMOLOL MALEATE, TIMOLOL MALEATE
 - VITRASERT, GANCICLOVIR
 - ZIRGAN, GANCICLOVIR
 - ZYLET, LOTEPRENDNOL ETABONATE
- * BAUSCH AND LOMB PHARMACEUTICALS INC
 - ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE, ACETIC ACID, GLACIAL
 - BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
 - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 - BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - CROLOM, CROMOLYN SODIUM
 - CROMOLYN SODIUM, CROMOLYN SODIUM
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAUSCH AND LOMB PHARMACEUTICALS INC
 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
 OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
 OTICAIR, HYDROCORTISONE
 PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROPARACAINA HYDROCHLORIDE, PROPARACAINA HYDROCHLORIDE
 SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 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

BAXTER HLTHCARE

* BAXTER HEALTHCARE CORP
 ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 ADENOSINE, ADENOSINE
 AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
 ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
 BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
 BRANCHAMIN 4% IN PLASTIC CONTAINER, AMINO ACIDS
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE
 CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
 CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 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, AMINO ACIDS
 CLINIMIX E 2.75/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/20 SULFITE-FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/15 SULFITE-FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/20 SULFITE-FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/35 SULFITE-FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CYTOXAN, CYCLOPHOSPHAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
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 75 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 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 (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
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER, DEXTROSE
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, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE
FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPATASOL 8%, AMINO ACIDS
IFEX, IFOSFAMIDE
ISOTONIC GENTAMICIN SULFATE IN PLASTIC CONTAINER, GENTAMICIN SULFATE
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
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
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
ONDANSETRON HYDROCHLORIDE AND SODIUM CHLORIDE IN PLASTIC CONTAINER, ONDANSETRON
HYDROCHLORIDE
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
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
PLASMA-LYTE 56 IN PLASTIC CONTAINER, MAGNESIUM ACETATE TETRAHYDRATE
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLASMA-LYTE R IN PLASTIC CONTAINER, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM
CHLORIDE
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
CALCIUM CHLORIDE
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
CALCIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
CALCIUM CHLORIDE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
CALCIUM CHLORIDE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP
 - POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% 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, CALCIUM CHLORIDE
 - POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 - POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 - PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 - RENAMIN W/O ELECTROLYTES, 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
 - SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
 - 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
 - THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 - TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
 - TIS-U-SOL, MAGNESIUM SULFATE
 - TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 10% W/O ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 3.5% W/ ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 5.5% W/ ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 5.5% W/O ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 8.5% W/ ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 8.5% W/O ELECTROLYTES, AMINO ACIDS
 - VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 - ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE, ACYCLOVIR SODIUM
 - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 - BUMETANIDE, BUMETANIDE
 - CEFOXITIN, CEFOXITIN SODIUM
 - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 - DIGOXIN, DIGOXIN
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DIPYRIDAMOLE, DIPYRIDAMOLE
 - DOPRAM, DOXAPRAM HYDROCHLORIDE
 - DURAMORPH PF, MORPHINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 - FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE
 - FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 - FLUMAZENIL, FLUMAZENIL
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - HEPARIN SODIUM, HEPARIN SODIUM
 - INFUMORPH, MORPHINE SULFATE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - MILRINONE LACTATE, MILRINONE LACTATE
 - MITOMYCIN, MITOMYCIN
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - ROBINUL, GLYCOPYRROLATE
 - SUFENTANIL CITRATE, SUFENTANIL CITRATE
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 - VINORELBINE TARTRATE, VINORELBINE TARTRATE
- * BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV
 - FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 - PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

BAXTER HLTHCARE CORP

- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 - MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
- * BAXTER HEALTHCARE CORP ANESTHESIA CRITICAL CARE
 - ATIVAN, LORAZEPAM
 - BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 - BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
 - BREVIBLOC, ESMOLOL HYDROCHLORIDE
 - ETHRANE, ENFLURANE
 - FORANE, ISOFLURANE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - OXYTOCIN, OXYTOCIN
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
 - REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
 - ROBAXIN, METHOCARBAMOL
 - SUPRANE, DESFLURANE

BAYER

- * BAYER HEALTHCARE LLC
 - ALEVE, NAPROXEN SODIUM (OTC)
 - ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)
 - FEMSTAT 3, BUTOCONAZOLE NITRATE (OTC)

BAYER HLTHCARE

- * BAYER HEALTHCARE CONSUMER CARE
 - RID MOUSSE, PIPERONYL BUTOXIDE (OTC)
- * BAYER HEALTHCARE PHARMACEUTICALS INC
 - ADALAT CC, NIFEDIPINE
 - ANGELIQ, DROSPIRENONE
 - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN HYDROCHLORIDE
 - AVELOX, MOXIFLOXACIN HYDROCHLORIDE
 - BETAPACE AF, SOTALOL HYDROCHLORIDE
 - BETAPACE, SOTALOL HYDROCHLORIDE
 - BEYAZ, DROSPIRENONE
 - CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 - CIPRO XR, CIPROFLOXACIN
 - CIPRO, CIPROFLOXACIN
 - CIPRO, CIPROFLOXACIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BAYER HEALTHCARE PHARMACEUTICALS INC
 CLIMARA PRO, ESTRADIOL
 CLIMARA, ESTRADIOL
 DTIC-DOME, DACARBAZINE
 EOVISt, GADOXETATE DISODIUM
 GADAVIST, GADOBUTROL
 LEVITRA, VARDENAFIL HYDROCHLORIDE
 MAGNEVIST, GADOPENTETATE DIMEGLUMINE
 MENOSTAR, ESTRADIOL
 MIRENA, LEVONORGESTREL
 MYCELEX, CLOTRIMAZOLE
 NATAZIA, ESTRADIOL VALERATE
 NEXAVAR, SORAFENIB TOSYLATE
 PRECOSE, ACARBOSE
 REFLUDAN, LEPIRUDIN RECOMBINANT
 SAFYRAL, DROSPIRENONE
 STAXYN, VARDENAFIL HYDROCHLORIDE
 TRASYLOL, APROTININ
 ULTRAVIST (PHARMACY BULK), IOPROMIDE
 ULTRAVIST 150, IOPROMIDE
 ULTRAVIST 240, IOPROMIDE
 ULTRAVIST 300 IN PLASTIC CONTAINER, IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 YASMIN, DROSPIRENONE
 YAZ, DROSPIRENONE

BAYER PHARMA AG

* BAYER PHARMA AG
 BILTRICIDE, PRAZIQUANTEL

BAYER PHARMS

* BAYER PHARMACEUTICALS CORP
 MYCELEX-7 COMBINATION PACK, CLOTRIMAZOLE (OTC)
 MYCELEX-7, CLOTRIMAZOLE (OTC)

BECTON DICKINSON

* BECTON DICKINSON AND CO
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
 E-Z SCRUB 201, POVIDONE-IODINE (OTC)
 E-Z SCRUB 241, POVIDONE-IODINE (OTC)

BEDFORD

* BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ADENOSINE, ADENOSINE
 ALPROSTADIL, ALPROSTADIL
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMRINONE LACTATE, INAMRINONE LACTATE
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 AZTREONAM, AZTREONAM
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BUMETANIDE, BUMETANIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CARBOPLATIN, CARBOPLATIN
 CEFTRIAXONE, CEFTRIAXONE SODIUM
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CISPLATIN, CISPLATIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC
CLADRIBINE, CLADRIBINE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CYCLOSPORINE, CYCLOSPORINE
CYTARABINE, CYTARABINE
DACARBAZINE, DACARBAZINE
DACTINOMYCIN, DACTINOMYCIN
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE-FREE), DICYCLOMINE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXYCYCLINE, DOXYCYCLINE HYCLATE
ENALAPRILAT, ENALAPRILAT
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
ETOMIDATE, ETOMIDATE
ETOPOSIDE, ETOPOSIDE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE
FLOXURIDINE, FLOXURIDINE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEVOCARNITINE, LEVOCARNITINE
LORAZEPAM, LORAZEPAM
MESNA, MESNA
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
METHOTREXATE SODIUM, METHOTREXATE SODIUM
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE, MILRINONE LACTATE
MITOMYCIN, MITOMYCIN
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NOREpinephrine BITARTRATE, NOREPINEPHRINE BITARTRATE
OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PACLITAXEL, PACLITAXEL
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RIFAMPIN, RIFAMPIN
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
THIOTEPA, THIOTEPA
VALPROATE SODIUM, VALPROATE SODIUM
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VINBLASTINE SULFATE, VINBLASTINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC
VINORELBINE TARTRATE, VINORELBINE TARTRATE

BEDFORD LABS

- * BEDFORD LABORATORIES
 - ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - CARBOPLATIN, CARBOPLATIN
 - DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 - FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
 - FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 - FLUMAZENIL, FLUMAZENIL
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 - IFOSFAMIDE, IFOSFAMIDE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - LORAZEPAM PRESERVATIVE FREE, LORAZEPAM
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 - METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 - ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 - PENTOSTATIN, PENTOSTATIN
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TORSEMIDE, TORSEMIDE

BEIJING DOUBLE CRANE

- * BEIJING DOUBLE CRANE PHARMACEUTICAL CO LTD
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - MELOXICAM, MELOXICAM

BELCHER PHARMS

- * BELCHER PHARMACEUTICALS LLC
 - CEPHALEXIN, CEPHALEXIN

BEN VENUE

- * BEN VENUE LABORATORIES INC
 - FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

BEXIMCO USA

- * BEXIMCO PHARMACEUTICALS USA INC
 - CARVEDILOL, CARVEDILOL

BIO NUCLEONICS

- * BIO NUCLEONICS INC
 - STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89

BIOALLIANCE PHARMA

- * BIOALLIANCE PHARMA
 - ORAVIG, MICONAZOLE

BIOKEY

- * BIOKEY INC
 - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 - LEVETIRACETAM, LEVETIRACETAM

BIOMARIN PHARM

- * BIOMARIN PHARMACEUTICAL INC
 - KUVAN, SAPROPTERIN DIHYDROCHLORIDE

BIONICHE PHARMA

- * BIONICHE PHARMA USA LLC
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 - COSYNTROPIN, COSYNTROPIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BIONICHE PHARMA USA LLC
 - DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
 - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - ENLON, EDROPHONIUM CHLORIDE
 - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
 - FLUOROURACIL, FLUOROURACIL
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 - MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
 - METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 - RIMSO-50, DIMETHYL SULFOXIDE
 - SOTRADECOL, SODIUM TETRADECYL SULFATE

BIONICHE PHARMA USA

- * BIONICHE PHARMA USA LLC
 - CARBOPLATIN, CARBOPLATIN
 - DURACLON, CLONIDINE HYDROCHLORIDE
 - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - FOMEPIZOLE, FOMEPIZOLE
 - IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
 - METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
 - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

BIOVAIL TECHNOLOGIES

- * BIOVAIL TECHNOLOGIES LTD
 - MEPHYTON, PHYTONADIONE

BLAIREX

- * BLAIREX LABORATORIES INC
 - BRONCHO SALINE, SODIUM CHLORIDE (OTC)

BLU CARIBE

- * BLU CARIBE INC
 - GEMFIBROZIL, GEMFIBROZIL
 - SIMVASTATIN, SIMVASTATIN

BOCA PHARMA

- * BOCA PHARMACAL INC
 - ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
 - ALPRAZOLAM, ALPRAZOLAM
 - CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - LEVETIRACETAM, LEVETIRACETAM
 - METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE

BOEHRINGER INGELHEIM

- * BOEHRINGER INGELHEIM
 - CATAPRES, CLONIDINE HYDROCHLORIDE
 - CATAPRES-TTS-1, CLONIDINE
 - CATAPRES-TTS-2, CLONIDINE
 - CATAPRES-TTS-3, CLONIDINE
 - MICARDIS HCT, HYDROCHLOROTHIAZIDE
 - MICARDIS, TELMISARTAN
 - MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
 - ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
 - ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)
- * BOEHRINGER INGELHEIM PHARMACEUTICALS INC
 - AGGRENOX, ASPIRIN
 - APTIVUS, TIPRANAVIR
 - ATROVENT HFA, IPRATROPIUM BROMIDE
 - ATROVENT, IPRATROPIUM BROMIDE
 - COMBIVENT RESPIMAT, ALBUTEROL SULFATE
 - COMBIVENT, ALBUTEROL SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

- * BOEHRINGER INGELHEIM PHARMACEUTICALS INC
FLOMAX, TAMSULOSIN HYDROCHLORIDE
MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE
MOBIC, MELOXICAM
PERSANTINE, DIPYRIDAMOLE
PRADAXA, DABIGATRAN ETEXILATE MESYLATE
SPIRIVA, TIOTROPiUM BROMIDE MONOHYDRATE
TRADJENTA, LINAGLiptin
TWYNSTA, AMLODiPINE BESYLATE
VIRAMUNE XR, NEViRAPiNE
VIRAMUNE, NEViRAPiNE

BRACCO

- * BRACCO DIAGNOSTICS INC
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
CHOLETEC, TECHNETiUM TC-99M MEBROFENiN KIT
CHOLOGRAFiN MEGlUMiNE, iODiPAMiDE MEGlUMiNE
CHROMITOPE SODiUM, SODiUM CHROMATE CR-51
CYSTOGRAFiN DiLiTE, DiATRiZOATE MEGlUMiNE
CYSTOGRAFiN, DiATRiZOATE MEGlUMiNE
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
MDP-BRACCO, TECHNETiUM TC-99M MEDRONATE KIT
MULTiHANCE MULTiPACK, GADOBENaTE DiMEGLUMiNE
MULTiHANCE, GADOBENaTE DiMEGLUMiNE
PROHANCE MULTiPACK, GADOTERiDOL
PROHANCE, GADOTERiDOL
RENOGRAFiN-76, DiATRiZOATE MEGlUMiNE
SiNOGRAFiN, DiATRiZOATE MEGlUMiNE

BRAINTREE

- * BRAINTREE LABORATORIES INC
AXiD, NiZATiDiNE
GOlyTELY, POLyETHYLENE GLYCOL 3350
HALFLyTELY, BiSACODYL
NULyTELY, POLyETHYLENE GLYCOL 3350
NULyTELY-FLAVORED, POLyETHYLENE GLYCOL 3350

BRAINTREE LABS

- * BRAINTREE LABORATORIES INC
SUPREP BOWEL PREP KIT, MAGNEsiUM SULFATE ANHYDROUS

BRECKENRIDGE PHARM

- * BRECKENRIDGE PHARMACEUTICAL INC
CiLOSTAZOL, CiLOSTAZOL
ESTRADIOL AND NORETHiNDRONE ACETATE, ESTRADIOL
LEVEtiRACETAM, LEVEtiRACETAM
MELOXiCAM, MELOXiCAM
METHSCOPOLAMiNE BROMiDE, METHSCOPOLAMiNE BROMiDE
OXCARBAZEPiNE, OXCARBAZEPiNE
POLyETHYLENE GLYCOL 3350, POLyETHYLENE GLYCOL 3350
PRAMIPEXOLE DiHYDROCHLORiDE, PRAMIPEXOLE DiHYDROCHLORiDE
TERBiNAFiNE HYDROCHLORiDE, TERBiNAFiNE HYDROCHLORiDE

BRIGHTSTONE

- * BRIGHTSTONE PHARMA INC
iSOSORBiDE MONONiTRATE, iSOSORBiDE MONONiTRATE

BRiSTOL

- * BRiSTOL LABORATORiES INC DiV BRiSTOL MYERS CO
BiCNU, CARMuSTiNE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ******BRISTOL MYERS SQUIBB**

- * BRISTOL MYERS SQUIBB
 - AZACTAM, AZTREONAM
 - BARACLUDE, ENTECAVIR
 - GLUCOVANCE, GLYBURIDE
 - KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE
 - LYSODREN, MITOTANE
 - MEGACE, MEGESTROL ACETATE
 - PRAVACHOL, PRAVASTATIN SODIUM
- * BRISTOL MYERS SQUIBB CO
 - AZACTAM IN PLASTIC CONTAINER, AZTREONAM
 - CEENU, LOMUSTINE
 - DROXIA, HYDROXYUREA
 - GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
 - HYDREA, HYDROXYUREA
 - IXEMpra KIT, IXABEPILONE
 - ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE
 - REYATAZ, ATAZANAVIR SULFATE
 - SPRYCEL, DASATINIB
 - SUSTIVA, EFAVIRENZ
 - VIDEX EC, DIDANOSINE
- * BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
 - ESTRACE, ESTRADIOL
 - ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
 - GLUCOPHAGE, METFORMIN HYDROCHLORIDE
 - MONOPRIL, FOSINOPRIL SODIUM
 - VIDEX, DIDANOSINE
 - VUMON, TENIPOSIDE
 - ZERIT, STAVUDINE
- * BRISTOL MYERS SQUIBB PHARMA CO
 - COUMADIN, WARFARIN SODIUM
 - SUSTIVA, EFAVIRENZ

BRYAN

- * BRYAN CORP
 - SCLEROSOL, TALC
 - TALC, TALC

CADENCE PHARMS

- * CADENCE PHARMACEUTICALS INC
 - OFIRMEV, ACETAMINOPHEN

CADISTA PHARMS

- * CADISTA PHARMACEUTICALS INC
 - ALENDRONATE SODIUM, ALENDRONATE SODIUM
 - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 - LAMOTRIGINE, LAMOTRIGINE
 - OXCARBAZEPINE, OXCARBAZEPINE

CANYON

- * CANYON PHARMACEUTICALS INC
 - IPRIVASK, DESIRUDIN RECOMBINANT

CARACO

- * CARACO PHARMACEUTICAL LABORATORIES LTD
 - ALLOPURINOL, ALLOPURINOL
 - AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - ATENOLOL, ATENOLOL
 - BACLOFEN, BACLOFEN
 - CARVEDILOL, CARVEDILOL
 - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLONAZEPAM, CLONAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

- * CARACO PHARMACEUTICAL LABORATORIES LTD
 - CLOZAPINE, CLOZAPINE
 - DIGOXIN, DIGOXIN
 - ELIXOPHYLLIN, THEOPHYLLINE
 - FLUMADINE, RIMANTADINE HYDROCHLORIDE
 - FLURBIPROFEN, FLURBIPROFEN
 - FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 - GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 - GLIPIZIDE, GLIPIZIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 - MELOXICAM, MELOXICAM
 - MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METHIMAZOLE, METHIMAZOLE
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - MIRTAZAPINE, MIRTAZAPINE
 - OXaprozin, OXaprozin
 - PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
 - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 - SYNALGOS-DC, ASPIRIN
 - THEOCHRON, THEOPHYLLINE
 - TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CARDINAL HEALTH 414

- * CARDINAL HEALTH 414 LLC
 - TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

CARDINAL HLTH

- * CARDINAL HEALTH
 - CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
 - CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
 - CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

CAREFUSION

- * CAREFUSION 213 LLC
 - CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
 - CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
 - CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)
 - PHARMASEAL SCRUB CARE, CHLORHEXIDINE GLUCONATE (OTC)

CARLSBAD

- * CARLSBAD TECHNOLOGY INC
 - ACYCLOVIR, ACYCLOVIR
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - FAMOTIDINE, FAMOTIDINE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - GLIMEPIRIDE, GLIMEPIRIDE
 - LOVASTATIN, LOVASTATIN
 - MELOXICAM, MELOXICAM
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

CAROLINA MEDCL

- * CAROLINA MEDICAL PRODUCTS CO
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** C **

CATALENT

* CATALENT PHARMA SOLUTIONS LLC
NIFEDIPINE, NIFEDIPINE
VALPROIC ACID, VALPROIC ACID

CEDAR PHARMS

* CEDAR PHARMACEUTICALS LLC
METHIMAZOLE, METHIMAZOLE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

CELGENE

* CELGENE CORP
ISTODAX, ROMIDEPSIN
REVLIMID, LENALIDOMIDE
THALOMID, THALIDOMIDE
VIDAZA, AZACITIDINE

CEPHALON

* CEPHALON INC
ACTIQ, FENTANYL CITRATE
FENTORA, FENTANYL CITRATE
GABITRIL, TIAGABINE HYDROCHLORIDE
NUVIGIL, ARMODAFINIL
PROVIGIL, MODAFINIL
TREANDA, BENDAMUSTINE HYDROCHLORIDE
TRISENOX, ARSENIC TRIOXIDE

CEPHAZONE PHARMA

* CEPHAZONE PHARMA LLC
CEFАЗОЛИН СODIUM, CEFАЗОЛИН SODIUM
CEFOTАКСИМ SODIUM, CEFOTАКСИМ SODIUM
CEFTRIАКСИМ, CEFTRIАКСИМ SODIUM

CEREXA

* CEREXA INC
TEFLARO, CEFTAROLINE FOSAMIL

CHATTEM

* CHATTEM INC
SELSUN, SELENIUM SULFIDE
UNISOM, DOXYLAMINE SUCCINATE (OTC)

CHEMISCH FBRK KRSSLR

* CHEMISCHE FABRIK KREUSSLER & CO. GMBH
ASCLERA, POLIDOCANOL

CHIRHOCLIN

* CHIRHOCLIN INC
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

CIPHER PHARMS INC

* CIPHER PHARMACEUTICALS INC
CONZIP, TRAMADOL HYDROCHLORIDE
LIPOFEN, FENOFIBRATE

CIPLA

* CIPLA LTD
RAMIPRIL, RAMIPRIL
RISPERIDONE, RISPERIDONE
TRANDOLAPRIL, TRANDOLAPRIL
ZIDOVUDINE, ZIDOVUDINE

CIPLA LTD

* CIPLA LTD
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
STAVUDINE, STAVUDINE
TOPIRAMATE, TOPIRAMATE
ZALEPLON, ZALEPLON
ZIDOVUDINE, ZIDOVUDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******CITIUS PHARMS**

- * CITIUS PHARMACEUTICALS LLC
SUPRENZA, PHENTERMINE HYDROCHLORIDE

CITRUSPHRMA

- * CITRUSPHRMA LLC
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

CLARIS LIFESCIENCES

- * CLARIS LIFESCIENCES LTD
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPROFLOXACIN, CIPROFLOXACIN
DIPYRIDAMOLE, DIPYRIDAMOLE
FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK), FAMOTIDINE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUMAZENIL, FLUMAZENIL
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FULCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE
MILRINONE LACTATE, MILRINONE LACTATE
ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE

CLINIGEN HLTHCARE

- * CLINIGEN HEALTHCARE LTD
FOSCAVIR, FOSCARNET SODIUM

CLINTEC NUTR

- * CLINTEC NUTRITION CO SUB CLINIQUE
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

CLONMEL HLTHCARE

- * CLONMEL HEALTHCARE LTD
MORPHINE SULFATE, MORPHINE SULFATE

CNS THERAPS INC

- * CNS THERAPEUTICS INC
GABLOFEN, BACLOFEN

CNTY LINE PHARMS

- * COUNTY LINE PHARMACEUTICALS LLC
UREX, METHENAMINE HIPPURATE

COASTAL PHARMS

- * COASTAL PHARMACEUTICALS
BROMFENAC SODIUM, BROMFENAC SODIUM
NYSTATIN, NYSTATIN
OXYCODONE AND ASPIRIN, ASPIRIN
- * COASTAL PHARMACEUTICALS INC
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

COBALT LABS INC

- * COBALT LABORATORIES INC
LEVETIRACETAM, LEVETIRACETAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ******COLGATE**

* COLGATE ORAL PHARMACEUTICALS INC
 ORABASE HCA, HYDROCORTISONE ACETATE
 PERIOGARD, CHLORHEXIDINE GLUCONATE

COLGATE PALMOLIVE

* COLGATE PALMOLIVE
 COLGATE TOTAL, SODIUM FLUORIDE (OTC)

CONNECTICS

* CONNECTICS CORP
 LUXIQ, BETAMETHASONE VALERATE

CONTRACT PHARMA

* CONTRACT PHARMACAL CORP
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)

CONTRACT PHARMACAL

* CONTRACT PHARMACAL CORP
 CIMETIDINE, CIMETIDINE (OTC)
 FOLIC ACID, FOLIC ACID
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
 PREDNISONE, PREDNISONE
 PROFEN, IBUPROFEN (OTC)
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

CONTROLLED THERAP

* CONTROLLED THERAPEUTICS (SCOTLAND) LTD
 CERVIDIL, DINOPROSTONE

COREPHARMA

* COREPHARMA LLC
 ALBENZA, ALBENDAZOLE
 ALPRAZOLAM, ALPRAZOLAM
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CARISOPRODOL, CARISOPRODOL
 CILOSTAZOL, CILOSTAZOL
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DARAPRIM, PYRIMETHAMINE
 DECLOMYCIN, DEMECLOCYCLINE HYDROCHLORIDE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
 DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 E-GLADES, ERYTHROMYCIN
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 GLCOPYRROLATE, GLCOPYRROLATE
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 HYDROCORTISONE, HYDROCORTISONE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LEVOCARNITINE, LEVOCARNITINE
 MELOXICAM, MELOXICAM
 METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 POTASSIUM CITRATE, POTASSIUM CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** C ****

- * COREPHARMA LLC
 - PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 - RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TRANDOLAPRIL, TRANDOLAPRIL
 - URSODIOL, URSDIOL
 - ZONISAMIDE, ZONISAMIDE

CORIA

- * CORIA LABORATORIES LTD
 - NUTRACORT, HYDROCORTISONE

CORNERSTONE THERAP

- * CORNERSTONE THERAPEUTICS INC
 - CUROSURF, PORACTANT ALFA
 - FACTIVE, GEMIFLOXACIN MESYLATE
 - SPECTRAZEF, CEFEDITOREN PIVOXIL
 - ZYFLO CR, ZILEUTON
 - ZYFLO, ZILEUTON

COVIDIEN

- * COVIDIEN
 - LYMPHAZURIN, ISOSULFAN BLUE
 - THERMAZENE, SILVER SULFADIAZINE

COVIS PHARMA

- * COVIS PHARMA SARL
 - LANOXIN PEDIATRIC, DIGOXIN
 - LANOXIN, DIGOXIN

CPPI CV

- * CP PHARMACEUTICALS INTERNATIONAL CV
 - LYRICA, PREGABALIN
 - SUTENT, SUNITINIB MALATE

CROWN LABS

- * CROWN LABORATORIES INC
 - ALA-CORT, HYDROCORTISONE
 - ALA-SCALP, HYDROCORTISONE
 - TRIDERM, TRIAMCINOLONE ACETONIDE

CSL BEHRING

- * CSL BEHRING LLC
 - STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE

CUBIST

- * CUBIST PHARMACEUTICALS INC
 - CUBICIN, DAPTOMYCIN

CUMBERLAND PHARMS

- * CUMBERLAND PHARMACEUTICALS INC
 - ACETADOTE, ACETYL CYSTEINE
 - CALDOLOR, IBUPROFEN

CYPRESS BIOSCIENCE

- * CYPRESS BIOSCIENCE INC
 - SAVELLA, MILNACIPRAN HYDROCHLORIDE

CYPRESS PHARM

- * CYPRESS PHARMACEUTICAL INC
 - CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - ELIPHOS, CALCIUM ACETATE
 - EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 - LEVETIRACETAM, LEVETIRACETAM
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - REZIRA, HYDROCODONE BITARTRATE
 - ZUTRIPRO, CHLORPHENIRAMINE MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******DAIICHI**

- * DAIICHI PHARMACEUTICAL CORP
FLOXIN OTIC, OFLOXACIN

DAIICHI SANKYO

- * DAIICHI SANKYO INC
AZOR, AMLODIPINE BESYLATE
BENICAR HCT, HYDROCHLOROTHIAZIDE
BENICAR, OLMESARTAN MEDOXOMIL
TRIBENZOR, AMLODIPINE BESYLATE
WELCHOL, COLESEVELAM HYDROCHLORIDE

DAIICHI SANKYO CO

- * DAIICHI SANKYO CO LTD
EVOXAC, CEVIMELINE HYDROCHLORIDE

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
CIMETIDINE, CIMETIDINE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DIAZEPAM, DIAZEPAM
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
FUROSEMIDE, FUROSEMIDE
GEMFIBROZIL, GEMFIBROZIL
GLYBURIDE (MICRONIZED), GLYBURIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
NAPROXEN, NAPROXEN
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
PROPYLTIIOURACIL, PROPYLTIIOURACIL
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

DENCO ASSET

- * DENCO ASSET LLC
DENAVIR, PENCICLOVIR SODIUM

DENTSPLY PHARM

- * DENTSPLY PHARMACEUTICAL
CITANESE PLAIN DENTAL, PRILOCAINE HYDROCHLORIDE
ORAQIX, LIDOCAINE
POLOCaine, MEPIVACAINE HYDROCHLORIDE
XYLOCAINE DENTAL WITH EPINEPHRINE, EPINEPHRINE
XYLOCAINE DENTAL, LIDOCAINE HYDROCHLORIDE
- * DENTSPLY PHARMACEUTICAL INC
CITANESE FORTE DENTAL, EPINEPHRINE BITARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ******DEPOMED INC**

- * DEPOMED INC
GRALISE, GABAPENTIN

DEPROCO

- * DEPROCO INC
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
SCANDONEST L, LEVONORDEFRIN
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SEPTOCAIN, ARTICAINE HYDROCHLORIDE

DEXCEL LTD

- * DEXCEL LTD
DICLOFENAC SODIUM, DICLOFENAC SODIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

DEXCEL PHARMA

- * DEXCEL PHARMA TECHNOLOGIES LTD
OMEPRAZOLE, OMEPRAZOLE (OTC)
PERIOCHIP, CHLORHEXIDINE GLUCONATE

DEY

- * DEY LP
ACCUNEB, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
CROMOLYN SODIUM, CROMOLYN SODIUM
DUONEB, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE

DEY PHARMA

- * DEY PHARMA LP
PERFOROMIST, FORMOTEROL FUMARATE

DIALYSIS SUPS

- * DIALYSIS SUPPLIES INC
NORMOCARB HF 25, MAGNESIUM CHLORIDE
NORMOCARB HF 35, MAGNESIUM CHLORIDE

DORC

- * DORC INTERNATIONAL BV
MEMBRANEBLUE, TRYPLAN BLUE
VISIONBLUE, TRYPLAN BLUE

DOW PHARM SCI

- * DOW PHARMACEUTICAL SCIENCES
ACANYA, BENZOYL PEROXIDE

DOW PHARM SCIENCES

- * DOW PHARMACEUTICAL SCIENCES INC
AKNE-MYCIN, ERYTHROMYCIN
ATRALIN, TRETINOIN
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE

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

- * DR REDDYS LABORATORIES LTD
OMEPRAZOLE, OMEPRAZOLE

DR REDDYS LABS INC

- * DR REDDYS LABORATORIES INC
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DR REDDYS LABORATORIES INC
 - AMOXIL, AMOXICILLIN
 - AUGMENTIN '125', AMOXICILLIN
 - AUGMENTIN '200', AMOXICILLIN
 - AUGMENTIN '250', AMOXICILLIN
 - AUGMENTIN '400', AMOXICILLIN
 - AUGMENTIN '500', AMOXICILLIN
 - AUGMENTIN '875', AMOXICILLIN
 - AUGMENTIN ES-600, AMOXICILLIN
 - AUGMENTIN XR, AMOXICILLIN
 - FINASTERIDE, FINASTERIDE
 - FLUCONAZOLE, FLUCONAZOLE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 - GRANisetron HYDROCHLORIDE PRESERVATIVE FREE, GRANisetron HYDROCHLORIDE
 - GRANisetron HYDROCHLORIDE, GRANisetron 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)
 - NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 - SIMVASTATIN, SIMVASTATIN
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

DR REDDYS LABS LTD

- * DR REDDYS LABORATORIES LIMITED
 - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- * DR REDDYS LABORATORIES LTD
 - ALENDRONATE SODIUM, ALENDRONATE SODIUM
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - ANASTROZOLE, ANASTROZOLE
 - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 - CARVEDILOL, CARVEDILOL
 - 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)
 - 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
 - DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
 - DESLORATADINE, DESLORATADINE
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 - FAMOTIDINE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 - FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE (OTC)
 - FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 - FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 - FINASTERIDE, FINASTERIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DR REDDYS LABORATORIES LTD
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - GLIMEPIRIDE, GLIMEPIRIDE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 - IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - LAMOTRIGINE, LAMOTRIGINE
 - LANSOPRAZOLE, LANSOPRAZOLE
 - LETROZOLE, LETROZOLE
 - LEVETIRACETAM, LEVETIRACETAM
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 - NABUMETONE, NABUMETONE
 - NAPROXEN SODIUM, NAPROXEN SODIUM
 - NATEGLINIDE, NATEGLINIDE
 - NIZATIDINE, NIZATIDINE
 - OFLOXACIN, OFLOXACIN
 - OLANZAPINE, OLANZAPINE
 - OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
 - OMEPRAZOLE, OMEPRAZOLE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXaprozin, OXaprozin
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - PRIMIDONE, PRIMIDONE
 - RAMIPRIL, RAMIPRIL
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 - RISPERIDONE, RISPERIDONE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - TACROLIMUS, TACROLIMUS
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
 - TRANDOLAPRIL, TRANDOLAPRIL
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - ZAFIRLUKAST, ZAFIRLUKAST
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 - ZONISAMIDE, ZONISAMIDE

DRAXIMAGE

- * DRAXIMAGE INC
 - DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE
 - DTPA, TECHNETIUM TC-99M PENTETATE KIT
 - HICON, SODIUM IODIDE I-131
 - SODIUM IODIDE I-131, SODIUM IODIDE I-131
 - TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 - TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

DURAMED

- * DURAMED PHARMACEUTICALS INC
 - NORDETTE-28, ETHINYL ESTRADIOL
 - PLAN B ONE-STEP, LEVONORGESTREL
 - PLAN B ONE-STEP, LEVONORGESTREL (OTC)
 - REVIA, NALTREXONE HYDROCHLORIDE

DURAMED PHARMS BARR

- * DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
 - AVIANE-28, ETHINYL ESTRADIOL
 - CRYSELLE, ETHINYL ESTRADIOL
 - DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 - DIAMOX, ACETAZOLAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** D ****

- * DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
DIAMOX, ACETAZOLAMIDE SODIUM
ENPRESSE-28, ETHINYL ESTRADIOL
METHYLPREDNISOLONE, METHYLPREDNISOLONE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VELIVET, DESOGESTREL
ZEBETA, BISOPROLOL FUMARATE
ZIAC, BISOPROLOL FUMARATE

DURAMED RES

- * DURAMED RESEARCH INC
AYGESTIN, NORETHINDRONE ACETATE
PARAGARD T 380A, COPPER
SEASONALE, ETHINYL ESTRADIOL

DUSA

- * DUSA PHARMACEUTICALS INC
LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

EAGLE PHARMS

- * EAGLE PHARMACEUTICALS INC
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN

EASTMAN KODAK

- * EASTMAN KODAK CO
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE

EBEWE PHARMA

- * EBEWE PHARMA
GRANisetron Hydrochloride, GRANisetron Hydrochloride
- * EBEWE PHARMA GES MBH NFG KG
CARBOPLATIN, CARBOPLATIN
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
FLUOROURACIL, FLUOROURACIL
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
OXALIPLATIN, OXALIPLATIN
PACLITAXEL, PACLITAXEL
VINORELBINE TARTRATE, VINORELBINE TARTRATE

ECOLAB

- * ECOLAB INC
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

ECR

- * ECR PHARMACEUTICALS
DEXAMETHASONE, DEXAMETHASONE

EDGEMONT PHARMS LLC

- * EDGEMONT PHARMACEUTICALS LLC
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE

EGIS

- * EGIS PHARMACEUTICALS
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE

EI INC

- * EI INC
THEROXIDIL, MINOXIDIL (OTC)

EISAI INC

- * EISAI INC
ACIPHEX, RABEPRAZOLE SODIUM
ARICEPT ODT, DONEPEZIL HYDROCHLORIDE
ARICEPT, DONEPEZIL HYDROCHLORIDE
BANZEL, RUFINAMIDE
DACOGEN, DECITABINE
FRAGMIN, DALTEPARIN SODIUM
GLIADEL, CARMUSTINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

- * EISAI INC
 - HALAVEN, ERIBULIN MESYLATE
 - HEXALEN, ALTRETAMINE
 - LUSEDRA, FOSPROPOFOL DISODIUM
 - PANRETIN, ALITRETNINOIN
 - SALAGEN, PILOCARPINE HYDROCHLORIDE
 - TARGRETIN, BEXAROTENE
 - ZONEGRAN, ZONISAMIDE

EKR THERAP

- * EKR THERAPEUTICS INC
 - 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 SR, NICARDIPINE HYDROCHLORIDE
 - CARDENE, NICARDIPINE HYDROCHLORIDE
 - DEPODUR, MORPHINE SULFATE

ELAN DRUG

- * ELAN DRUG DELIVERY INC
 - VERELAN PM, VERAPAMIL HYDROCHLORIDE

ELAN PHARM

- * ELAN PHARMACEUTICAL RESEARCH CORP
 - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 - KETOPROFEN, KETOPROFEN

ELI LILLY AND CO

- * ELI LILLY AND CO
 - AXIRON, TESTOSTERONE
 - EFFIENT, PRASUGREL HYDROCHLORIDE

ELI LILLY CO

- * 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

ELKINS SINN

- * ELKINS SINN DIV AH ROBINS CO INC
 - MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE

EMCURE PHARMS

- * EMCURE PHARMACEUTICALS LTD
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE

EMCURE PHARMS USA

- * EMCURE PHARMACEUTICALS USA INC
 - FOSINOPRIL SODIUM AND HYDROCHLORTIAZIDE, FOSINOPRIL
 - METHIMAZOLE, METHIMAZOLE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

EMCURE USA

- * EMCURE PHARMACEUTICALS USA INC
 - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE

EMD SERONO

- * EMD SERONO INC
 - CETROTIDE, CETRORELIX
 - GONAL-F RFF PEN, FOLLITROPIN ALFA/BETA
 - GONAL-F RFF, FOLLITROPIN ALFA/BETA
 - GONAL-F, FOLLITROPIN ALFA/BETA
 - LUVERIS, LUTROPIN ALFA
 - OVIDREL, CHORIOGONADOTROPIN ALFA
 - SAIZEN, SOMATROPIN RECOMBINANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

- * EMD SERONO INC
SEROPHENE, CLOMIPHENE CITRATE
SEROSTIM, SOMATROPIN RECOMBINANT
ZORBTIVE, SOMATROPIN RECOMBINANT

EMMAUS MEDCL

- * EMMAUS MEDICAL INC
NUTRESTORE, GLUTAMINE

ENDO PHARM

- * ENDO PHARMACEUTICAL SOLUTIONS INC
DELATESTRYL, TESTOSTERONE ENANTHATE
SUPPRELIN LA, HISTRELIN ACETATE
VALSTAR PRESERVATIVE FREE, VALRUBICIN
VANTAS, HISTRELIN ACETATE

ENDO PHARMS

- * ENDO PHARMACEUTICALS INC
FELODIPINE, FELODIPINE
FORTESTA, TESTOSTERONE
FROVA, FROVATRIPTAN SUCCINATE
LETROZOLE, LETROZOLE
MORPHINE SULFATE, MORPHINE SULFATE
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
OPANA ER, OXYMORPHONE HYDROCHLORIDE
OPANA, OXYMORPHONE HYDROCHLORIDE
PERCOSET, ACETAMINOPHEN
PERCODAN, ASPIRIN
ZYDONE, ACETAMINOPHEN

EPIC PHARMA

- * EPIC PHARMA INC
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
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 LLC

- * EPIC PHARMA LLC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

ERGOJECT

- * ERGOJECT LLC
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE

ETHYPHARM

- * ETHYPHARM
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

EURAND

- * EURAND AMERICA INC
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

EUSA PHARMA USA

- * EUSA PHARMA (USA) INC
QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

EXALENZ BIOSCIENCE

- * EXALENZ BIOSCIENCE LTD
IDKIT:HP, UREA C-13

EXCELLIUM

- * EXCELLIUM PHARMACEUTICAL INC
FOLIC ACID, FOLIC ACID
FUROSEMIDE, FUROSEMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** E ****

- * EXCELLIUM PHARMACEUTICAL INC
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LORAZEPAM, LORAZEPAM

EXELA PHARMA SCIENCE

- * EXELA PHARMA SCIENCES
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

EYETECH INC

- * EYETECH INC
MACUGEN, PEGAPTANIB SODIUM

FACTA FARMA

- * FACTA FARMACEUTICI SPA
CEPHALEXIN, CEPHALEXIN

FALCON PHARMS

- * FALCON PHARMACEUTICALS INC
TIMOLOL MALEATE, TIMOLOL MALEATE
- * FALCON PHARMACEUTICALS LTD
DIPIVEFRIN HYDROCHLORIDE, DIPIVEFRIN HYDROCHLORIDE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
MAXITROL, DEXAMETHASONE
METIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TOBREX, TOBRAMYCIN
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

FDC LTD

- * FDC LTD
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
OFLOXACIN, OFLOXACIN
TIMOLOL MALEATE, TIMOLOL MALEATE

FEINSTEIN

- * FEINSTEIN INSTITUTE MEDICAL RESEARCH
AMMONIA N 13, AMMONIA, N-13
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

FERA PHARMS

- * FERA PHARMACEUTICALS LLC
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
OFLOXACIN, OFLOXACIN
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TOBRAMYCIN, TOBRAMYCIN

FERNDALE LABS

- * FERNDALE LABORATORIES INC
HYDROCORTISONE ACETATE, HYDROCORTISONE ACETATE
MICORT-HC, HYDROCORTISONE ACETATE
PRAMOSONE, HYDROCORTISONE ACETATE

FERRING

- * FERRING PHARMACEUTICALS INC
ACTHREL, CORTICORELIN OVINE TRIFLUTATE
BRAVELLE, UROFOLLITROPIN
CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
ENDOMETRIN, PROGESTERONE
FIRMAGON, DEGARELIX ACETATE
MENOPUR, LUTEINIZING HORMONE
MINIRIN, DESMOPRESSIN ACETATE
REPRONEX, LUTEINIZING HORMONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

- * FERRING PHARMACEUTICALS INC
TEV-TROPIN, SOMATROPIN RECOMBINANT

FERRING PHARMS AS

- * FERRING PHARMACEUTICALS AS
LYSTEDA, TRANEXAMIC ACID

FLEMING

- * FLEMING AND CO PHARMACEUTICALS
THYROSHIELD, POTASSIUM IODIDE (OTC)

FOREST LABS

- * FOREST LABORATORIES INC
BYSTOLIC, NEBIVOLOL HYDROCHLORIDE
CAMPRAL, ACAMPROSATE CALCIUM
CELEXA, CITALOPRAM HYDROBROMIDE
LEXAPRO, ESCITALOPRAM OXALATE
NAMENDA XR, MEMANTINE HYDROCHLORIDE
NAMENDA, MEMANTINE HYDROCHLORIDE
TESSALON, BENZONATATE
THYROLAR-0.25, LIOTHYRONINE SODIUM
THYROLAR-0.5, LIOTHYRONINE SODIUM
THYROLAR-1, LIOTHYRONINE SODIUM
THYROLAR-2, LIOTHYRONINE SODIUM
THYROLAR-3, LIOTHYRONINE SODIUM

FOREST LABS INC

- * FOREST LABORATORIES INC
VIIBRYD, VILAZODONE HYDROCHLORIDE

FOREST RES INST INC

- * FOREST RESEARCH INSTITUTE INC
DALIRESP, ROFLUMILAST

FOUGERA

- * E FOUGERA DIV ALTANA INC
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

FOUGERA PHARMS

- * FOUGERA PHARMACEUTICALS INC
ACLOVATE, ALCLOMETASONE DIPROPIONATE
CUTIVATE, FLUTICASONE PROPIONATE
ERYTHROMYCIN, ERYTHROMYCIN
OXISTAT, OXICONAZOLE NITRATE
PAMINE FORTE, METHSCOPOLAMINE BROMIDE
PAMINE, METHSCOPOLAMINE BROMIDE
PANDEL, HYDROCORTISONE PROBUTATE
SOLARAZE, DICLOFENAC SODIUM
TEMOVATE E, CLOBETASOL PROPIONATE

FRESENIUS

- * FRESENIUS KABI DEUTSCHLAND GMBH
INTRALIPID 10%, SOYBEAN OIL
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
- * FRESENIUS USA INC
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** F ****

- * FRESENIUS USA INC
 - INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

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
 - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 - OXALIPLATIN, OXALIPLATIN
 - PACLITAXEL, PACLITAXEL
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

FRESENIUS MEDCL

- * FRESENIUS MEDICAL CARE NORTH AMERICA
 - CALCITRIOL, CALCITRIOL
 - 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 3.5% 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
 - DELFLEX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - DELFLEX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - DELFLEX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - DELFLEX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 - PHOSLO GELCAPS, CALCIUM ACETATE
 - PHOSLYRA, CALCIUM ACETATE
 - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

FSC

- * FSC LABORATORIES INC
 - PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE

FUTURE PAK

- * FUTURE PAK LTD
 - K+10, POTASSIUM CHLORIDE

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)
 - MIGERGOT, CAFFEINE
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - PROCHLORPERAZINE, PROCHLORPERAZINE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** G **

G AND W LABS INC

- * G AND W LABORATORIES INC
CALCIPOTRIENE, CALCIPOTRIENE
CICLOPIROX, CICLOPIROX
METRONIDAZOLE, METRONIDAZOLE

GALDERMA LABS

- * GALDERMA LABORATORIES INC
CLOBEX, CLOBETASOL PROPIONATE
EPIDUO, ADAPALENE

GALDERMA LABS LP

- * GALDERMA LABORATORIES L P
CLOBEX, CLOBETASOL PROPIONATE
- * GALDERMA LABORATORIES LP
CAPEX, FLUOCINOLONE ACETONIDE
CLINDAGEL, CLINDAMYCIN PHOSPHATE
CLOBEX, CLOBETASOL PROPIONATE
DESOWEN, DESONIDE
DIFFERIN, ADAPALENE
LIDOCAINE AND TETRACAIN, LIDOCAINE
METROCREAM, METRONIDAZOLE
METROGEL, METRONIDAZOLE
METROLOTION, METRONIDAZOLE
METVIXIA, METHYL AMINOLEVULINATE HYDROCHLORIDE
ORACEA, DOXYCYCLINE
PERIOSTAT, DOXYCYCLINE HYCLATE
TRI-LUMA, FLUOCINOLONE ACETONIDE
VECTICAL, CALCITRIOL

GALDERMA R AND D

- * GALDERMA RESEARCH AND DEVELOPMENT INC
DIFFERIN, ADAPALENE

GALEN (UK)

- * GALEN LTD
DAUNOXOME, DAUNORUBICIN CITRATE

GALEN LTD

- * GALEN LTD
FEMRING, ESTRADIOL ACETATE

GAMBRO RENAL PRODS

- * GAMBRO RENAL PRODUCTS
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

GASTROENTERO

- * GASTROENTERO LOGIC LLC
OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN

GAVIS PHARMS

- * GAVIS PHARMACEUTICALS LLC
ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE, ACETAMINOPHEN
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PENTAZOCINE AND NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
QUINARETIC, HYDROCHLOROTHIAZIDE
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******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
 COVERA-HS, 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
 CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE
 INDICLOR, INDIUM IN-111 CHLORIDE
 INDIUM IN-111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
 METASTRON, STRONTIUM CHLORIDE SR-89
 MPI DMSA KIDNEY REAGENT, TECHNETIUM TC-99M SUCCIMER KIT
 MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
 MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
 MYOVIEW, 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
 SODIUM IODIDE I 123, SODIUM IODIDE I-123
 THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
 VISIPAQ 270, IODIXANOL
 VISIPAQ 320, IODIXANOL

GE HLTHCARE INC

* GE HEALTHCARE INC
 DATSCAN, IOFLUPANE I-123

GEDEON RICHTER USA

* GEDEON RICHTER USA INC
 FINASTERIDE, FINASTERIDE

GENENTECH

* GENENTECH INC
 NUTROPIN AQ PEN, SOMATROPIN RECOMBINANT
 NUTROPIN AQ, SOMATROPIN RECOMBINANT
 NUTROPIN, SOMATROPIN RECOMBINANT

GENERAMEDIX

* GENERAMEDIX INC
 SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX

GENTA

* GENTA INC
 GANITE, GALLIUM NITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******GENZYME**

- * GENZYME CORP
 - CEREDASE, ALGLUCERASE
 - CEREZYME, IMIGLUCERASE
 - CLOLAR, CLOFARABINE
 - FLUDARA, FLUDARABINE PHOSPHATE
 - MOZOBIL, PLERIXAFOR
 - RENAGEL, SEVELAMER HYDROCHLORIDE
 - RENVELA, SEVELAMER CARBONATE
 - THYROGEN, THYROTROPIN ALFA

GENZYME CORP

- * GENZYME CORP
 - HECTOROL, DOXERCALCIFEROL
- * GENZYME CORPORATION
 - HECTOROL, DOXERCALCIFEROL

GILEAD

- * GILEAD SCIENCES INC
 - ATRIPLA, EFAVIRENZ
 - CAYSTON, AZTREONAM
 - EMTRIVA, EMTRICITABINE
 - HEPSERA, ADEFOVIR DIPIVOXIL
 - LETAIRIS, AMBRISENTAN
 - RANEXA, RANOLAZINE
 - TRUVADA, EMTRICITABINE
 - VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 - VISTIDE, CIDOFOVIR

GILEAD SCIENCES INC

- * GILEAD SCIENCES INC
 - COMPLERA, EMTRICITABINE

GLAND PHARMA LTD

- * GLAND PHARMA LTD
 - ADENOSINE, ADENOSINE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - AZITHROMYCIN, AZITHROMYCIN
 - HALOPERIDOL, HALOPERIDOL LACTATE
 - MILRINONE LACTATE, MILRINONE LACTATE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE

GLAXO GRP LTD

- * GLAXO GROUP LTD
 - HORIZANT, GABAPENTIN ENACARBIL
- * GLAXO GROUP LTD DBA GLAXOSMITHKLINE
 - ALTABAX, RETAPAMULIN
 - FLOVENT HFA, FLUTICASONE PROPIONATE
 - SEREVENT, SALMETEROL XINAFOATE

GLAXOSMITHKLINE

- * GLAXOSMITHKLINE
 - ABREVA, DOCOSANOL (OTC)
 - ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
 - ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
 - ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
 - ADVAIR HFA, FLUTICASONE PROPIONATE
 - ALKERAN, MELPHALAN
 - ALKERAN, MELPHALAN HYDROCHLORIDE
 - AMERGE, NARatriptan HYDROCHLORIDE
 - ARIIXTRA, FONDAPARINUX SODIUM
 - AVODART, DUTASTERIDE
 - BACTROBAN, MUPIROCIN
 - BACTROBAN, MUPIROCIN CALCIUM
 - BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 - CEFTIN, CEFUROXIME AXETIL
 - EPIVIR-HBV, LAMIVUDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ****

- * GLAXOSMITHKLINE
 - FLONASE, FLUTICASONE PROPIONATE
 - FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 - FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 - FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
 - FORTAZ IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
 - FORTAZ, CEFTAZIDIME
 - HYCAMTIN, TOPOTECAN HYDROCHLORIDE
 - IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
 - IMITREX, SUMATRIPTAN
 - IMITREX, SUMATRIPTAN SUCCINATE
 - JALYN, DUTASTERIDE
 - LAMICTAL CD, LAMOTRIGINE
 - LAMICTAL, LAMOTRIGINE
 - MALARONE PEDIATRIC, ATOVAQUONE
 - MALARONE, ATOVAQUONE
 - NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
 - NICORETTE, NICOTINE POLACRILEX (OTC)
 - PARNATE, TRANYLCYPROMINE SULFATE
 - PAXIL CR, PAROXETINE HYDROCHLORIDE
 - PAXIL, PAROXETINE HYDROCHLORIDE
 - PROMACTA, ELTROMBOPAG OLAMINE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - RELENZA, ZANAMIVIR
 - REQUIP, ROPINIROLE HYDROCHLORIDE
 - TAGAMET HB, Cimetidine (OTC)
 - TAGAMET, Cimetidine
 - THIOGUANINE, THIOGUANINE
 - TIMENTIN IN PLASTIC CONTAINER, CLAVULANATE POTASSIUM
 - TIMENTIN, CLAVULANATE POTASSIUM
 - TREXIMET, NAPROXEN SODIUM
 - VALTREX, VALACYCLOVIR HYDROCHLORIDE
 - VENTOLIN HFA, ALBUTEROL SULFATE
 - VERAMYST, FLUTICASONE FUROATE
 - VOTRIENT, PAZOPANIB HYDROCHLORIDE
 - WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
 - WELLBUTRIN, BUPROPION HYDROCHLORIDE
 - ZANTAC 150, RANITIDINE HYDROCHLORIDE
 - ZANTAC 25, RANITIDINE HYDROCHLORIDE
 - ZANTAC 300, RANITIDINE HYDROCHLORIDE
 - ZANTAC IN PLASTIC CONTAINER, RANITIDINE HYDROCHLORIDE
 - ZANTAC, RANITIDINE HYDROCHLORIDE
 - ZINACEF IN PLASTIC CONTAINER, CEFUROXIME SODIUM
 - ZINACEF, CEFUROXIME SODIUM
 - ZOFRAN ODT, ONDANSETRON
 - ZOFRAN PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ZOFRAN, ONDANSETRON HYDROCHLORIDE
 - ZOVIRAX, ACYCLOVIR
 - ZYBAN, BUPROPION HYDROCHLORIDE

GLAXOSMITHKLINE CONS

- * GLAXOSMITHKLINE CONSUMER HEALTHCARE
 - ALLI, ORLISTAT (OTC)
 - COMMIT, NICOTINE POLACRILEX (OTC)
 - NICORETTE, NICOTINE POLACRILEX (OTC)

GLAXOSMITHKLINE LLC

- * GLAXOSMITHKLINE LLC
 - DYAZIDE, HYDROCHLOROTHIAZIDE
 - DYNACIRC CR, ISRADIPINE
 - FLOLAN, EPOPROSTENOL SODIUM
 - INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
 - MEPRON, ATOVAQUONE
 - RYTHMOL SR, PROPAFENONE HYDROCHLORIDE
 - RYTHMOL, PROPAFENONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******GLENMARK GENERICS**

- * GLENMARK GENERICS INC USA
ADAPALENE, ADAPALENE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CALCIPOTRIENE, CALCIPOTRIENE
DIPYRIDAMOLE, DIPYRIDAMOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
NILSTAT, NYSTATIN
NIZATIDINE, NIZATIDINE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
ZONISAMIDE, ZONISAMIDE
- * GLENMARK GENERICS LIMITED
BRIELLYN, ETHINYLMESTRADIOL
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- * GLENMARK GENERICS LTD
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
CARVEDILOL, CARVEDILOL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
FELODIPINE, FELODIPINE
FLUCONAZOLE, FLUCONAZOLE
GABAPENTIN, GABAPENTIN
HEATHER, NORETHINDRONE
LAMOTRIGINE, LAMOTRIGINE
LEVOFLOXACIN, LEVOFLOXACIN
LITHIUM CARBONATE, LITHIUM CARBONATE
MELOXICAM, MELOXICAM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
NORETHIDRONE, NORETHINDRONE
NORGESTIMATE AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON
OXCARBAZEPINE, OXCARBAZEPINE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
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, URSDIOL
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
- * GLENMARK GENERICS LTD INDIA
INDOMETHACIN, INDOMETHACIN
LITHIUM CARBONATE, LITHIUM CARBONATE
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 PHARMS LTD

- * GLENMARK PHARMACEUTICALS LTD
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******GRACEWAY**

- * GRACEWAY PHARMACEUTICALS LLC
MINITRAN, NITROGLYCERIN
THEOLAIR, THEOPHYLLINE

GRANULES INDIA

- * GRANULES INDIA LTD
IBUPROFEN, IBUPROFEN (OTC)
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

GRIFFEN

- * KW GRIFFEN CO
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

GTX INC

- * GTX INC
FARESTON, TOREMIFENE CITRATE

GUARDIAN DRUG

- * GUARDIAN DRUG CO INC
FOAMCOAT, ALUMINUM HYDROXIDE (OTC)

GUERBET

- * GUERBET LLC
HEXBABIX, IOXAGLATE MEGLUMINE
OXILAN-300, IOXILAN
OXILAN-350, IOXILAN

HAEMONETICS

- * HAEMONETICS CORP
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HALOCARBON PRODS

- * HALOCARBON PRODUCTS CORP
ISOFLURANE, ISOFLURANE
SEVOFLURANE, SEVOFLURANE

HALOZYME THERAP

- * HALOZYME THERAPEUTICS INC
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN

HANFORD GC

- * GC HANFORD MANUFACTURING CO
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

HARRIS PHARM

- * HARRIS PHARMACEUTICAL INC
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

HEC PHARM INC

- * HEC PHARM INC
ZIDOVUDINE, ZIDOVUDINE

HELSINN HLTHCARE

- * HELSINN HEALTHCARE SA
ALOXI, PALONOSETRON HYDROCHLORIDE

HEMOFARM

- * HEMOFARM AD
A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE

HERCON LABS

- * HERCON LABORATORIES CORP
NITROGLYCERIN, NITROGLYCERIN

HERITAGE PHARMS INC

- * HERITAGE PHARMACEUTICALS INC
ACETAZOLAMIDE, ACETAZOLAMIDE
CARISOPRODOL AND ASPIRIN, ASPIRIN
DOXYCYCLINE, DOXYCYCLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HERITAGE PHARMACEUTICALS INC
 - GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 - GLYBURIDE, GLYBURIDE
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - INDOMETHACIN, INDOMETHACIN
 - KETOPROFEN, KETOPROFEN
 - LEFLUNOMIDE, LEFLUNOMIDE
 - NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 - PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE

HETERO DRUGS

- * HETERO DRUGS LTD
 - METHOCARBAMOL, METHOCARBAMOL
 - STAVUDINE, STAVUDINE
 - TORSEMIDE, TORSEMIDE

HETERO DRUGS LTD

- * HETERO DRUGS LTD UNIT III
 - FINASTERIDE, FINASTERIDE
 - LEVETIRACETAM, LEVETIRACETAM
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - ZIDOVUDINE, ZIDOVUDINE

HETERO LABS UNIT III

- * HETERO LABS LTD UNIT III
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - INDOMETHACIN, INDOMETHACIN

HEYL CHEMISCH

- * HEYL CHEMISCH PHARMAZEUTISCHE FABRIK
 - RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

HH AND P

- * HH AND P LLC
 - CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 - FLUNISOLIDE, FLUNISOLIDE

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
 - ERYTHRO-STATIN, ERYTHROMYCIN
 - 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)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HI TECH PHARMACAL CO INC
 - MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 - NYSTATIN, NYSTATIN
 - OFLOXACIN, OFLOXACIN
 - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 - PREDNISOLONE, PREDNISOLONE
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE
 - 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

HIGH TECH PHARMA

- * HIGH TECHNOLOGY PHARMACAL CO INC
 - VALPROIC ACID, VALPROIC ACID

HIKMA

- * HIKMA FARMACEUTICA LDA
 - CEFOTAXIME, CEFOTAXIME SODIUM
- * HIKMA PHARMACEUTICALS
 - AMOXICILLIN, AMOXICILLIN
 - CARVEDILOL, CARVEDILOL
 - CEFACLOR, CEFACLOR
 - CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 - CEPHALEXIN, CEPHALEXIN
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - GABAPENTIN, GABAPENTIN
 - GLYBURIDE (MICRONIZED), GLYBURIDE
 - NAPROXEN SODIUM, NAPROXEN SODIUM
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

HIKMA FARMACEUTICA

- * HIKMA FARMACEUTICA (PORTUGAL) SA
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 - CEFOXITIN, CEFOXITIN SODIUM
 - CEFTRIAZONE, CEFTRIAZONE SODIUM
 - CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
 - CIPROFLOXACIN, CIPROFLOXACIN
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - ENALAPRILAT, ENALAPRILAT
 - FLUMAZENIL, FLUMAZENIL
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - GLYCOPYRRROLATE, GLYCOPYRRROLATE
 - GRANISTERON HYDROCHLORIDE, GRANISTERON 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
 - VALPROATE SODIUM, VALPROATE SODIUM
- * HIKMA FARMACEUTICA PORTUGAL LDA
 - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 - CEFUXIME SODIUM, CEFUXIME SODIUM
 - FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HIKMA FARMACEUTICA SA
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE

HIKMA PHARMS

- * HIKMA PHARMACEUTICALS
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
CEPHALEXIN, CEPHALEXIN
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
GEMFIBROZIL, GEMFIBROZIL
LAMOTRIGINE, LAMOTRIGINE
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

HILL DERMAC

- * HILL DERMACEUTICALS INC
DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
DERMOTIC, FLUOCINOLONE ACETONIDE

HISAMITSU

- * HISAMITSU PHARMACEUTICAL CO INC
SALONPAS, MENTHOL (OTC)

HITECH PHARMA

- * HITECH PHARMACAL CORP
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE

HI-TECH PHARMA CO

- * HI-TECH PHARMACAL CO INC
TUSSICAPS, CHLORPHENIRAMINE POLISTIREX

HOFFMANN LA ROCHE

- * HOFFMANN LA ROCHE INC
INVIRASE, SAQUINAVIR MESYLATE
ROCEPHIN, CEFTRIAZONE SODIUM
ROMAZICON, FLUMAZENIL
XELODA, CAPECITABINE
XENICAL, ORLISTAT
ZELBORAF, VEMURAFENIB

HOPE PHARMS

- * HOPE PHARMACEUTICALS
NITHIODETE, SODIUM NITRITE

HORIZON PHARMA

- * HORIZON PHARMA INC
DUEXIS, FAMOTIDINE

HOSPIRA

- * HOSPIRA INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ACETYLCYSTEINE, ACETYLCYSTEINE
A-HYDROCORT, HYDROCORTISONE SODIUM SUCCINATE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE
AMIDATE, ETOMIDATE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOPHYLLINE, AMINOPHYLLINE
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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HOSPIRA INC
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
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
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
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
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPROFLOXACIN, CIPROFLOXACIN
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CORLOPAM, FENOLDOPAM MESYLATE
CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
CYTARABINE, CYTARABINE
DACARBAZINE, DACARBAZINE
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
DEMEROL, MEPERIDINE HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 25%, 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 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
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
DIAZEPAM, DIAZEPAM
DIGOXIN, DIGOXIN
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%, 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
ERYC, ERYTHROMYCIN
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
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FOSCARNET SODIUM, FOSCARNET SODIUM
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
FUROSEMIDE, FUROSEMIDE
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
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 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
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
ISOFLURANE, ISOFLURANE
ISUPREL, ISOPROTERENOL HYDROCHLORIDE
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
LEVOPHED, NOREPINEPHRINE BITARTRATE
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.8% IN 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 IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LIPOSYN II 10%, SAFFLOWER OIL
LIPOSYN II 20%, SAFFLOWER OIL
LIPOSYN III 10%, SOYBEAN OIL
LIPOSYN III 20%, SOYBEAN OIL
LIPOSYN III 30%, SOYBEAN OIL
LORAZEPAM, LORAZEPAM
LTA II KIT, LIDOCAINE HYDROCHLORIDE
M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
M.V.I. ADULT, ASCORBIC ACID
M.V.I. PEDIATRIC, ASCORBIC ACID
M.V.I.-12 (WITHOUT VITAMIN K), ASCORBIC ACID
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE, MAGNESIUM SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

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
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
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE 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
NITROGLYCERIN, NITROGLYCERIN
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
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PACLITAXEL, PACLITAXEL
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
POTASSIUM ACETATE IN PLASTIC CONTAINER, POTASSIUM ACETATE
POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 10EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 10EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 10EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 10EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 10EQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 15EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 15EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 15EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 15EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20EQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 20EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20EQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ******* HOSPIRA INC**

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
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 SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
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, CALCIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
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 SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPOFOL, PROPOFOL
QUELICIN PRESERVATIVE FREE, SUCCINYLCHOLINE CHLORIDE
QUELICIN, SUCCINYLCHOLINE CHLORIDE
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SODIUM ACETATE IN PLASTIC CONTAINER, SODIUM ACETATE ANHYDROUS
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
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
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
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** H ****

- * HOSPIRA INC
 - 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
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 - NITROPRESS, SODIUM NITROPRUSSIDE
 - ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 - VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

HOSPIRA INC

- * HOSPIRA INC
 - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 - CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
 - CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
 - CEFOXITIN, CEFOXITIN SODIUM
 - CEFTRIAXONE, CEFTRIAXONE SODIUM
 - CEFUROXIME SODIUM, CEFUROXIME SODIUM
 - DOCETAXEL, DOCETAXEL
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - GEMCITABINE, GEMCITABINE HYDROCHLORIDE
 - HEPARIN SODIUM, HEPARIN SODIUM
 - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 - IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
 - MAXIPIME, CEFEPIME HYDROCHLORIDE
 - MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
 - MEROPENEM, MEROPENEM
 - MORPHINE SULFATE, MORPHINE SULFATE
 - NIPENT, PENTOSTATIN
 - OXALIPLATIN, OXALIPLATIN
 - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
 - THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 - TOPOTECAN, TOPOTECAN HYDROCHLORIDE
 - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

HOSPIRA WORLDWIDE

- * HOSPIRA WORLDWIDE PTY
 - OXALIPLATIN, OXALIPLATIN

HQ SPECIALITY PHARMA

- * HQ SPECIALITY PHARMA LLC
 - LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM

HUAHAI US INC

- * HUAHAI US INC
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE

IBI

- * ISTITUTO BIOCHIMICO ITALIANO SPA
 - AMPICILLIN SODIUM, AMPICILLIN SODIUM
 - NAFCILLIN SODIUM, NAFCILLIN SODIUM

IDENTI PHARMS INC

- * IDENTI PHARMACEUTICALS INC
 - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

IMPAX LABS

- * IMPAX LABORATORIES INC
 - ACARBOSE, ACARBOSE
 - ALPRAZOLAM, ALPRAZOLAM
 - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 - BACLOFEN, BACLOFEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

- * IMPAX LABORATORIES INC
 - BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
 - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - CARBIDOPA AND LEVODOPA, CARBIDOPA
 - CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
 - DANTROLENE SODIUM, DANTROLENE SODIUM
 - DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 - DIGOXIN, DIGOXIN
 - DIPYRIDAMOLE, DIPYRIDAMOLE
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - FENOFIBRATE (MICRONIZED), FENOFIBRATE
 - FENOFIBRATE, FENOFIBRATE
 - FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 - LETROZOLE, LETROZOLE
 - LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 - LORATADINE, LORATADINE (OTC)
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METHYLTESTOSTERONE, METHYLTESTOSTERONE
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE
 - OMEPRAZOLE, OMEPRAZOLE
 - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 - PENTOXIFYLLINE, PENTOXIFYLLINE
 - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 - PRIMIDONE, PRIMIDONE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
 - RILUZOLE, RILUZOLE
 - RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
 - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 - TERBUTALINE SULFATE, TERBUTALINE SULFATE
 - TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

IMPAX LABS INC

- * IMPAX LABORATORIES INC
 - CABERGOLINE, CABERGOLINE
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DOXYCYCLINE, DOXYCYCLINE
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

IMPAX PHARMS

- * IMPAX PHARMACEUTICALS
 - FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 - FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 - GEMFIBROZIL, GEMFIBROZIL
 - MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

INALCO

- * INALCO SPA
 - LACTULOSE, LACTULOSE

INCYTE CORP

- * INCYTE CORP
 - JAKAFI, RUXOLITINIB PHOSPHATE

INDICUS PHARMA

- * INDICUS PHARMA LLC
 - LETROZOLE, LETROZOLE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******INNOPHARMA LLC**

- * INNOPHARMA LLC
 - LEVETIRACETAM, LEVETIRACETAM
 - OLANZAPINE, OLANZAPINE

INO

- * INO THERAPEUTICS INC
 - INOMAX, NITRIC OXIDE

INSIGHT PHARMS

- * INSIGHT PHARMACEUTICALS CORP
 - 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)
 - MONISTAT-3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 - NIX, PERMETHRIN (OTC)

INSPIRE

- * INSPIRE PHARMACEUTICALS INC
 - AZASITE, AZITHROMYCIN

INST BIOCHEM

- * INSTITUT BIOCHEMIQUE SA
 - FLECTOR, DICLOFENAC EPOLAMINE

INST BIOCHIMIQUE

- * INSTITUTE BIOCHIMIQUE SA (IBSA)
 - TIROSINT, LEVOTHYROXINE SODIUM

INSTITUT BIOCHIMIQUE

- * INSTITUT BIOCHIMIQUE SA IBSA
 - TIROSINT, LEVOTHYROXINE SODIUM

INSTITUTO BIOCHEMICO

- * INSTITUTO BIOCHEMICO ITALIANO SPA
 - AMPICILLIN SODIUM, AMPICILLIN SODIUM
 - NAFCILLIN SODIUM, NAFCILLIN SODIUM
 - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

INSTITUTO BIOCHIMICO

- * INSTITUTO BIOCHIMICO ITALIANO SPA
 - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM

INSYS THERAP

- * INSYS THERAPEUTICS INC
 - DRONABINOL, DRONABINOL

INTELGENX CORP

- * INTELGENX CORP
 - FORFIVO XL, BUPROPION HYDROCHLORIDE

INTENDIS

- * INTENDIS INC
 - DESONATE, DESONIDE
 - FINACEA, AZELAIC ACID

INTERGEL PHARM

- * INTERGEL PHARMACEUTICAL INC
 - NIFEDIPINE, NIFEDIPINE

INTL MEDICATED

- * INTERNATIONAL MEDICATED SYSTEMS LTD
 - BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - MILRINONE LACTATE, MILRINONE LACTATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ******INTL MEDICATION**

- * INTERNATIONAL MEDICATION SYSTEM
 - AMINOPHYLLINE, AMINOPHYLLINE
 - DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 - FUROSEMIDE, FUROSEMIDE
 - ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 - LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 - LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - MANNITOL 25%, MANNITOL
 - MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
 - NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
 - PHYTONADIONE, PHYTONADIONE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
- * INTERNATIONAL MEDICATION SYSTEMS LTD
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE

INTL MEDICATION SYS

- * INTERNATIONAL MEDICATION SYSTEMS LTD
 - LORAZEPAM, LORAZEPAM

INVAGEN PHARMS

- * INVAGEN PHARMACEUTICALS INC
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 - FOLIC ACID, FOLIC ACID
 - FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 - FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 - GABAPENTIN, GABAPENTIN
 - GEMFIBROZIL, GEMFIBROZIL
 - GLIMEPIRIDE, GLIMEPIRIDE
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - LEVETIRACETAM, LEVETIRACETAM
 - MEPROBAMATE, MEPROBAMATE
 - NABUMETONE, NABUMETONE
 - NAPROXEN, NAPROXEN
 - QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 - RAMIPRIL, RAMIPRIL
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - TOPIRAMATE, TOPIRAMATE
 - TRANDOLAPRIL, TRANDOLAPRIL
 - WARFARIN SODIUM, WARFARIN SODIUM
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 - ZONISAMIDE, ZONISAMIDE

INWOOD LABS

- * INWOOD LABORATORIES INC SUB FOREST LABORATORIES INC
 - THEOPHYLLINE, THEOPHYLLINE

IPCA LABS LTD

- * IPCA LABORATORIES LTD
 - ALLOPURINOL, ALLOPURINOL
 - ATENOLOL, ATENOLOL
 - CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 - FUROSEMIDE, FUROSEMIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

- * IPCA LABORATORIES LTD
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

IPR

- * IPR PHARMACEUTICALS INC
 - ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 - ATENOLOL, ATENOLOL
 - CRESTOR, ROSUVASTATIN CALCIUM
 - ZOMIG, ZOLMITRIPTAN

IPR PHARMS INC

- * IPR PHARMACEUTICALS INC
 - VANDETANIB, VANDETANIB

IPSEN PHARMS

- * IPSEN PHARMACEUTICALS INC
 - SOMATULINE DEPOT, LANREOTIDE ACETATE

IROKO PHARMS

- * IROKO PHARMACEUTICALS LLC
 - INDOCIN, INDOMETHACIN

ISO TEX

- * ISO TEX DIAGNOSTICS INC
 - JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 - MEGATOPE, ALBUMIN IODINATED I-131 SERUM

ISTA PHARMS

- * ISTA PHARMACEUTICALS
 - BEPREVE, BEPOTASTINE BESILATE
 - ISTALOL, TIMOLOL MALEATE
 - VITRASE, HYALURONIDASE

ISTA PHARMS INC

- * ISTA PHARMACEUTICALS INC
 - BROMDAY, BROMFENAC SODIUM

ISTITUTO BIOCHIMICO

- * ISTITUTO BIOCHIMICO ITALIANO GIOVANNI LORENZINI
 - PIPERACILLIN, PIPERACILLIN SODIUM

IVAX PHARMS

- * IVAX PHARMACEUTICALS INC
 - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE

IVAX SUB TEVA PHARMS

- * IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA
 - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 - BACLOFEN, BACLOFEN
 - BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 - BUMETANIDE, BUMETANIDE
 - CABERGOLINE, CABERGOLINE
 - CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 - CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
 - CIMETIDINE, CIMETIDINE
 - CIMETIDINE, CIMETIDINE (OTC)
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLOZAPINE, CLOZAPINE
 - CYCLOSPORINE, CYCLOSPORINE
 - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 - DIAZEPAM, DIAZEPAM
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DOXYCYCLINE, DOXYCYCLINE
 - FAMOTIDINE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** I ****

* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUTAMIDE, FLUTAMIDE
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GRISEOFULVIN, GRISEOFULVIN, MICROCRYSTALLINE
 GUANABENZ ACETATE, GUANABENZ ACETATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 INDAPAMIDE, INDAPAMIDE
 INDOMETHACIN, INDOMETHACIN
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LORAZEPAM, LORAZEPAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHYLDOPA, METHYLDOPA
 MISOPROSTOL, MISOPROSTOL
 NADOLOL, NADOLOL
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OXaprozin, OXaprozin
 OXAZEPAM, OXAZEPAM
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PROBENECID, PROBENECID
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 SIMVASTATIN, SIMVASTATIN
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

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
 DITROPAN, OXYBUTYNIN CHLORIDE
 DORIBAX, DORIPENEM
 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, PALIPERIDONE
 LEUSTATIN, CLADRIBINE
 LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
 LEVAQUIN, LEVOFLOXACIN
 MICRONOR, NORETHINDRONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ****

- * JANSEN PHARMACEUTICALS INC
MODICON 28, ETHINYL ESTRADIOL
NIZORAL, KETOCONAZOLE
NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
NUCYNTA, TAPENTADOL HYDROCHLORIDE
ORTHO CYCLEN-28, ETHINYL ESTRADIOL
ORTHO EVRA, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
ORTHO-CEPT, DESOGESTREL
ORTHO-NOVUM 1/35-28, ETHINYL ESTRADIOL
ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
PANCREAZE, LIPASE
RAZADYNE ER, GALANTAMINE HYDROBROMIDE
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
TYLOX, ACETAMINOPHEN
ULTRACET, ACETAMINOPHEN
ULTRAM, TRAMADOL HYDROCHLORIDE
XARELTO, RIVAROXABAN

JANSEN R AND D

- * JANSEN RESEARCH AND DEVELOPMENT LLC
FLEXERIL, CYCLOBENZAPRINE HYDROCHLORIDE
PARAFON FORTE DSC, CHLORZOXAZONE

JAZZ

- * JAZZ PHARMACEUTICALS
LUVOX CR, FLUVOXAMINE MALEATE
XYREM, SODIUM OXYBATE

JHP PHARMS

- * JHP PHARMACEUTICALS LLC
BREVITAL SODIUM, METHOHEXITAL SODIUM
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
DANTRIUM, DANTROLENE SODIUM
DELESTROGEN, ESTRADIOL VALERATE
KETALAR, KETAMINE HYDROCHLORIDE
PITOCIN, OXYTOCIN
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
TRIOSTAT, LIOTHYRONINE SODIUM

JIANGSU HENGRI MED

- * JIANGSU HENGRI MEDICINE CO LTD
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

JOHN O BUTLER CO

- * JOHN O BUTLER CO
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE

JOHNSON AND JOHNSON

- * JOHNSON AND JOHNSON GROUP CONSUMER COMPANIES
MEN'S ROGAINE, MINOXIDIL (OTC)
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISINE-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** J ******JUBILANT CADISTA**

- * JUBILANT CADISTA PHARMACEUTICALS INC
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
PREDNISONE, PREDNISONE
PROCOMP, PROCHLORPERAZINE MALEATE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

JUBILANT LIFE

- * JUBILANT LIFE SCIENCES LTD
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
LAMOTRIGINE, LAMOTRIGINE

JUBILANT ORGANOSYS

- * JUBILANT ORGANOSYS LTD
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
RISPERIDONE, RISPERIDONE

KEY PHARMS

- * KEY PHARMACEUTICALS INC SUB SCHERING PLOUGH CORP
NITRO-DUR, NITROGLYCERIN

KIEL

- * KIEL LABORATORIES INC
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE

KING PHARMS

- * KING PHARMACEUTICALS INC
ALTACE, RAMIPRIL
AVINZA, MORPHINE SULFATE
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
CORGARD, NADOLOL
CORZIDE, BENDROFLUMETHIAZIDE
CYTOMEL, LIOTHYRONINE SODIUM
INTAL, CROMOLYN SODIUM
LEVOXYL, LEVOTHYROXINE SODIUM
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
SILVADENE, SILVER SULFADIAZINE
SKELAXIN, METAXALONE
SYNERCID, DALFOPRISTIN
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
- * KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT INC SUB KING PHARMACEUTICALS INC
SONATA, ZALEPLON
TAPAZOLE, METHIMAZOLE
TUSSIGON, HOMATROPINE METHYLBROMIDE

KING PHARMS R AND D

- * KING PHARMACEUTICALS RESEARCH DEVELOPMENT INC
OXECTA, OXYCODONE HYDROCHLORIDE

KOWA CO

- * KOWA CO LTD
LIVALO, PITAVASTATIN CALCIUM

KREMERS URBAN PHARMS

- * KREMERS URBAN PHARMACEUTICALS INC
GLYCOLAX, POLYETHYLENE GLYCOL 3350
GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
NITROGLYCERIN, NITROGLYCERIN
OMEPRAZOLE, OMEPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** K ******KUDCO IRELAND**

- * KUDCO IRELAND LTD
 - ANASTROZOLE, ANASTROZOLE
 - BICALUTAMIDE, BICALUTAMIDE
 - LETROZOLE, LETROZOLE
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

KV PHARM

- * KV PHARMACEUTICAL CO
 - CLINDESSE, CLINDAMYCIN PHOSPHATE
 - EVAMIST, ESTRADIOL
 - GYNAZOLE-1, BUTOCONAZOLE NITRATE
 - MAKENA, HYDROXYPROGESTERONE CAPROATE

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

LAB HRA PHARMA

- * LABORATOIRE HRA PHARMA
 - ELLA, ULIPRISTAL ACETATE

LANDELA PHARM

- * LANDELA PHARMACEUTICAL
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - 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
 - LANORINAL, ASPIRIN
 - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 - PRIMIDONE, PRIMIDONE
 - PROBALAN, PROBENECID
 - SERPALAN, SERPINE
- * LANNETT HOLDINGS INC
 - BACLOFEN, BACLOFEN
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 - DANAZOL, DANAZOL
 - DIPYRIDAMOLE, DIPYRIDAMOLE
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DOXYCYCLINE, DOXYCYCLINE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 - RIFAMPIN, RIFAMPIN
 - TERBUTALINE SULFATE, TERBUTALINE SULFATE
 - URSODIOL, URSDIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ******LANNETT HOLDINGS INC**

- * LANNETT HOLDINGS INC
 - DIETHYLPROMION HYDROCHLORIDE, DIETHYLPROMION HYDROCHLORIDE
 - LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 - METHOCARBAMOL, METHOCARBAMOL
 - MORPHINE SULFATE, MORPHINE SULFATE
 - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 - PHENTERMINE RESIN COMPLEX, PHENTERMINE RESIN COMPLEX
 - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

LANTHEUS MEDCL

- * LANTHEUS MEDICAL IMAGING INC
 - ABLAVAR, GADOFOSVESET TRISODIUM
 - 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

LAVIPHARM LABS

- * LAVIPHARM LABORATORIES INC
 - FENTANYL-100, FENTANYL
 - FENTANYL-25, FENTANYL
 - FENTANYL-50, FENTANYL
 - FENTANYL-75, FENTANYL

LEHIGH VALLEY

- * LEHIGH VALLEY TECHNOLOGIES INC
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

LEK PHARMS

- * LEK PHARMACEUTICALS D D
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 - ENALAPRIL MALEATE, ENALAPRIL MALEATE
 - LISINOPRIL, LISINOPRIL
 - OMEPRAZOLE, OMEPRAZOLE

LEK PHARMS DD

- * LEK PHARMACEUTICALS DD
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

LEO PHARM

- * LEO PHARMACEUTICAL PRODUCTS LTD
 - DOVONEX, CALCIPOTRIENE
 - TACLONEX, BETAMETHASONE DIPROPIONATE

LEO PHARM PRODS

- * LEO PHARMACEUTICAL PRODUCTS LTD
 - TACLONEX SCALP, BETAMETHASONE DIPROPIONATE

LEO PHARMA AS

- * LEO PHARMA AS
 - INNOHEP, TINzaparin SODIUM

LG LIFE

- * LG LIFE SCIENCES LTD
 - VALTROPIN, SOMATROPIN RECOMBINANT

LILLY

- * ELI LILLY AND CO
 - ALIMTA, PEMETREXED DISODIUM
 - CIALIS, Tadalafil
 - CYMBALTA, DULOXETINE HYDROCHLORIDE
 - EVISTA, RALOXIFENE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

- * ELI LILLY AND CO
 - 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 PEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 - HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
 - HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 - HUMALOG MIX 75/25 PEN, INSULIN LISPRO PROTAMINE RECOMBINANT
 - HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
 - HUMALOG PEN, INSULIN LISPRO 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 PEN, INSULIN RECOMBINANT HUMAN (OTC)
 - HUMULIN R, INSULIN RECOMBINANT HUMAN
 - HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
 - PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
 - QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 - STRATTERA, ATOMOXETINE HYDROCHLORIDE
 - SYMBYAX, FLUOXETINE HYDROCHLORIDE
 - ZYPREXA ZYDIS, OLANZAPINE
 - ZYPREXA, OLANZAPINE
- * LILLY RESEARCH LABORATORIES DIV ELI LILLY AND CO
 - PROZAC, FLUOXETINE HYDROCHLORIDE
 - SARAFEM, FLUOXETINE HYDROCHLORIDE

LLOYD

- * LLOYD INC
 - LEVOOTHROID, LEVOTHYROXINE SODIUM

LNK

- * LNK INTERNATIONAL INC
 - DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

LOREAL USA

- * LOREAL USA PRODUCTS INC
 - ANTHELIOS 20, AVOBENZONE (OTC)
 - ANTHELIOS 40, AVOBENZONE (OTC)
 - ANTHELIOS SX, AVOBENZONE (OTC)
 - CAPITAL SOLEIL 15, AVOBENZONE (OTC)

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
 - CALCITRIOL, CALCITRIOL
 - CEFTRIAXONE, CEFTRIAXONE 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

- * LUITPOLD PHARMACEUTICALS INC
 - DROPERIDOL, DROPERIDOL
 - ESTRADIOL VALERATE, ESTRADIOL VALERATE
 - ETOMIDATE, ETOMIDATE
 - FOMEPIZOLE, FOMEPIZOLE
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - FUROSEMIDE, FUROSEMIDE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - GRANisetron hydrochloride, GRANisetron hydrochloride
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - LATANOPROST, LATANOPROST
 - LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 - LEVOCARNITINE, LEVOCARNITINE
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - MANNITOL 25%, MANNITOL
 - METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
 - METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
 - NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
 - NITROGLYCERIN, NITROGLYCERIN
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 - PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - PROGESTERONE, PROGESTERONE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - SPRIX, KETOROLAC TROMETHAMINE
 - TORSEMIDE, TORSEMIDE
 - TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 - TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
 - VENOFER, IRON SUCROSE
 - ZIDOVUDINE, ZIDOVUDINE

LUNDBECK INC

- * LUNDBECK INC
 - CHEMET, SUCCIMER
 - COGETINT, BENZTROPINE MESYLATE
 - COSMEGEN, DACTINOMYCIN
 - DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
 - DIURIL, CHLOROTHIAZIDE SODIUM
 - INDOCIN, INDOMETHACIN SODIUM
 - MUSTARGEN, MECHLORETHAMINE HYDROCHLORIDE
 - NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
 - NEOPROFEN, IBUPROFEN LYSINE
 - ONFI, CLOBAZAM
 - PEGANONE, ETHOTOIN
 - SABRIL, VIGABATRIN
 - TRANXENE, CLORAZEPATE DIPOTASSIUM

LUPIN

- * LUPIN LTD
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** L ****

- * LUPIN LTD
 - 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
 - TRANDOLAPRIL, TRANDOLAPRIL

LUPIN ATLANTIS

- * LUPIN ATLANTIS HOLDINGS SA
 - ANTARA (MICRONIZED), FENOFLIBRATE

LUPIN LTD

- * LUPIN LIMITED
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
- * LUPIN LTD
 - FAMOTIDINE, FAMOTIDINE
 - FENOFLIBRATE, FENOFLIBRATE
 - IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 - IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 - LAMOTRIGINE, LAMOTRIGINE
 - LEVETIRACETAM, LEVETIRACETAM
 - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 - MEFENAMIC ACID, MEFENAMIC ACID
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - NABUMETONE, NABUMETONE
 - NORETHINDRONE, NORETHINDRONE
 - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 - SUPRAX, CEFIXIME
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

LUPIN PHARMS

- * LUPIN PHARMACEUTICALS INC
 - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 - DESLORATADINE, DESLORATADINE
 - MELOXICAM, MELOXICAM
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - SUPRAX, CEFIXIME

LYNE

- * LYNE LABORATORIES INC
 - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 - DEXAMETHASONE, DEXAMETHASONE
 - HYDROCORTISONE, HYDROCORTISONE
 - LEVOCARNITINE, LEVOCARNITINE
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

LYNROSE LABS

- * LYNROSE LABS LLC
 - CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE

MALLINCKRODT

- * MALLINCKRODT CHEMICAL INC
 - ANEXSIA 7.5/650, ACETAMINOPHEN
 - ANEXSIA, ACETAMINOPHEN
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - METHADOSE, METHADONE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MALLINCKRODT CHEMICAL INC
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
- * MALLINCKRODT INC
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANEXSIA, ACETAMINOPHEN
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
FENTANYL CITRATE, FENTANYL CITRATE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
OPTIMARK IN PLASTIC CONTAINER, GADOVERSETAMIDE
OPTIMARK, GADOVERSETAMIDE
OXYCET, ACETAMINOPHEN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PENNSAID, DICLOFENAC SODIUM
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
TECHNETIUM TC-99 SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
- * MALLINCKRODT MEDICAL INC
CONRAY 30, IOTHALAMATE MEGLUMINE
CONRAY 43, IOTHALAMATE MEGLUMINE
CONRAY, IOTHALAMATE MEGLUMINE
CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67
INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE
MD-76R, DIATRIZOATE MEGLUMINE
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
OPTIRAY 240, IOVERSOL
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL
OPTIRAY 350, IOVERSOL
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
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201
ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

MALLINCKRODT INC

- * MALLINCKRODT INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
EXALGO, HYDROMORPHONE HYDROCHLORIDE
FENTANYL-100, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
RESTORIL, TEMAZEPAM
TOFRANIL, IMIPRAMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MALLINCKRODT INC
TOFRANIL-PM, IMIPRAMINE PAMOATE

MALLINCKRODT LLC

* MALLINCKRODT LLC
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
PAMELOR, NORTRIPTYLINE HYDROCHLORIDE

MARATHON PHARMS

* MARATHON PHARMACEUTICALS LLC
SECONAL SODIUM, SECOBARBITAL SODIUM

MARKSANS PHARMA

* MARKSANS PHARMA LTD
GABAPENTIN, GABAPENTIN
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NAPROXEN, NAPROXEN

MARSAM PHARMS LLC

* MARSAM PHARMACEUTICALS LLC
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

MATRIX LABS LTD

* MATRIX LABORATORIES LTD
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
BACLOFEN, BACLOFEN
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
DIDANOSINE, DIDANOSINE
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
LANSOPRAZOLE, LANSOPRAZOLE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
NABUMETONE, NABUMETONE
NIFEDIPINE, NIFEDIPINE
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
RAMIPRIL, RAMIPRIL
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
SIMVASTATIN, SIMVASTATIN
STAVUDINE, STAVUDINE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VORICONAZOLE, VORICONAZOLE
ZIDOVUDINE, ZIDOVUDINE

MAYNE PHARMA

* MAYNE PHARMA INTERNATIONAL PTY LTD
DORYX, DOXYCYCLINE HYCLATE

MCNEIL

* MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC
CHILDREN'S MOTRIN, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
IMODIUM A-D EZ CHEWS, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)
MOTRIN IB, IBUPROFEN (OTC)
MOTRIN MIGRAINE PAIN, IBUPROFEN (OTC)
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

MCNEIL CONS

* MCNEIL CONSUMER HEALTHCARE
BENADRYL PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
BENADRYL, DIPHENHYDRAMINE HYDROCHLORIDE
CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
CHILDREN'S MOTRIN, IBUPROFEN (OTC)
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MCNEIL CONSUMER HEALTHCARE
 - IMODIUM, LOPERAMIDE HYDROCHLORIDE
 - JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
 - NIZORAL A-D, KETOCONAZOLE (OTC)
 - SINE-AID IB, IBUPROFEN (OTC)
 - SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 - SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 - TYLENOL (CAPLET), ACETAMINOPHEN (OTC)
 - TYLENOL (GELTAB), ACETAMINOPHEN (OTC)

MCNEIL CONSUMER

- * MCNEIL CONSUMER HEALTHCARE DIV MCNEIL PPC INC
 - CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - ZYRTEC, CETIRIZINE HYDROCHLORIDE

MEAD JOHNSON

- * MEAD JOHNSON AND CO
 - CAFCIT, CAFFEINE CITRATE

MEDA PHARMS

- * MEDA PHARMACEUTICALS
 - EDLUAR, ZOLPIDEM TARTRATE
- * MEDA PHARMACEUTICALS INC
 - ANADROL-50, OXYMETHOLONE
 - ASTELIN, AZELASTINE HYDROCHLORIDE
 - ASTEPRO, AZELASTINE HYDROCHLORIDE
 - BUTISOL SODIUM, BUTABARBITAL SODIUM
 - CESAMET, NABILONE
 - COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 - COLYTE, POLYETHYLENE GLYCOL 3350
 - COLYTE-FLAVORED, POLYETHYLENE GLYCOL 3350
 - CORTIFOAM, HYDROCORTISONE ACETATE
 - DEMADEX, TORSEMIDE
 - DEPEN, PENICILLAMINE
 - DIPENTUM, OLSALAZINE SODIUM
 - EPIFOAM, HYDROCORTISONE ACETATE
 - FELBATOL, FELBAMATE
 - LUFYLLIN, DYPHYLLINE
 - MUSE, ALPROSTADIL
 - ONSOLIS, FENTANYL CITRATE
 - OPTIVAR, AZELASTINE HYDROCHLORIDE
 - PROCTOFOAM HC, HYDROCORTISONE ACETATE
 - ROWASA, MESALAMINE
 - SFROWASA, MESALAMINE
 - TRILYTE, POLYETHYLENE GLYCOL 3350
- * MEDA PHARMACEUTICALS MEDA PHARMACEUTICALS INC
 - SOMA COMPOUND W/ CODEINE, ASPIRIN
 - SOMA COMPOUND, ASPIRIN
 - SOMA, CARISOPRODOL

MEDICINES CO

- * THE MEDICINES CO
 - ANGIOMAX, BIVALIRUDIN
 - CLEVIPREX, CLEVIDIPIINE BUTYRATE

MEDICIS

- * MEDICIS PHARMACEUTICAL CORP
 - ALDARA, IMIQUIMOD
 - BUPHENYL, SODIUM PHENYLBUTYRATE
 - CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DYNACIN, MINOCYCLINE HYDROCHLORIDE
 - ESTRASORB, ESTRADIOL HEMIHYDRATE
 - LIDEX, FLUOCINONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

* MEDICIS PHARMACEUTICAL CORP
LIDEX-E, FLUOCINONIDE
LOPROX, CICLOPIROX
MAXAIR, PIRBUTEROL ACETATE
METROGEL-VAGINAL, METRONIDAZOLE
NORFLEX, ORPHENADRINE CITRATE
NORGESIC FORTE, ASPIRIN
NORGESIC, ASPIRIN
SOLODYN, MINOCYCLINE HYDROCHLORIDE
SYNACORT, HYDROCORTISONE
TAMBOCOR, FLECAINIDE ACETATE
VANOS, FLUOCINONIDE
ZIANA, CLINDAMYCIN PHOSPHATE
ZYCLARA, IMIQUIMOD

MEDICURE

* MEDICURE INTERNATIONAL INC
AGGRASTAT, TIROFIBAN HYDROCHLORIDE

MEDIGENE AG

* MEDIGENE AG
VEREGEN, SINECATECHINS

MEDIMETRIKS PHARMS

* MEDIMETRIKS PHARMACEUTICALS INC
SYNALAR, FLUOCINOLONE ACETONIDE

MEDIMMUNE

* MEDIMMUNE
ETHYOL, AMIFOSTINE

MEDTRONIC

* MEDTRONIC INC
LIORESAL, BACLOFEN

MERCK

* MERCK AND CO INC
AMINOHIPPURATE SODIUM, AMINOHIPPURATE SODIUM
CANCIDAS, CASPOFUNGIN ACETATE
EMEND, APREPITANT
FOSAMAX PLUS D, ALENDRONATE SODIUM
FOSAMAX, ALENDRONATE SODIUM
HYZAAR, HYDROCHLOROTHIAZIDE
INVANZ, ERTPENEM SODIUM
JANUMET, METFORMIN HYDROCHLORIDE
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
PRIMAXIN, CILASTATIN SODIUM
PROSCAR, FINASTERIDE
SINGULAIR, MONTELUKAST SODIUM
STROMECTOL, IVERMECTIN
ZOLINZA, VORINOSTAT
* MERCK RESEARCH LABORATORIES DIV MERCK CO INC
CLINORIL, SULINDAC
COSOPT, DORZOLAMIDE HYDROCHLORIDE
COZAAR, LOSARTAN POTASSIUM
MEVACOR, LOVASTATIN
NOROXIN, NORFLOXACIN
PEPCID, FAMOTIDINE
PRINIVIL, LISINOPRIL
PRINZIDE, HYDROCHLOROTHIAZIDE
PROPECIA, FINASTERIDE
SINGULAIR, MONTELUKAST SODIUM
TRUSOPT, DORZOLAMIDE HYDROCHLORIDE
ZOCOR, SIMVASTATIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MERCK AND CO INC**

- * MERCK AND CO INC
 - EMEND, FOSAPREPITANT DIMEGLUMINE
 - FOSAMAX, ALENDRONATE SODIUM

MERCK CO INC

- * MERCK CO INC
 - JANUVIA, SITAGLIPTIN PHOSPHATE

MERCK KGAA

- * MERCK KGAA
 - LEVOHYROXINE SODIUM, LEVOTHYROXINE SODIUM

MERCK SANTE SAS

- * MERCK SANTE SAS
 - CYANOKIT, HYDROXOCOBALAMIN

MERCK SHARP DOHME

- * MERCK SHARP AND DOHME CORP
 - CRIVAN, INDINAVIR SULFATE
 - ISENTRESS, Raltegravir Potassium
 - JUVISYNC, SIMVASTATIN
 - PEPCID AC (GELTAB), FAMOTIDINE (OTC)
 - PEPCID AC, FAMOTIDINE (OTC)
 - PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
 - SINEMET CR, CARBIDOPA
 - SINEMET, CARBIDOPA

MERIDIAN MEDCL

- * MERIDIAN MEDICAL TECHNOLOGIES INC
 - DUODOTE, ATROPINE
- * MERIDIAN MEDICAL TECHNOLOGIES INC SUB KING PHARMACEUTICALS INC
 - ALSUMA, SUMATRIPTAN SUCCINATE

MERIDIAN MEDCL TECHN

- * MERIDIAN MEDICAL TECHNOLOGIES INC
 - ATROOPEN, ATROPINE
 - EPIPEN JR., EPINEPHRINE
 - EPIPEN, EPINEPHRINE
 - LIDOPEN, LIDOCAINE HYDROCHLORIDE
 - MORPHINE SULFATE, MORPHINE SULFATE
 - PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE

MERRO PHARM

- * MERRO PHARMACEUTICAL CO LTD
 - IBUPROFEN, IBUPROFEN (OTC)

MERZ PHARMS

- * MERZ PHARMACEUTICALS LLC
 - ERYGEL, ERYTHROMYCIN
 - NAFTIN, NAFTIFINE HYDROCHLORIDE

METHAPHARM

- * METHAPHARM INC
 - LEVETIRACETAM, LEVETIRACETAM
 - PROVOCHOLINE, METHACHOLINE CHLORIDE

MICRO LABS LTD

- * MICRO LABS LTD
 - MEFENAMIC ACID, MEFENAMIC ACID
 - SIMVASTATIN, SIMVASTATIN

MIKAH PHARMA

- * MIKAH PHARMA LLC
 - ISRADIPINE, ISRADIPINE
 - PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 - TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

MIKART

- * MIKART INC
 - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******* MIKART INC**

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 AMINOCAPROIC ACID, AMINOCAPROIC ACID
 AMINOCAPROIC, AMINOCAPROIC ACID
 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
 ESGIC-PLUS, ACETAMINOPHEN
 ETHOSUXIMIDE, ETHOSUXIMIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 ISONIAZID, ISONIAZID
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 METHAZOLAMIDE, METHAZOLAMIDE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
 PYRAZINAMIDE, PYRAZINAMIDE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

MILLENNIUM PHARMS

* MILLENNIUM PHARMACEUTICALS INC
 VELCADE, BORTEZOMIB

MIRROR PHARMS

* MIRROR PHARMACEUTICALS LLC
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 CARISOPRODOL AND ASPIRIN, ASPIRIN
 CARISOPRODOL, CARISOPRODOL
 CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
 PROBENECID AND COLCHICINE, COLCHICINE

MISSION PHARMA

* MISSION PHARMACAL CO
 LITHOSTAT, ACETOHYDROXAMIC ACID
 TEXACORT, HYDROCORTISONE
 TINDAMAX, TINIDAZOLE
 TIOPRONIN, TIOPRONIN
 UROCIT-K, POTASSIUM CITRATE

MN PHARMS

* MN PHARMACEUTICALS
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

MOLNLYCKE HLTH

* MOLNLYCKE HEALTH CARE
 HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
 HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

MONARCH PHARMS

* MONARCH PHARMACEUTICALS INC
 CORTISPORIN, BACITRACIN ZINC
 CORTISPORIN, HYDROCORTISONE
 CORTISPORIN, HYDROCORTISONE ACETATE
 MENEST, ESTROGENS, ESTERIFIED
 NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
 NEOSPORIN, GRAMICIDIN
 PEDIOTIC, HYDROCORTISONE
 SEPTRA DS, SULFAMETHOXAZOLE
 SEPTRA, SULFAMETHOXAZOLE
 THALITONE, CHLORTHALIDONE
 VIROPTIC, TRIFLURIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MONOSOL RX LLC**

- * MONOSOL RX LLC
ZUPLENZ, ONDANSETRON

MORTON GROVE

- * MORTON GROVE PHARMACEUTICALS INC
LACTULOSE, LACTULOSE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

MORTON GROVE PHARMS

- * MORTON GROVE PHARMACEUTICALS INC
GENERLAC, LACTULOSE

MSD CONSUMER

- * MSD CONSUMER CARE INC
ZEGERID OTC, OMEPRAZOLE (OTC)

MSD INTL

- * MSD INTERNATIONAL GMBH
VYTORIN, EZETIMIBE

MSP SINGAPORE

- * MSP SINGAPORE CO LLC
ZETIA, EZETIMIBE

MURTY PHARMS

- * MURTY PHARMACEUTICALS INC
DIPYRIDAMOLE, DIPYRIDAMOLE

MUSTAFA NEVSAT

- * MUSTAFA NEVSAT ILAC SANAYII AS
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
METHYLREDNISOLONE SODIUM SUCCINATE, METHYLREDNISOLONE SODIUM SUCCINATE

MUSTAFA NEVZAT

- * MUSTAFA NEVZAT ILAC SANAYII AS
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
VECURONIUM BROMIDE, VECURONIUM BROMIDE

MUTUAL PHARM

- * MUTUAL PHARMACEUTICAL CO INC
ACETAZOLAMIDE, ACETAZOLAMIDE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
BACTRIM DS, SULFAMETHOXAZOLE
BACTRIM, SULFAMETHOXAZOLE
CARISOPRODOL, CARISOPRODOL
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ERGOLOID MESYLATES, ERGOLOID MESYLATES
FELODIPINE, FELODIPINE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
LOVASTATIN, LOVASTATIN
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE, METRONIDAZOLE
MINOXIDIL, MINOXIDIL
NYSTATIN, NYSTATIN
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PIROXICAM, PIROXICAM
PREDNISONE, PREDNISONE
PRIMIDONE, PRIMIDONE
PROPafenone HYDROCHLORIDE, PROPafenone HYDROCHLORIDE
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MUTUAL PHARMACEUTICAL CO INC
 - QUINIDINE SULFATE, QUINIDINE SULFATE
 - SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - SPIRONOLACTONE, SPIRONOLACTONE
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 - SULINDAC, SULINDAC
 - TEMAZEPAM, TEMAZEPAM
 - THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 - TOLMETIN SODIUM, TOLMETIN SODIUM
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

MUTUAL PHARM CO INC

- * MUTUAL PHARMACEUTICAL CO INC
 - LEVETIRACETAM, LEVETIRACETAM

MUTUAL PHARMA

- * MUTUAL PHARMACAL CO
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - KETOCONAZOLE, KETOCONAZOLE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

MYLAN

- * MYLAN LABORATORIES INC
 - ACYCLOVIR, ACYCLOVIR
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
 - CAPTOPRIL, CAPTOPRIL
 - ETODOLAC, ETODOLAC
 - TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
- * MYLAN PHARMACEUTICALS
 - METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- * MYLAN PHARMACEUTICALS INC
 - ACARBOSE, ACARBOSE
 - ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 - ACYCLOVIR, ACYCLOVIR
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - ALENDRONATE SODIUM, ALENDRONATE SODIUM
 - ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
 - ALLOPURINOL, ALLOPURINOL
 - ALPRAZOLAM, ALPRAZOLAM
 - AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 - AMNESTEEM, ISOTRETINOIN
 - ANASTROZOLE, ANASTROZOLE
 - ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 - ATENOLOL, ATENOLOL
 - ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
 - 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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
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)
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLOROTHIAZIDE, CHLOROTHIAZIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORTHALIDONE, CHLORTHALIDONE
CILOSTAZOL, CILOSTAZOL
CIMETIDINE, CIMETIDINE
CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLARITHROMYCIN, CLARITHROMYCIN
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLORAZEPATE DIPOTASSUM, CLORAZEPATE DIPOTASSUM
CLORPRES, CHLORTHALIDONE
CLOZAPINE, CLOZAPINE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
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
DOXEPIН HYDROCHLORIDE, DOXEPIН HYDROCHLORIDE
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
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
FAMCICLOVIR, FAMCICLOVIR
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
FELODIPINE, FELODIPINE
FENOFIBRATE, FENOFIBRATE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
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
FLUTAMIDE, FLUTAMIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
FUROSEMIDE, FUROSEMIDE
GABAPENTIN, GABAPENTIN
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
GLIMEPIRIDE, GLIMEPIRIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLIPIZIDE, GLIPIZIDE
GLYBURIDE (MICRONIZED), GLYBURIDE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
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
LEVOFLOXACIN, LEVOFLOXACIN
LEVOTHYROXINE SODIUM, LEVOTHYROXINE 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
MENTAX-TC, BUTENAFINE HYDROCHLORIDE
MERCAPTOPURINE, MERCAPTOPURINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHIMAZOLE, METHIMAZOLE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHYCLOTHIAZIDE, METHYCLOTHIAZIDE
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPA, METHYLDOPA
METOLAZONE, METOLAZONE
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NADOLOL AND BENDROFLUMETHAZIDE, BENDROFLUMETHAZIDE
NADOLOL, NADOLOL
NAPROXEN, NAPROXEN
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NIFEDIPINE, NIFEDIPINE
NISOLDIPINE, NISOLDIPINE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NIZATIDINE, NIZATIDINE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
OMEPRAZOLE, OMEPRAZOLE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ****

- * MYLAN PHARMACEUTICALS INC
OXAPROZIN, OXAPROZIN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PACLITAXEL, PACLITAXEL
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
PEG-3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE, POLYETHYLENE GLYCOL 3350
PENTOXIFYLLINE, PENTOXIFYLLINE
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PHENYTEK, PHENYTOIN SODIUM
PINDOLOL, PINDOLOL
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PIROXICAM, PIROXICAM
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PROBENECID, PROBENECID
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
RISPERIDONE, RISPERIDONE
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SERTRALINE HYDROCHLORIDE, SERTRALINE 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
TEMAZEPM, TEMAZEPM
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTHIXENE, THIOTHIXENE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TOPIRAMATE, TOPIRAMATE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRANDOLAPRIL, TRANDOLAPRIL
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
URSODIOL, URSODIOL
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
WARFARIN SODIUM, WARFARIN SODIUM
ZALEPLON, ZALEPLON
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
ZONISAMIDE, ZONISAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** M ******MYLAN BERTEK**

- * MYLAN BERTEK PHARMACEUTICALS INC
 - AVITA, TRETINOIN
 - MAXZIDE, HYDROCHLOROTHIAZIDE
 - MAXZIDE-25, HYDROCHLOROTHIAZIDE

MYLAN INSTITUTIONAL

- * MYLAN INSTITUTIONAL LLC
 - ALOPRIM, ALLOPURINOL SODIUM
 - DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
 - DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - ENLON-PLUS, ATROPINE SULFATE
 - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
 - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 - TRANEXAMIC ACID, TRANEXAMIC ACID
 - ULTIVA, REMIFENTANIL HYDROCHLORIDE

MYLAN PHARMS INC

- * MYLAN PHARMACEUTICALS INC
 - EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
 - FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LEVETIRACETAM, LEVETIRACETAM
 - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 - MORPHINE SULFATE, MORPHINE SULFATE

MYLAN TECHNOLOGIES

- * MYLAN TECHNOLOGIES INC
 - CLONIDINE, CLONIDINE
 - ESTRADIOL, ESTRADIOL
 - FENTANYL-100, FENTANYL
 - FENTANYL-12, FENTANYL
 - FENTANYL-25, FENTANYL
 - FENTANYL-50, FENTANYL
 - FENTANYL-75, FENTANYL
 - NITROGLYCERIN, NITROGLYCERIN

NATCO PHARMA

- * NATCO PHARMA LTD
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

NATCO PHARMA LTD

- * NATCO PHARMA LIMITED
 - CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
- * NATCO PHARMA LTD
 - ANASTROZOLE, ANASTROZOLE
 - CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - LETROZOLE, LETROZOLE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

NAUTILUS NEUROSCIENC

- * NAUTILUS NEUROSCIENCES INC
 - CAMBIA, DICLOFENAC POTASSIUM

NAVINTA LLC

- * NAVINTA LLC
 - FAMOTIDINE, FAMOTIDINE
 - FOMEPIZOLE, FOMEPIZOLE
 - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NESHER PHARMS**

- * NESHER PHARMACEUTICALS USA LLC
 - BENZONATATE, BENZONATATE
 - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 - DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 - MICRO-K 10, POTASSIUM CHLORIDE
 - MICRO-K, POTASSIUM CHLORIDE
 - MORPHINE SULFATE, MORPHINE SULFATE
 - NYSTATIN, NYSTATIN
 - ONDANSETRON, ONDANSETRON
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 - PREDNISOLONE, PREDNISOLONE
 - PROPafenone HYDROCHLORIDE, PROPafenone HYDROCHLORIDE

NEUROGESX

- * NEUROGESX INC
 - QUTENZA, CAPSAICIN

NEW RIVER

- * NEW RIVER PHARMACEUTICALS INC
 - PROFERDEX, IRON DEXTRAN

NEXGEN PHARMA

- * NEXGEN PHARMA INC
 - BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - CHENODIOL, CHENODIOL
 - 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

NEXUS PHARMS

- * NEXUS PHARMACEUTICALS INC
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVOFLOXACIN, LEVOFLOXACIN

NIAGARA PHARMS

- * NIAGARA PHARMACEUTICALS INC
 - PUR-WASH, PURIFIED WATER (OTC)

NORTHSTAR HILTHCARE

- * NORTHSTAR HEALTHCARE HOLDINGS LTD
 - ALLOPURINOL, ALLOPURINOL
 - ATENOLOL, ATENOLOL
 - BACLOFEN, BACLOFEN
 - GEMFIBROZIL, GEMFIBROZIL
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - IBUPROFEN, IBUPROFEN
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NOSTRUM**

- * NOSTRUM PHARMACEUTICALS INC
CARBAMAZEPINE, CARBAMAZEPINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
THEOPHYLLINE, THEOPHYLLINE

NOSTRUM LABS

- * NOSTRUM LABORATORIES INC
PIROXICAM, PIROXICAM
SUCRALFATE, SUCRALFATE

NOVADEL

- * NOVADEL PHARMA INC
NITROMIST, NITROGLYCERIN
ZOLPIMIST, ZOLPIDEM TARTRATE

NOVARTIS

- * NOVARTIS CONSUMER HEALTH INC
EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
HABITROL, NICOTINE (OTC)
LAMISIL AT, TERBINAFINE (OTC)
LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
NEXCIDE, KETOPROFEN (OTC)
PREVACID 24 HR, LANSOPRAZOLE (OTC)
TAVIST ALLERGY/SINUS/HEADACHE, ACETAMINOPHEN (OTC)
THRIVE, NICOTINE POLACRILEX (OTC)
TRANSDERM SCOP, SCOPOLAMINE
VAGISTAT-1, TIOCONAZOLE (OTC)
VOLTAREN, DICLOFENAC SODIUM
- * NOVARTIS PHARMACEUTICALS CORP
AFINITOR, EVEROLIMUS
AMTURNIDE, ALISKIREN HEMIFUMARATE
ARCAPTA NEOHALER, INDACATEROL MALEATE
AREDIA, PAMIDRONATE DISODIUM
CATAFLAM, DICLOFENAC POTASSIUM
CLOZARIL, CLOZAPINE
COARTEM, ARTEMETHER
COMBIPATCH, ESTRADIOL
DESFERAL, DEFEROXAMINE MESYLATE
DIOVAN HCT, HYDROCHLOROTHIAZIDE
DIOVAN, VALSARTAN
ESTRADERM, ESTRADIOL
EXELON, RIVASTIGMINE
EXELON, RIVASTIGMINE TARTRATE
EXFORGE HCT, AMLODIPINE BESYLATE
EXFORGE, AMLODIPINE BESYLATE
EXJADE, DEFERASIROX
FAMVIR, FAMCICLOVIR
FANAPT, ILOPERIDONE
FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
FORADIL, FORMOTEROL FUMARATE
GILENYA, FINGOLIMOD
GLEEVEC, IMATINIB MESYLATE
HYDERGINE, ERGOLOID MESYLATES
LAMISIL, TERBINAFINE HYDROCHLORIDE
LAMPRENE, CLOFAZIMINE
LESCOL XL, FLUVASTATIN SODIUM
LESCOL, FLUVASTATIN SODIUM
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPRESSOR, METOPROLOL TARTRATE
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
LOTENSIN, BENAZEPRIL HYDROCHLORIDE
LOTREL, AMLODIPINE BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** N **

* NOVARTIS PHARMACEUTICALS CORP
 METHERGINE, METHYLERGONOVINE MALEATE
 METOPIRONE, METYRAPONE
 MIACALCIN, CALCITONIN SALMON
 MYFORTIC, MYCOPHENOLIC ACID
 NEORAL, CYCLOSPORINE
 OCUPRESS, CARTEOLOL HYDROCHLORIDE
 PARLODEL, BROMOCRIPTINE MESYLATE
 RECLAST, ZOLEDRONIC ACID
 REGITINE, PHENTOLAMINE MESYLATE
 RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN, METHYLPHENIDATE HYDROCHLORIDE
 RITALIN-SR, METHYLPHENIDATE HYDROCHLORIDE
 SANDIMMUNE, CYCLOSPORINE
 SANDOSTATIN LAR, OCTREOTIDE ACETATE
 SANDOSTATIN, OCTREOTIDE ACETATE
 STARLIX, NATEGLINIDE
 TASIGNA, NILOTINIB HYDROCHLORIDE MONOHYDRATE
 TAVIST-1, CLEMASTINE FUMARATE (OTC)
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TEKAMLO, ALISKIREN HEMIFUMARATE
 TEKTURN A HCT, ALISKIREN HEMIFUMARATE
 TEKTURN A, ALISKIREN HEMIFUMARATE
 TRILEPTAL, OXCARBAZEPINE
 TYZEKA, TELBIVUDINE
 VALTURNA, ALISKIREN HEMIFUMARATE
 VASOCIDIN, PREDNISOLONE SODIUM PHOSPHATE
 VIVELLE, ESTRADIOL
 VIVELLE-DOT, ESTRADIOL
 VOLTAREN, DICLOFENAC SODIUM
 VOLTAREN-XR, DICLOFENAC SODIUM
 ZOMETA, ZOLEDRONIC ACID
 ZORTRESS, EVEROLIMUS

NOVARTIS PHARMS

* NOVARTIS PHARMACEUTICALS CORP
 FEMARA, LETROZOLE
 TOBI, TOBRAMYCIN

NOVAST LABS LTD

* NOVAST LABORATORIES LTD
 DASETTA 1/35, ETHINYL ESTRADIOL
 DASETTA 7/7/7, ETHINYL ESTRADIOL
 LEVONEST, ETHINYL ESTRADIOL
 PHILITH, ETHINYL ESTRADIOL

NOVEL LABS INC

* NOVEL LABORATORIES INC
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
 MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE, SODIUM PHOSPHATE, DIBASIC,
 ANHYDROUS
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 PEG-3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE, POLYETHYLENE GLYCOL 3350
 PHENELZINE SULFATE, PHENELZINE SULFATE
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 SULFAMETHOPRIM, SULFAMETHOXAZOLE
 SULFAMETHOPRIM-DS, SULFAMETHOXAZOLE
 TEMAZEPAM, TEMAZEPAM
 TRIMETHOPRIM, TRIMETHOPRIM

NOVEN

* NOVEN PHARMACEUTICALS INC
 FENTANYL-100, FENTANYL
 FENTANYL-25, FENTANYL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ****

* NOVEN PHARMACEUTICALS INC
 FENTANYL-50, FENTANYL
 FENTANYL-75, FENTANYL

NOVEN PHARMS INC

* NOVEN PHARMACEUTICALS INC
 DAYTRANA, METHYLPHENIDATE

NOVEN THERAP

* NOVEN THERAPEUTICS LLC
 LITHOBID, LITHIUM CARBONATE
 PEDEXA, PAROXETINE MESYLATE

NOVEX

* NOVEX PHARMA
 ALBUTEROL SULFATE, ALBUTEROL SULFATE
 BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 CYCLOSPORINE, CYCLOSPORINE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LACTULOSE, LACTULOSE
 LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 OFLOXACIN, OFLOXACIN
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 TOBRAMYCIN, TOBRAMYCIN

NOVO NORDISK

* NOVO NORDISK PHARMACEUTICALS INC
 GLUCAGEN, GLUCAGON HYDROCHLORIDE RECOMBINANT

NOVO NORDISK INC

* NOVO NORDISK INC
 ACTIVELLA, ESTRADIOL
 LEVEMIR, INSULIN DETEMIR RECOMBINANT
 NORDITROPIN FLEXPRO, SOMATROPIN RECOMBINANT
 NORDITROPIN NORDIFLEX, SOMATROPIN RECOMBINANT
 NORDITROPIN, SOMATROPIN RECOMBINANT
 NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
 NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
 NOVOLOG, INSULIN ASPART RECOMBINANT
 PRANDIMET, METFORMIN HYDROCHLORIDE
 PRANDIN, REPAGLINIDE
 VAGIFEM, ESTRADIOL
 VICTOZA, LIRAGLUTIDE RECOMBINANT

NOVOCOL

* NOVOCOL PHARMACEUTICAL INC
 ISOCOCAINE HYDROCHLORIDE W/ LEVONORDEFRIN, LEVONORDEFRIN
 ISOCOCAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE

NOVOCOL INC

* NOVOCOL INC
 LIDOCAINE, LIDOCAINE

NU PHARM

* NU PHARM INC
 DIVALPROEX SODIUM, DIVALPROEX SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** N ******NYCOMED US**

- * NYCOMED US INC
 - ADAPALENE, ADAPALENE
 - ALVESCO, CICLESONIDE
 - CALCIPOTRIENE, CALCIPOTRIENE
 - CARMOL HC, HYDROCORTISONE ACETATE
 - CICLOPIROX, CICLOPIROX
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - DESOXIMETASONE, DESOXIMETASONE
 - IMIQUIMOD, IMIQUIMOD
 - LIDOCaine AND PRILOCAINE, LIDOCaine
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - OMNARIS, CICLESONIDE
 - TEMOVATE, CLOBETASOL PROPIONATE
 - TERCONAZOLE, TERCONAZOLE
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 - TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE
 - ZONALON, DOXEpin HYDROCHLORIDE

OAK PHARMS

- * OAK PHARMACEUTICALS INC
 - EMLA, LIDOCaine
 - XYLOCAINE, LIDOCaine HYDROCHLORIDE

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)
 - IBUPROFEN, IBUPROFEN
 - IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
 - IBUPROHM, IBUPROFEN (OTC)
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

OLTA PHARMS

- * OLTA PHARMACEUTICALS CORP
 - LINDANE, LINDANE

OMAN PHARM PRODUCTS

- * OMAN PHARMACEUTICAL PRODUCTS CO LLC
 - NEOMYCIN SULFATE, NEOMYCIN SULFATE

ONCO THERAPIES LTD

- * ONCO THERAPIES LTD
 - CARBOPLATIN, CARBOPLATIN
 - CLADRIBINE, CLADRIBINE
 - CYTARABINE, CYTARABINE
 - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - PACLITAXEL, PACLITAXEL

ONY

- * ONY INC
 - INFASURF PRESERVATIVE FREE, CALFACTANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** O ******OPTIMER PHARMS**

- * OPTIMER PHARMACEUTICALS INC
DIFICID, FIDAXOMICIN

ORAPHARMA

- * ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

ORCHID HLTHCARE

- * ORCHID HEALTHCARE
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
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
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
LEVETIRACETAM, LEVETIRACETAM
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- * ORCHID HEALTHCARE DIV ORCHID CHEMICALS AND PHARMACEUTICALS LTD
ZALEPLON, ZALEPLON

ORGANON USA INC

- * ORGANON USA INC
CYCLESSA, DESOGESTREL
DESOGEN, DESOGESTREL
FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
GANIRELIX ACETATE INJECTION, GANIRELIX ACETATE
IMPLANON, ETONOGESTREL
NEXPLANON, ETONOGESTREL
NUVARING, ETHINYLMESTRADIOL
PREGNYL, GONADOTROPIN, CHORIONIC
REMERON SOLTAB, MIRTAZAPINE
REMERON, MIRTAZAPINE
SAPHRIS, ASENAPINE MALEATE

ORION

- * ORION CORP
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
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
ERGOCALCIFEROL, ERGOCALCIFEROL

ORPHAN EUROPE

- * ORPHAN EUROPE
CARBAGLU, CARGLUMIC ACID

ORTHO BIOTECH

- * ORTHO BIOTECH PRODUCTS LP
DOXIL, DOXORUBICIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** O **

ORTHO JANSSEN

- * ORTHO DERMATOLOGICS DIV JANSSEN PHARMACEUTICALS INC
ERTACZO, SERTACONAZOLE NITRATE
- RENOVA, TRETINOIN
- RETIN-A, TRETINOIN

ORTHO MCNEIL JANSSEN

- * ORTHO MCNEIL JANSSEN PHARMACEUTICALS INC
NIZORAL, KETOCONAZOLE

ORTHONEUTROGENA

- * ORTHONEUTROGENA
GRIFULVIN V, GRISEOFULVIN, MICROCRYSTALLINE

OSI PHARMS

- * OSI PHARMACEUTICALS INC
TARCEVA, ERLOTINIB HYDROCHLORIDE

OSMOTICA PHARM

- * OSMOTICA PHARMACEUTICAL CORP
NIFEDIPINE, NIFEDIPINE
- OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
- VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

OTSUKA

- * OTSUKA PHARMACEUTICAL CO LTD
ABILIFY, ARIPIPRAZOLE
- PLETAL, CILOSTAZOL
- * OTSUKA PHARMACEUTICAL DEVELOPMENT AND COMMERCIALIZATION INC
ABILIFY, ARIPIPRAZOLE

OTSUKA AMERICA

- * OTSUKA AMERICA PHARMACEUTICALS INC
BREATHTEK UBT FOR H-PYLORI, UREA C-13

OTSUKA AMERICA PHARM

- * OTSUKA AMERICA PHARMACEUTICAL INC
SAMSCA, TOLVAPTA

OTSUKA PHARM

- * OTSUKA PHARMACEUTICAL CO LTD
BUSULFEX, BUSULFAN

OUTLOOK PHARMS

- * OUTLOOK PHARMACEUTICALS INC
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

PACIFIC PHARMA

- * PACIFIC PHARMA
TIMOLOL MALEATE, TIMOLOL MALEATE
- * PACIFIC PHARMA INC
TIMOLOL MALEATE, TIMOLOL MALEATE

PACIRA PHARMS INC

- * PACIRA PHARMACEUTICALS INC
DEPOCYT, CYTARABINE
- EXPAREL, BUPIVACAINE

PACK PHARMS LLC

- * PACK PHARMACEUTICALS LLC
CROMOLYN SODIUM, CROMOLYN SODIUM

PADDOCK

- * PADDOCK LABORATORIES INC
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE

PADDOCK LABS

- * PADDOCK LABORATORIES INC
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE

PADDOCK LLC

- * PADDOCK LABORATORIES LLC
AMMONIUM LACTATE, AMMONIUM LACTATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

* PADDOCK LABORATORIES LLC
 BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
 CAFFEINE CITRATE, CAFFEINE CITRATE
 CALCIUM ACETATE, CALCIUM ACETATE
 CICLOPIROX, CICLOPIROX
 CLENZ-LYTE, POLYETHYLENE GLYCOL 3350
 CLINDA-DERM, CLINDAMYCIN PHOSPHATE
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 CLOTRIMAZOLE, CLOTRIMAZOLE
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 COLOCORT, HYDROCORTISONE
 COMPRO, PROCHLORPERAZINE
 DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
 ERYTHRA-DERM, ERYTHROMYCIN
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 INDOMETHACIN, INDOMETHACIN
 KIONEX, SODIUM POLYSTYRENE SULFONATE
 LATANOPROST, LATANOPROST
 LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
 LORAZEPAM, LORAZEPAM
 MIDAMOR, AMILORIDE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
 NYSTOP, NYSTATIN
 PODOFILOX, PODOFILOX
 POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
 TROSPiUM CHLORIDE, TROSPiUM CHLORIDE

PALADIN LABS

* PALADIN LABS USA INC
 ANTIZOL, FOMEPIZOLE

PAR PHARM

* PAR PHARMACEUTICAL
 NABUMETONE, NABUMETONE
 OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
 PROPafenone HYDROCHLORIDE, PROPafenone HYDROCHLORIDE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 * PAR PHARMACEUTICAL INC
 ALPRAZOLAM, ALPRAZOLAM
 AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 AMLODIPINE BEsylate AND BENazepril HYDROCHLORIDE, AMLODIPINE BEsylate
 CABERGOLINE, CABERGOLINE
 CALCITONIN-SALMON, CALCITONIN SALMON
 CAPOTEN, CAPTOPRIL
 CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 CHOLESTYRAMINE, CHOLESTYRAMINE
 CLOMIPHENe CITRATE, CLOMIPHENE CITRATE
 CLONAZEPAM, CLONAZEPAM
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

* PAR PHARMACEUTICAL INC
 ESTAZOLAM, ESTAZOLAM
 FLUTAMIDE, FLUTAMIDE
 GLYCOPYRROLATE, GLYCOPYRROLATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYUREA, HYDROXYUREA
 IBUPROFEN, IBUPROFEN (OTC)
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 LEVETIRACETAM, LEVETIRACETAM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MEGACE ES, MEGESTROL ACETATE
 MEGESTROL ACETATE, MEGESTROL ACETATE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METRONIDAZOLE, METRONIDAZOLE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 NASCOBAL, CYANOCOBALAMIN
 NATEGLINIDE, NATEGLINIDE
 NYSTATIN, NYSTATIN
 OLANZAPINE, OLANZAPINE
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 ONDANSETRON, ONDANSETRON
 OXANDROLONE, OXANDROLONE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RISPERIDONE, RISPERIDONE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 TORSEMIDE, TORSEMIDE
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 TRANLYLCYPROMINE SULFATE, TRANLYLCYPROMINE SULFATE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

PARAPRO PHARMS

* PARAPRO PHARMACEUTICALS LLC
 NATROBA, SPINOSAD

PARKE DAVIS

* PARKE DAVIS DIV WARNER LAMBERT CO
 CELONTIN, METHSUXIMIDE
 DILANTIN, PHENYTOIN SODIUM
 DILANTIN-125, PHENYTOIN
 NARDIL, PHENELZINE SULFATE
 NEURONTIN, GABAPENTIN
 ZARONTIN, ETHOSUXIMIDE
* PARKE DAVIS PHARMACEUTICAL RESEARCH DIV WARNER LAMBERT CO
 ZARONTIN, ETHOSUXIMIDE

PBS

* PBS REGULATORY CONSULTING SERVICES INC
 LEUKERAN, CHLORAMBUCIL

PEDIATRX

* PEDIATRX INC
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE

PEDINOL

* PEDINOL PHARMACAL INC
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 NALFON, FENOPROFEN CALCIUM

PERNIX THERAP

* PERNIX THERAPEUTICS LLC
 CEDAX, CEFTIBUTEN DIHYDRATE

PERRIGO

* L PERRIGO CO
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * L PERRIGO CO
 - 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)
- * PERRIGO CO
 - CICLOPIROX, CICLOPIROX
 - CIMETIDINE, CIMETIDINE (OTC)
 - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
 - CLINDETS, CLINDAMYCIN PHOSPHATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - ERYTHROMYCIN, ERYTHROMYCIN
 - FAMOTIDINE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 - IBUPROFEN, IBUPROFEN (OTC)
 - JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 - MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 - PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 - STIE-CORT, HYDROCORTISONE

PERRIGO CO TENNESSEE

- * PERRIGO CO TENNESSEE INC
 - GRISEOFULVIN, GRISEOFULVIN, MICROCRYSTALLINE
 - SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM

PERRIGO ISRAEL

- * PERRIGO ISRAEL PHARMACEUTICALS LTD
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 - IMIQUIMOD, IMIQUIMOD
 - KETOCONAZOLE, KETOCONAZOLE
 - MESALAMINE, MESALAMINE
 - MINOXIDIL, MINOXIDIL (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

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PERRIGO NEW YORK INC
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - GENTAMICIN SULFATE, GENTAMICIN SULFATE
 - HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 - HYDROCORTISONE, HYDROCORTISONE
 - KETOCONAZOLE, KETOCONAZOLE
 - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 - 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 R AND D

- * PERRIGO R AND D CO
 - CALCIUM CARBONATE, FAMOTIDINE AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
 - 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
 - GUAIFENESIN, GUAIFENESIN (OTC)
 - IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
 - IBUPROFEN, IBUPROFEN
 - IBUPROFEN, IBUPROFEN (OTC)
 - LEVONORGESTREL, LEVONORGESTREL
 - LEVONORGESTREL, LEVONORGESTREL (OTC)
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 - NAPROXEN, NAPROXEN
 - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 - POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
- * PERRIGO R AND D COMPANY
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

PERRIGO UK FINCO

- * PERRIGO UK FINCO LTD PARTNERSHIP
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

PETNET

- * PETNET SOLUTIONS INC
 - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

PFIZER

- * PFIZER CENTRAL RESEARCH
 - DIFLUCAN, FLUCONAZOLE
 - ZITHROMAX, AZITHROMYCIN
- * PFIZER CHEMICALS DIV PFIZER INC
 - DIFLUCAN, FLUCONAZOLE
 - ZITHROMAX, AZITHROMYCIN
- * PFIZER INC
 - ANTIVERT, MECLIZINE HYDROCHLORIDE
 - ARGATROBAN, ARGATROBAN
 - CADUET, AMLODIPINE BESYLATE
 - CARDURA XL, DOXAZOSIN MESYLATE
 - CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 - DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
 - DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 - DIFLUCAN IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PFIZER INC
 - FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE
 - FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 - FLUMAZENIL, FLUMAZENIL
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - GEODON, ZIPRASIDONE HYDROCHLORIDE
 - GEODON, ZIPRASIDONE MESYLATE
 - GLUCOTROL XL, GLIPIZIDE
 - GLUCOTROL, GLIPIZIDE
 - HALOPERIDOL, HALOPERIDOL LACTATE
 - HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 - HEPARIN SODIUM, HEPARIN SODIUM
 - I索丙丁SR, VERAPAMIL HYDROCHLORIDE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - LIPITOR, ATORVASTATIN CALCIUM
 - NAVANE, THIOTHIXENE
 - NORVASC, AMLODIPINE BESYLATE
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 - PROCARDIA, NIFEDIPINE
 - REVATIO, SILDENAFIL CITRATE
 - RIFAMPIN, RIFAMPIN
 - TOVIAZ, FESOTERODINE FUMARATE
 - UNASYN, AMPICILLIN SODIUM
 - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 - VECURONIUM BROMIDE, VECURONIUM BROMIDE
 - VFEND, VORICONAZOLE
 - XALKORI, CRIZOTINIB
 - ZITHROMAX, AZITHROMYCIN
- * PFIZER LABORATORIES DIV PFIZER INC
 - CARDURA, DOXAZOSIN MESYLATE
 - DIABINESE, CHLORPROPAMIDE
 - FELDENE, PIROXICAM
 - MINIPRESS, PRAZOSIN HYDROCHLORIDE
 - PERMAPEN, PENICILLIN G BENZATHINE
 - PFIZERPEN, PENICILLIN G POTASSIUM
 - PROCARDIA XL, NIFEDIPINE
 - TERRAMYCIN W/ POLYMYXIN B SULFATE, OXYTETRACYCLINE HYDROCHLORIDE
 - UNASYN, AMPICILLIN SODIUM
 - VIBRAMYCIN, DOXYCYCLINE
 - VIBRAMYCIN, DOXYCYCLINE CALCIUM
 - VIBRAMYCIN, DOXYCYCLINE HYCLATE
 - VISTARIL, HYDROXYZINE PAMOATE
- * PFIZER PHARMACEUTICALS INC
 - ZOLOFT, SERTRALINE HYDROCHLORIDE
- * PFIZER PHARMACEUTICALS PRODUCTION CORP LTD
 - TIKOSYN, DOFETILIDE

PFIZER CONS HLTHCARE

- * PFIZER CONSUMER HEALTHCARE
 - ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)
 - ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)

PFIZER GLOBAL

- * PFIZER GLOBAL RESEARCH DEVELOPMENT
 - ZMAX, AZITHROMYCIN

PFIZER INC

- * PFIZER INC
 - CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 - CHANTIX, VARENICLINE TARTRATE
 - ELLENCE, EPIRUBICIN HYDROCHLORIDE
 - GEODON, ZIPRASIDONE HYDROCHLORIDE
 - NICOTROL, NICOTINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

PFIZER IRELAND

- * PFIZER IRELAND PHARMACEUTICALS
RELPAX, ELETRIPTAN HYDROBROMIDE
VIAGRA, SILDENAFIL CITRATE

PFIZER PHARMS

- * PFIZER PHARMACEUTICALS LTD
ACCUPRIL, QUINAPRIL HYDROCHLORIDE
ACCURETIC, HYDROCHLORTHIAZIDE
DILANTIN, PHENYTOIN
LOPID, GEMFIBROZIL
NEURONTIN, GABAPENTIN
NITROSTAT, NITROGLYCERIN

PHARM ASSOC

- * PHARMACEUTICAL ASSOC INC
DOXEPEPIN HYDROCHLORIDE, DOXEPEPIN HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LORAZEPAM, LORAZEPAM
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
- * PHARMACEUTICAL ASSOC INC DIV BEACH PRODUCTS
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
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PREDNISOLONE, PREDNISOLONE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
VALPROIC ACID, VALPROIC ACID

PHARMA SERVE NY

- * PHARMA SERVE INC SUB TORIGIAN LABORATORIES
AMINOPHYLLINE, AMINOPHYLLINE

PHARMACHEMIE

- * PHARMACHEMIE BV
CARBOPLATIN, CARBOPLATIN
CISPLATIN, CISPLATIN
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
ETOPOSIDE, ETOPOSIDE

PHARMACHEMIE BV

- * PHARMACHEMIE BV
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
CARBOPLATIN, CARBOPLATIN
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

PHARMACIA AND UPJOHN

- * PHARMACIA AND UPJOHN
XANAX XR, ALPRAZOLAM
- * PHARMACIA AND UPJOHN CO
ANSAID, FLURBIPROFEN
AROMASIN, EXEMESTANE
AZULFIDINE EN-TABS, SULFASALAZINE
AZULFIDINE, SULFASALAZINE
BACITRACIN, BACITRACIN
CAVERJECT IMPULSE, ALPROSTADIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PHARMACIA AND UPJOHN CO
 - 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
 - COLESTID, COLESTIPOL HYDROCHLORIDE
 - CORTEF, HYDROCORTISONE
 - CONVERT, IBUTILIDE FUMARATE
 - CYKLOKAPRON, TRANEXAMIC ACID
 - DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 - DEPO-MEDROL, METHYLPPREDNISOLONE ACETATE
 - DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 - DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 - DETROL LA, TOLTERODINE TARTRATE
 - DETROL, TOLTERODINE TARTRATE
 - DIDREX, BENZPHETAMINE HYDROCHLORIDE
 - EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 - ESTRING, ESTRADIOL
 - FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 - 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, METHYLPPREDNISOLONE
 - MYCOBUTIN, RIFABUTIN
 - NICOTROL, NICOTINE
 - OGEN .625, ESTROPIPATE
 - OGEN 1.25, ESTROPIPATE
 - OGEN 2.5, ESTROPIPATE
 - 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, METHYLPPREDNISOLONE SODIUM SUCCINATE
 - SOMAVERT, PEGVISOMANT
 - XALATAN, LATANOPROST
 - XANAX, ALPRAZOLAM
 - ZINECARD, DEXRAZOXANE HYDROCHLORIDE
 - ZYVOX, LINEZOLID
- * PHARMACIA AND UPJOHN SUB PFIZER INC
 - DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE

PHARMAFORCE

- * PHARMAFORCE INC
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
 - NANDROLONE DECANOATE, NANDROLONE DECANOATE
 - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 - TRANEXAMIC ACID, TRANEXAMIC ACID

PHARMLUCENCE

- * PHARMLUCENCE INC
 - AN-DTPA, TECHNETIUM TC-99M PENTETATE KIT
 - AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
 - CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ****

- * PHARMALUCENCE INC
 - CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 - HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
 - PULMOLITE, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 - TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
 - TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT

PHARMAX

- * PHARMAX GROUP INC
 - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE

PHARMAXIS

- * PHARMAXIS INC
 - ARIDOL KIT, MANNITOL

PIERRE FABRE

- * PIERRE FABRE MEDICAMENT
 - NAVELBINE, VINORELBINE TARTRATE

PIERREL

- * PIERREL S.P.A.
 - ARTICAINE HYDROCHLORIDE WITH EPINEPHRINE, ARTICAINE HYDROCHLORIDE

PINNACLE BIOLGS

- * PINNACLE BIOLOGICS INC
 - PHOTOFIRIN, PORFIMER SODIUM

PIRAMAL CRITICAL

- * PIRAMAL CRITICAL CARE INC
 - ENFLURANE, ENFLURANE
 - ISOFLURANE, ISOFLURANE
 - SOJOURN, SEVOFLURANE

PLANTEX

- * PLANTEX USA INC DIV IKAPHARM INC
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE

PLIVA

- * PLIVA INC
 - AZITHROMYCIN, AZITHROMYCIN
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - CARBOPLATIN, CARBOPLATIN
 - CHLORPROPAMIDE, CHLORPROPAMIDE
 - CHLORTHALIDONE, CHLORTHALIDONE
 - CIMETIDINE, CIMETIDINE
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 - CYCLOSPORINE, CYCLOSPORINE
 - DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 - FLUCONAZOLE, FLUCONAZOLE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - KETOCONAZOLE, KETOCONAZOLE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - METRONIDAZOLE, METRONIDAZOLE
 - NAPROXEN, NAPROXEN
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - PENTOXIFYLLINE, PENTOXIFYLLINE
 - PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 - PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - THEOPHYLLINE, THEOPHYLLINE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - WARFARIN SODIUM, WARFARIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** P ******PLIVA HRVATSKA DOO**

- * PLIVA HRVATSKA DOO
 - ADAPALENE, ADAPALENE
 - AZITHROMYCIN, AZITHROMYCIN
 - CARVEDILOL, CARVEDILOL
 - CILOSTAZOL, CILOSTAZOL
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - 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

POHL BOSKAMP

- * POHL BOSKAMP
 - NITROLINGUAL PUMPSpray, NITROGLYCERIN

POLYMEDICA

- * POLYMEDICA INDUSTRIES INC
 - ANESTACON, LIDOCaine HYDROCHLORIDE
 - NEOPAP, ACETAMINOPHEN (OTC)

PRASCO

- * PRASCO LLC DBA PRASCO LABORATORIES
 - ECONAZOLE NITRATE, ECONAZOLE NITRATE

PRECISION DERMAT

- * PRECISION DERMATOLOGY INC
 - PODOFILOX, PODOFILOX

PRECISION DOSE

- * PRECISION DOSE INC
 - RISPERIDONE, RISPERIDONE

PRINSTON INC

- * PRINSTON PHARMACEUTICAL INC
 - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 - CAPTOPRIL, CAPTOPRIL
 - LISINOPRIL AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE
 - LISINOPRIL, LISINOPRIL
 - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 - RISPERIDONE, RISPERIDONE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

PROCTER AND GAMBLE

- * PROCTER AND GAMBLE PHARMACEUTICALS INC SUB PROCTER AND GAMBLE CO
 - DIDRONEL, ETIDRONATE DISODIUM

PROGENICS

- * PROGENICS PHARMACEUTICALS INC
 - RELISTOR, METHYLNALTREXONE BROMIDE

PROMETHEUS LABS

- * PROMETHEUS LABORATORIES INC
 - HELIDAC, BISMUTH SUBSALICYLATE
 - IMURAN, AZATHIOPRINE
 - LOTRONEX, ALOSETRON HYDROCHLORIDE
 - MERCAPTOPURINE, MERCAPTOPURINE
 - RIDAURA, AURANOFIN
 - TRANDATE, LABETALOL HYDROCHLORIDE
 - ZYLOPRIM, ALLOPURINOL

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

PROSAM LABS

- * PROSAM LABS LLC
BACLOFEN, BACLOFEN
CARISOPRODOL AND ASPIRIN, ASPIRIN
CARISOPRODOL, CARISOPRODOL
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE

PROSTRAKAN INC

- * PROSTRAKAN INC
ABSTRAL, FENTANYL CITRATE
RECTIV, NITROGLYCERIN
SANCUSO, GRANisetron

PROVIDENT PHARM

- * PROVIDENT PHARMACEUTICAL INC
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

PULSION MEDCL

- * PULSION MEDICAL SYSTEMS AG
INDOCYANINE GREEN, INDOCYANINE GREEN

PURACAP PHARM

- * PURACAP PHARMACEUTICAL LLC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
MELOXICAM, MELOXICAM

PURDUE GMP

- * PURDUE GMP CENTER LLC DBA THE CHAO CENTER INDUSTRIAL PHARMACY
SEROMYCIN, CYCLOSERINE

PURDUE PHARM PRODS

- * PURDUE PHARMACEUTICAL PRODUCTS LP
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILAUDID-HP, HYDROMORPHONE HYDROCHLORIDE

PURDUE PHARMA

- * PURDUE PHARMA PRODUCTS LP
INTERMEZZO, ZOLPIDEM TARTRATE
RYZOLT, TRAMADOL HYDROCHLORIDE

PURDUE PHARMA LP

- * PURDUE PHARMA LP
BUTRANS, BUPRENORPHINE
MS CONTIN, MORPHINE SULFATE
OXYCONTIN, OXYCODONE HYDROCHLORIDE

PUREPAC PHARM

- * PUREPAC PHARMACEUTICAL CO
DIPYRIDAMOLE, DIPYRIDAMOLE

QLT

- * QLT INC
VISUDYNE, VERTEPORFIN

QOL MEDCL

- * QOL MEDICAL LLC
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
ETHAMOLIN, ETHANOLAMINE OLEATE
SUCRAID, SACROSIDASE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** Q **

QUESTCOR PHARMS

* QUESTCOR PHARMACEUTICALS INC
 DORAL, QUAZEPAM
 H.P. ACTHAR GEL, CORTICOTROPIN

RANBAXY

* RANBAXY INC
 GLYCOPYRROLATE, GLYCOPYRROLATE
 * RANBAXY LABORATORIES INC
 EURAX, CROTAMITON
 EXELDERM, SULCONAZOLE NITRATE
 FAMOTIDINE, FAMOTIDINE (OTC)
 HALOG, HALCINONIDE
 KENALOG, TRIAMCINOLONE ACETONIDE
 LAC-HYDRIN, AMMONIUM LACTATE
 RISPERIDONE, RISPERIDONE
 ULTRAVATE, HALOBETASOL PROPIONATE
 WESTCORT, HYDROCORTISONE VALERATE
 * RANBAXY LABORATORIES LTD
 ACYCLOVIR, ACYCLOVIR
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 AMOXICILLIN, AMOXICILLIN
 BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 CARVEDILOL, CARVEDILOL
 CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
 CEFPROZIL, CEFPROZIL
 CEFUROXIME AXETIL, CEFUROXIME AXETIL
 CEPHALEXIN, CEPHALEXIN
 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)
 CLARITHROMYCIN, CLARITHROMYCIN
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLORAZEPATE DIPOTASSUM, CLORAZEPATE DIPOTASSUM
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DISPERMOX, AMOXICILLIN
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 DOXYCYCLINE, DOXYCYCLINE
 FENOFLIBRATE, FENOFLIBRATE
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 GANCICLOVIR, GANCICLOVIR
 GLIMEPIRIDE, GLIMEPIRIDE
 LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
 LORATADINE REDIDOSE, LORATADINE (OTC)
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
 OFLOXACIN, OFLOXACIN
 ONDANSETRON, ONDANSETRON
 OXCARBAZEPINE, OXCARBAZEPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * RANBAXY LABORATORIES LTD
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
 - QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 - RAMIPRIL, RAMIPRIL
 - RANICLOR, CEFACLOR
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RISPERIDONE, RISPERIDONE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - SIMVASTATIN, SIMVASTATIN
 - SOTRET, ISOTRETINOIN
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 - TOPIRAMATE, TOPIRAMATE
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - ZIDOVUDINE, ZIDOVUDINE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- * RANBAXY PHARMACEUTICALS INC
 - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
 - AMOXICILLIN, AMOXICILLIN
 - AVENTYL HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 - CEFACLOR, CEFACLOR
 - CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LISINOPRIL, LISINOPRIL
 - LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - PENTAZOCINE AND NALOXONE HYDROCHLORIDES, NALOXONE HYDROCHLORIDE
 - RIOMET, METFORMIN HYDROCHLORIDE
 - SOTRET, ISOTRETINOIN

RANBAXY LABS LTD

- * RANBAXY LABORATORIES LIMITED
 - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
- * RANBAXY LABORATORIES LTD
 - ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

RARE DIS

- * RARE DISEASE THERAPEUTICS INC
 - ORFADIN, NITISINONE

RARE DIS THERAP

- * RARE DISEASE THERAPEUTICS INC
 - CYSTADANE, BETAINE HYDROCHLORIDE

RECIP

- * RECIP AB
 - THYROSAFE, POTASSIUM IODIDE (OTC)

RECKITT BENCKISER

- * RECKITT BENCKISER
 - DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
- * RECKITT BENCKISER INC
 - MUCINEX D, GUAIFENESIN (OTC)
 - MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
 - MUCINEX, GUAIFENESIN (OTC)
- * RECKITT BENCKISER PHARMACEUTICALS INC
 - BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
 - SUBOXONE, BUPRENORPHINE
 - SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
 - SUBUTEX, BUPRENORPHINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** R **

REDDYS

- * DOCTOR REDDYS LABORATORIES LTD
DESLORATADINE, DESLORATADINE

RHODES PHARMS

- * RHODES PHARMACEUTICALS LP
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

RHODIA

- * RHODIA LTD
ISOFLURANE, ISOFLURANE

RITEDOSE CORP

- * THE RITEDOSE CORP
ALBUTEROL SULFATE, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

ROCHE

- * HOFFMANN LA ROCHE INC
BONIVA, IBANDRONATE SODIUM
COPEGUS, RIBAVIRIN
FUZEON, ENFUVIRTIDE
INVIRASE, SAQUINAVIR MESYLATE
KLONOPIN, CLONAZEPAM
TAMIFLU, OSeltamivir phosphate
VALIUM, DIAZEPAM

ROCHE PALO

- * ROCHE PALO ALTO LLC
AEROBID, FLUNISOLIDE
ANAPROX DS, NAPROXEN SODIUM
ANAPROX, NAPROXEN SODIUM
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CYTOVENE, GANCICLOVIR SODIUM
EC-NAPROSYN, NAPROXEN
NAPROSYN, NAPROXEN
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE

ROCKWELL MEDCL

- * ROCKWELL MEDICAL TECHNOLOGIES INC
CALCITRIOL, CALCITRIOL

ROMARK

- * ROMARK LABORATORIES
ALINIA, NITAZOXANIDE

ROSEDALE THERAPEUTIC

- * ROSEDALE THERAPEUTICS
ERGOMAR, ERGOTAMINE TARTRATE

ROSS LABS

- * ROSS LABORATORIES DIV ABBOTT LABORATORIES INC
SURVANTA, BERACTANT

ROXANE

- * ROXANE LABORATORIES INC
ACARBOSE, ACARBOSE
ALPRAZOLAM, ALPRAZOLAM
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ANASTROZOLE, ANASTROZOLE
AZATHIOPRINE, AZATHIOPRINE
BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
BICALUTAMIDE, BICALUTAMIDE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CALCITRIOL, CALCITRIOL
CALCIUM ACETATE, CALCIUM ACETATE
CILOSTAZOL, CILOSTAZOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * ROXANE LABORATORIES INC
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLARITHROMYCIN, CLARITHROMYCIN
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - CODEINE SULFATE, CODEINE SULFATE
 - CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 - DEXAMETHASONE INTENSOL, DEXAMETHASONE
 - DEXAMETHASONE, DEXAMETHASONE
 - DIAZEPAM INTENSOL, DIAZEPAM
 - DIAZEPAM, DIAZEPAM
 - DIGOXIN, DIGOXIN
 - DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
 - DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - DOXERCALCIFEROL, DOXERCALCIFEROL
 - EXEMESTANE, EXEMESTANE
 - FAMCICLOVIR, FAMCICLOVIR
 - FLECAINIDE ACETATE, FLECAINIDE ACETATE
 - FLUCONAZOLE, FLUCONAZOLE
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - FUROSEMIDE, FUROSEMIDE
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
 - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
 - IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE
 - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 - LACTULOSE, LACTULOSE
 - LETROZOLE, LETROZOLE
 - LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - LITHIUM CITRATE, LITHIUM CITRATE
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - LORAZEPAM INTENSOL, LORAZEPAM
 - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 - MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
 - MEGESTROL ACETATE, MEGESTROL ACETATE
 - MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 - MERCAPTOPURINE, MERCAPTOPURINE
 - METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
 - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
 - METHOTREXATE SODIUM, METHOTREXATE SODIUM
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - MORPHINE SULFATE, MORPHINE SULFATE
 - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
 - NAPROXEN, NAPROXEN
 - NARatriptan, NARatriptan HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXCARBAZEPINE, OXCARBAZEPINE
 - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
 - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
 - PREDNISONE INTENSOL, PREDNISONE
 - PREDNISONE, PREDNISONE
 - PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
 - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 - PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
 - RAMIPRIL, RAMIPRIL
 - RISPERIDONE, RISPERIDONE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** R ****

- * ROXANE LABORATORIES INC
 - ROXICET 5/500, ACETAMINOPHEN
 - ROXICET, ACETAMINOPHEN
 - ROXILOX, ACETAMINOPHEN
 - SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
 - TORSEMIDE, TORSEMIDE
 - TRIAZOLAM, TRIAZOLAM
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - ZALEPLON, ZALEPLON
 - ZIDOVUDINE, ZIDOVUDINE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

SAGE PRODS

- * SAGE PRODUCTS INC
 - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SAGENT PHARMS

- * SAGENT PHARMACEUTICALS INC
 - CALCITRIOL, CALCITRIOL
 - HALOPERIDOL, HALOPERIDOL LACTATE
 - HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
 - HEPARIN SODIUM, HEPARIN SODIUM
 - LEVOFLOXACIN, LEVOFLOXACIN
 - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

SAGENT STRIDES

- * SAGENT STRIDES LLC
 - AZITHROMYCIN, AZITHROMYCIN
 - BACITRACIN, BACITRACIN
 - BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 - BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 - MESNA, MESNA
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
 - MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - POLYMYCIN B SULFATE, POLYMYXIN B SULFATE
 - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

SALIX PHARMS

- * SALIX PHARMACEUTICALS INC
 - ANUSOL HC, HYDROCORTISONE
 - APRISO, MESALAMINE
 - COLAZAL, BALSALAZIDE DISODIUM
 - DIURIL, CHLOROTHIAZIDE
 - METOZOLOV ODT, METOCLOPRAMIDE HYDROCHLORIDE
 - MOVIPREP, ASCORBIC ACID
 - OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
 - PEPCID, FAMOTIDINE
 - VISICOL, SODIUM PHOSPHATE, DIBASIC ANHYDROUS
 - XIFAXAN, RIFAXIMIN

SAMSON MEDCL

- * SAMSON MEDICAL TECHNOLOGIES LLC
 - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 - CEFUROXIME SODIUM IN PLASTIC CONTAINER, CEFUROXIME SODIUM

SANDOZ

- * SANDOZ
 - DOCETAXEL, DOCETAXEL
- * SANDOZ CANADA INC
 - ANECTINE, SUCCINYLCHOLINE CHLORIDE
 - ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
 - BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SANDOZ CANADA INC
COSYNTROPIN, COSYNTROPIN
DIGOXIN, DIGOXIN
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
ESTRADIOL VALERATE, ESTRADIOL VALERATE
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FLUMAZENIL, FLUMAZENIL
GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INFUVITE PEDIATRIC, ASCORBIC ACID
ISONIAZID, ISONIAZID
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
METHYLprednisolone ACETATE, METHYLprednisolone ACETATE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
REGONOL, PYRIDOSTIGMINE BROMIDE
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
- * SANDOZ INC
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
ALPRAZOLAM, ALPRAZOLAM
ALTAVERA, ETHINYLN ESTRADOL
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
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANASTROZOLE, ANASTROZOLE
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
ATENOLOL, ATENOLOL
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
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
CAPTOPRIL, CAPTOPRIL
CARBOPLATIN, CARBOPLATIN
CARISOPRODOL AND ASPIRIN, ASPIRIN
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CARVEDILOL, CARVEDILOL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFDINIR, CEFDINIR
CEFEPEMINE HYDROCHLORIDE, CEFEPEMINE HYDROCHLORIDE
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFTRIAXONE, CEFTRIAXONE SODIUM
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SANDOZ INC
 - CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 - CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
 - CHOLESTYRAMINE, CHOLESTYRAMINE
 - CILOSTAZOL, CILOSTAZOL
 - CIMETIDINE, CIMETIDINE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLARITHROMYCIN, CLARITHROMYCIN
 - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
 - CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 - CLONAZEPAM, CLONAZEPAM
 - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 - CYCLOSPORINE, CYCLOSPORINE
 - DESIPIRAMINE HYDROCHLORIDE, DESIPIRAMINE HYDROCHLORIDE
 - DESLORATADINE, DESLORATADINE
 - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
 - DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
 - DOXYCYCLINE, DOXYCYCLINE
 - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 - ENALAPRIL MALEATE, ENALAPRIL MALEATE
 - ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 - EPLERENONE, EPLERENONE
 - ETODOLAC, ETODOLAC
 - FAMOTIDINE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 - FLUOROURACIL, FLUOROURACIL
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 - FLUTAMIDE, FLUTAMIDE
 - FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 - FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 - FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 - FUROSEMIDE, FUROSEMIDE
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 - GLIPIZIDE, GLIPIZIDE
 - HALOPERIDOL, HALOPERIDOL
 - HEPARIN SODIUM, HEPARIN SODIUM
 - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 - HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 - IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 - IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 - INDAPAMIDE, INDAPAMIDE
 - INDOMETHACIN, INDOMETHACIN
 - INTROVALE, ETHINYLMESTRADIOL
 - INVAGESIC FORTE, ASPIRIN
 - INVAGESIC, ASPIRIN
 - ISONIAZID, ISONIAZID
 - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 - ITRACONAZOLE, ITRACONAZOLE
 - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 - LAMOTRIGINE, LAMOTRIGINE
 - LANSOPRAZOLE, LANSOPRAZOLE
 - LEFLUNOMIDE, LEFLUNOMIDE
 - LEUPROLIDE ACETATE, LEUPROLIDE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SANDOZ INC
LEVETIRACETAM, LEVETIRACETAM
LEVOFLOXACIN, LEVOFLOXACIN
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LOCHOLEST LIGHT, CHOLESTYRAMINE
LOCHOLEST, CHOLESTYRAMINE
LONOX, ATROPINE SULFATE
LORATADINE, LORATADINE (OTC)
LORAZEPAM, LORAZEPAM
LORYNA, DROSPIRENONE
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
LOVASTATIN, LOVASTATIN
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEROPENEM, MEROPENEM
METAXALONE, METAXALONE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHAZOLAMIDE, METHAZOLAMIDE
METHIMAZOLE, METHIMAZOLE
METHOCARBAMOL, METHOCARBAMOL
METHYL PREDNISOLONE, METHYL PREDNISOLONE
METOLAZONE, METOLAZONE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NABUMETONE, NABUMETONE
NADOLOL, NADOLOL
NAFCILLIN SODIUM, NAFCILLIN SODIUM
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NAPROXEN, NAPROXEN
NARatriptan, NARatriptan HYDROCHLORIDE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NIZATIDINE, NIZATIDINE
OFLOXACIN, OFLOXACIN
OMEPRAZOLE, OMEPRAZOLE
OMNITROPE, SOMATROPIN RECOMBINANT
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
ONDANSETRON, ONDANSETRON
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
OXACILLIN SODIUM, OXACILLIN SODIUM
OXALIPLATIN, OXALIPLATIN
OXANDROLONE, OXANDROLONE
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
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
QUINIDINE SULFATE, QUINIDINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* SANDOZ INC
 RAMIPRIL, RAMIPRIL
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 RESERPINE, RESERPINE
 RIBAVIRIN, RIBAVIRIN
 RIFAMPIN, RIFAMPIN
 RISPERIDONE, RISPERIDONE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFADIAZINE, SULFADIAZINE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
 SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 SYEDA, DROSPIRENONE
 TACROLIMUS, TACROLIMUS
 TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 TEMAZEPAM, TEMAZEPAM
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 THIOTHIXENE, THIOTHIXENE
 TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 ZONISAMIDE, ZONISAMIDE

SANDOZ CANADA INC

* SANDOZ CANADA INC
 PARICALCITOL, PARICALCITOL

SANDOZ INC

* SANDOZ INC
 DOCETAXEL, DOCETAXEL
 ENOXAPARIN SODIUM, ENOXAPARIN SODIUM
 VORICONAZOLE, VORICONAZOLE

SANOHEMIA CORP USA

* SANOHEMIA CORP USA
 SCANLUX-300, IOPAMIDOL
 SCANLUX-370, IOPAMIDOL

SANOFI AVENTIS

* SANOFI AVENTIS
 AVALIDE, HYDROCHLOROTHIAZIDE

SANOFI AVENTIS US

* SANOFI AVENTIS US INC
 JEVTONA 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
 ANZEMET, DOLASSETRON MESYLATE
 APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
 APIDRA, INSULIN GLULISINE RECOMBINANT
 ARALEN, CHLOROQUINE PHOSPHATE
 ARAVA, LEFLUNOMIDE
 AVAPRO, IRBESARTAN
 BENZAACLIN, BENZOYL PEROXIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SANOFI AVENTIS US LLC
 - BENZAMYCIN PAK, BENZOYL PEROXIDE
 - BENZAMYCIN, BENZOYL PEROXIDE
 - CANTIL, MEPENZOLATE BROMIDE
 - CARAC, FLUOROURACIL
 - CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
 - CHILDREN'S ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
 - CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
 - CLAFORAN, CEFOTAXIME SODIUM
 - CLOMID, CLOMIPHENE CITRATE
 - DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 - DDAVP, DESMOPRESSIN ACETATE
 - DEMEROL, MEPERIDINE HYDROCHLORIDE
 - DERMATOP E EMOLLIENT, PREDNICARBATE
 - DERMATOP, PREDNICARBATE
 - DIABETA, GLYBURIDE
 - DRISDOL, ERGOCALCIFEROL
 - ELOXATIN, OXALIPLATIN
 - FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
 - GAVISCON, ALUMINUM HYDROXIDE (OTC)
 - HIPREX, METHENAMINE HIPPURATE
 - KAYEXALATE, SODIUM POLYSTYRENE SULFONATE
 - KETEK, TELITHROMYCIN
 - KLARON, SULFACETAMIDE SODIUM
 - LANTUS, INSULIN GLARGINE RECOMBINANT
 - LASIX, FUROSEMIDE
 - LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 - LOVENOX, ENOXAPARIN SODIUM
 - MULTAQ, DRONEDARONE HYDROCHLORIDE
 - MYTELASE, AMBENONIUM CHLORIDE
 - NASACORT AQ, TRIAMCINOLONE ACETONIDE
 - NICODERM CQ, NICOTINE (OTC)
 - NILANDRON, NILUTAMIDE
 - NORITATE, METRONIDAZOLE
 - NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
 - OFORTA, FLUDARABINE PHOSPHATE
 - PENLAC, CICLOPIROX
 - PHISOHEX, HEXACHLOROPHENE
 - PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
 - PLAVIX, CLOPIDOGREL BISULFATE
 - PRIFTIN, RIFAPENTINE
 - PRIMAQUINE, PRIMAQUINE PHOSPHATE
 - RIFADIN, RIFAMPIN
 - RIFAMATE, ISONIAZID
 - RIFATER, ISONIAZID
 - RILUTEK, RILUZOLE
 - SKELID, TILUDRONATE DISODIUM
 - TAXOTERE, DOCETAXEL
 - TRENTAL, PENTOXIFYLLINE
 - UROXATRAL, ALFUZOSIN HYDROCHLORIDE

SANTARUS

- * SANTARUS INC
 - FENOGLIIDE, FENOFIBRATE
 - GLUMETZA, METFORMIN HYDROCHLORIDE
 - MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE, MAGNESIUM HYDROXIDE
 - ZEGERID, OMEPRAZOLE

SANTEN

- * SANTEN INC
 - ALAMAST, PEMIROLAST POTASSIUM
 - IQUIX, LEVOFLOXACIN
 - QUIXIN, LEVOFLOXACIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******SANTEN OY**

- * SANTEN OY
BETIMOL, TIMOLOL

SANTOS BIOTECH

- * SANTOS BIOTECH INDUSTRIES INC
ANASTROZOLE, ANASTROZOLE

SAVAGE LABS

- * SAVAGE LABORATORIES INC DIV ALTANA INC
KAON CL-10, POTASSIUM CHLORIDE

SAVENT PHARMS

- * SAVIENT PHARMACEUTICALS INC
OXANDRIN, OXANDROLONE

SB PHARMCO

- * SB PHARMCO PUERTO RICO INC
AVANDAMET, METFORMIN HYDROCHLORIDE
AVANDARYL, GLIMEPIRIDE
AVANDIA, ROSIGLITAZONE MALEATE
COREG CR, CARVEDILOL PHOSPHATE

SCHERING

- * MERCK, INC.
CLARINEX D 24 HOUR, DESLORATADINE
- * SCHERING
CLARINEX-D 12 HOUR, DESLORATADINE
- * SCHERING CORP
ASMANEX TWISTHALER, MOMETASONE FUROATE
CLARINEX, DESLORATADINE
DIPROLENE AF, BETAMETHASONE DIPROPIONATE
DULERA, FORMOTEROL FUMARATE
GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
INTEGRILIN, EPTIFIBATIDE
LOTRISONE, BETAMETHASONE DIPROPIONATE
NOXAFL, POSACONAZOLE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
REBETOL, RIBAVIRIN
TEMODAR, TEMOZOLOMIDE
VICTRELIS, BOCEPREVIR
ZEMURON, ROCURONIUM BROMIDE
- * SCHERING CORP SUB SCHERING PLOUGH CORP
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELESTONE, BETAMETHASONE
DIPROLENE, BETAMETHASONE DIPROPIONATE
ELOCON, MOMETASONE FUROATE

SCHERING PLOUGH

- * SCHERING PLOUGH CORP
CLARINEX, DESLORATADINE
- * SCHERING PLOUGH HEALTHCARE PRODUCTS INC
AFRINOL, PSEUDOEPHEDRINE SULFATE (OTC)
CHILDREN'S CLARITIN, LORATADINE (OTC)
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)
CLARITIN HIVES RELIEF REDTAB, LORATADINE (OTC)
CLARITIN HIVES RELIEF, LORATADINE (OTC)
CLARITIN REDTABS, LORATADINE (OTC)
CLARITIN, LORATADINE (OTC)
CLARITIN-D 24 HOUR, LORATADINE (OTC)
CLARITIN-D, LORATADINE (OTC)
DISOPHROL, DEXBROMPHENIRAMINE MALEATE (OTC)
DRIXORAL PLUS, ACETAMINOPHEN (OTC)
DRIXORAL, DEXBROMPHENIRAMINE MALEATE (OTC)
GYNE-LOTRIMIN 3 COMBINATION PACK, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN 3, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN COMBINATION PACK, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN, CLOTRIMAZOLE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SCHERING PLOUGH HEALTHCARE PRODUCTS INC
LOTTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)
NASONEX, MOMETASONE FUROATE MONOHYDRATE
OCUCLEAR, OXYMETAZOLINE HYDROCHLORIDE (OTC)
SHADE UVAGUARD, AVOBENZONE (OTC)

SCHERING PLOUGH RES

- * SCHERING PLOUGH RESEARCH INSTITUTE
LOTRISONE, BETAMETHASONE DIPROPIONATE
REBETOL, RIBAVIRIN

SCHWARZ PHARMA

- * SCHWARZ PHARMA AG
EDEX, ALPROSTADIL
- * SCHWARZ PHARMA INC
DILATRATE-SR, ISOSORBIDE DINITRATE
LEVATOL, PENBUTOLOL SULFATE
NIRAVAM, ALPRAZOLAM
PARCOPA, CARBIDOPA
ROBAXIN, METHOCARBAMOL
ROBAXIN-750, METHOCARBAMOL

SCIOS

- * SCIOS INC
NATRECOR, NESIRITIDE RECOMBINANT

SECAN PHARMS

- * SECAN PHARMACEUTICALS INC
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

SEPTODONT

- * SEPTODONT INC
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
OCTOCAIN, EPINEPHRINE

SEPTODONT HOLDING

- * SEPTODONT HOLDING SAS
ORaverse, PHENTOLAMINE MESYLATE

SEPTODONT INC

- * SEPTODONT INC
PRILOCaine HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
PRILOCaine HYDROCHLORIDE, PRILOCaine HYDROCHLORIDE

SHASUN USA

- * SHASUN USA INC
IBUPROFEN, IBUPROFEN

SHIONOGI INC

- * SHIONOGI INC
ADRENAClick, EPINEPHRINE
CUVPOSA, GLYCOPYRROLATE
FURADANTIN, NITROFURANTOIN
KAPVAY, CLONIDINE HYDROCHLORIDE
KEFLEX, CEPHALEXIN
MOXATAG, AMOXICILLIN
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE
ORAPRED, PREDNISOLONE SODIUM PHOSPHATE
PONSTEL, MEFENAMIC ACID
ROBINUL FORTE, GLYCOPYRROLATE
ROBINUL, GLYCOPYRROLATE
SULAR, NISOLDIPINE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TWINJECT 0.15, EPINEPHRINE
TWINJECT 0.3, EPINEPHRINE
ULESFIA, BENZYL ALCOHOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******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
 - AGRYLIN, ANAGRELIDE HYDROCHLORIDE
 - CARBATROL, CARBAMAZEPINE
 - FOSRENOL, LANTHANUM CARBONATE
 - INTUNIV, GUANFACINE HYDROCHLORIDE
 - LIALDA, MESALAMINE
 - PENTASA, MESALAMINE
 - PROAMATINE, MIDODRINE HYDROCHLORIDE
 - SALURON, HYDROFLUMETHIAZIDE

SHIRE DEVELOPMENT

- * SHIRE DEVELOPMENT INC
 - VYVANSE, LISDEXAMFETAMINE Dimesylate

SHIRE HUMAN GENETIC

- * SHIRE HUMAN GENETIC THERAPIES INC
 - VPRIV, VELAGLUCERASE ALFA

SHIRE ORPHAN THERAP

- * SHIRE ORPHAN THERAPIES INC
 - FIRAZYR, ICATIBANT ACETATE

SIGMA TAU

- * SIGMA TAU PHARMACEUTICALS INC
 - ABELCET, AMPHOTERICIN B
 - ADAGEN, PEGADEMASE BOVINE
 - CARNITOR SF, LEVOCARNITINE
 - CARNITOR, LEVOCARNITINE
 - MATULANE, PROCARBAZINE HYDROCHLORIDE

SIGMAPHARM LABS LLC

- * SIGMAPHARM LABORATORIES LLC
 - AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
 - DISULFIRAM, DISULFIRAM
 - ERGOCALCIFEROL, ERGOCALCIFEROL
 - FLUCYTOSINE, FLUCYTOSINE
 - PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE

SILARX

- * SILARX PHARMACEUTICALS INC
 - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 - DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - HALOPERIDOL, HALOPERIDOL LACTATE
 - LEVETIRACETAM, LEVETIRACETAM
 - LORATADINE, LORATADINE (OTC)
 - METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - THEOPHYLLINE, THEOPHYLLINE

SKINMEDICA

- * SKINMEDICA INC
 - VANIQA, EFLORNITHINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ******SKYEPHARMA AG**

- * SKYEPHARMA AG
TRIGLIDE, FENOFLIBRATE

SLATE PHARMS

- * SLATE PHARMACEUTICALS INC
TESTOPEL, TESTOSTERONE

SMITHKLINE BEECHAM

- * SMITHKLINE BEECHAM
LOVASA, OMEGA-3-ACID ETHYL ESTERS
- * SMITHKLINE BEECHAM CORP
LAMICTAL XR, LAMOTRIGINE
- * SMITHKLINE BEECHAM CORP DBA GLAXOSMITHKLINE
ARRANON, NELARABINE
AXID, NIZATIDINE
COREG, CARVEDILOL
HYCAMTIN, TOPOTECAN HYDROCHLORIDE
LAMICTAL ODT, LAMOTRIGINE
REQUIP XL, ROPINIROLE HYDROCHLORIDE
TYKERB, LAPATINIB DITOSYLADE

SOAPCO

- * SOAPCO INC
BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)

SOLCO HLTHCARE

- * SOLCO HEALTHCARE US LLC
LEVETIRACETAM, LEVETIRACETAM

SOMAXON

- * SOMAXON PHARMACEUTICALS INC
SILENOR, DOXEPIN HYDROCHLORIDE

SOMERSET

- * SOMERSET PHARMACEUTICALS INC
ELDEPRYL, SELEGILINE HYDROCHLORIDE
EMSAM, SELEGILINE

SPEAR PHARMS

- * SPEAR PHARMACEUTICALS INC
FLUOROURACIL, FLUOROURACIL
TRETINOIN, TRETINOIN

SPECTRUM PHARMS

- * SPECTRUM PHARMACEUTICALS
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
- * SPECTRUM PHARMACEUTICALS INC
FUSILEV, LEVOLEUCOVORIN CALCIUM

STAND HOMEOPATH

- * STANDARD HOMEOPATHIC CO
IVY BLOCK, BENTOQUATAM (OTC)

STASON

- * STASON INDUSTRIAL CORP
ACYCLOVIR, ACYCLOVIR
CAPTOPRIL, CAPTOPRIL
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

STASON PHARMS

- * STASON PHARMACEUTICALS INC
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

STAT TRADE

- * STAT TRADE INC
NAPRELAN, NAPROXEN SODIUM

STERI PHARMA

- * STERI PHARMA LLC
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFTRIAXONE, CEFTRIAXONE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

* STERI PHARMA LLC
CEFUROXIME SODIUM, CEFUROXIME SODIUM

STEVENS J

* JEROME STEVENS PHARMACEUTICALS INC
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
CEPHALEXIN, CEPHALEXIN
DIGOXIN, DIGOXIN
METHOCARBAMOL AND ASPIRIN, ASPIRIN
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
UNITHROID, LEVOTHYROXINE SODIUM

STI PHARMA LLC

* STI PHARMA LLC
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
DEXAMETHASONE, DEXAMETHASONE
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
TRIACIN-C, CODEINE PHOSPHATE

STIEFEL

* STIEFEL LABORATORIES INC
DUAC, BENZOYL PEROXIDE

STIEFEL GSK

* STIEFEL A GSK CO
VELTIN, CLINDAMYCIN PHOSPHATE

STIEFEL LABS INC

* STIEFEL LABORATORIES INC
EVOCLIN, CLINDAMYCIN PHOSPHATE
EXTINA, KETOCONAZOLE
OLUX E, CLOBETASOL PROPIONATE
OLUX, CLOBETASOL PROPIONATE
ONMEL, ITRACONAZOLE
SORIATANE, ACITRETIN
SORILUX, CALCIPOTRIENE
VERDESO, DESONIDE
VUSION, MICONAZOLE NITRATE

STRIDES ARCOLAB

* STRIDES ARCOLAB LIMITED
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM

STRIDES ARCOLAB LTD

* STRIDES ARCOLAB LTD
ACARBOSE, ACARBOSE
ADENOSINE, ADENOSINE
ERGOCALCIFEROL, ERGOCALCIFEROL
ETOMIDATE, ETOMIDATE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
MELOXICAM, MELOXICAM
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

SUCAMPO PHARMS

* SUCAMPO PHARMACEUTICALS INC
AMITIZA, LUBIPROSTONE

SUMMERS

* SUMMERS LABORATORIES INC
CROTAN, CROTAMITON

SUN PHARM INDs

* SUN PHARMACEUTICAL INDUSTRIES LTD
CARBIDOPA AND LEVODOPA, CARBIDOPA
DESLORATADINE, DESLORATADINE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SUN PHARMACEUTICAL INDUSTRIES LTD
 - EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 - ONDANSETRON, ONDANSETRON
 - OXCARBAZEPINE, OXCARBAZEPINE
 - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE
 - TORSEMIDE, TORSEMIDE

SUN PHARM INDS (IN)

- * SUN PHARMACEUTICAL INDUSTRIES LTD
 - CEPHALEXIN, CEPHALEXIN
 - EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - ORTHO-EST, ESTROPIPATE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - ZONISAMIDE, ZONISAMIDE

SUN PHARM INDS INC

- * SUN PHARMACEUTICAL INDUSTRIES INC
 - BENZONATATE, BENZONATATE
 - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - ERGOCALCIFEROL, ERGOCALCIFEROL
 - GEMFIBROZIL, GEMFIBROZIL
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - LITHIUM CARBONATE, LITHIUM CARBONATE
 - NIMODIPINE, NIMODIPINE
 - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - VALPROIC ACID, VALPROIC ACID
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

SUN PHARM INDS LTD

- * SUN PHARMACEUTICAL INDUSTRIES LTD
 - CARISOPRODOL, CARISOPRODOL
 - GABAPENTIN, GABAPENTIN
 - LETROZOLE, LETROZOLE
 - LEVETIRACETAM, LEVETIRACETAM
 - NARatriptan, NARatriptan
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 - TOPIRAMATE, TOPIRAMATE
- * SUN PHARMACEUTICAL INDUSTRIES LTD.
 - ANASTROZOLE, ANASTROZOLE

SUN PHARMA GLOBAL

- * SUN PHARMA GLOBAL FZE
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - CAFFEINE CITRATE, CAFFEINE CITRATE
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
 - CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DOCEFREZ, DOCTETAXEL
 - FINASTERIDE, FINASTERIDE
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** S ****

- * SUN PHARMA GLOBAL FZE
OXALIPLATIN, OXALIPLATIN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- * SUN PHARMA GLOBAL INC
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALPRAZOLAM, ALPRAZOLAM
AMIFOSTINE, AMIFOSTINE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
BICALUTAMIDE, BICALUTAMIDE
CARBOPLATIN, CARBOPLATIN
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
VECURONIUM BROMIDE, VECURONIUM BROMIDE

SUNOVION

- * SUNOVION PHARMACEUTICALS INC
BROVANA, ARFORMOTEROL TARTRATE
LATUDA, LURASIDONE HYDROCHLORIDE
XOPENEX HFA, LEVALBUTEROL TARTRATE
XOPENEX, LEVALBUTEROL HYDROCHLORIDE

SUNOVION PHARMS INC

- * SUNOVION PHARMACEUTICALS INC
LUNESTA, ESZOPICLONE

SVADS HOLDINGS SA

- * SVADS HOLDINGS SA
IBUPROFEN, IBUPROFEN (OTC)
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

SVC PHARMA

- * SVC PHARMA LP
DRONABINOL, DRONABINOL

SYNCOR PHARMS

- * SYNCOR PHARMACEUTICALS INC
SODIUM IODIDE I 123, SODIUM IODIDE I-123

SYNERX

- * SYNERX PHARMA LLC
ISOSULFAN BLUE, ISOSULFAN BLUE
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE

SYNERX PHARMA

- * SYNERX PHARMA LLC
CICLOPIROX, CICLOPIROX
FOMEPIZOLE, FOMEPIZOLE
MALATHION, MALATHION
TESTOSTERONE CYPTIONATE, TESTOSTERONE CYPTIONATE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

SYNTTHON PHARMS

- * SYNTTHON PHARMACEUTICALS INC
BICALUTAMIDE, BICALUTAMIDE
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TAKEDA GLOBAL

- * TAKEDA GLOBAL RESEARCH DEVELOPMENT CENTER INC
ACTOPLUS MET XR, METFORMIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TAKEDA GLOBAL RESEARCH DEVELOPMENT CENTER INC
ACTOPLUS MET, METFORMIN HYDROCHLORIDE
DUETACT, GLIMEPIRIDE
ROZEREM, RAMELTEON

TAKEDA PHARMS

- * TAKEDA PHARMACEUTICALS NORTH AMERICA INC
DEXILANT, DEXLANSOPRAZOLE
EDARBEL, AZILSARTAN KAMEDOXOMIL
EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
ULORIC, FEBUXOSTAT

TAKEDA PHARMS NA

- * TAKEDA PHARMACEUTICALS NORTH AMERICA INC
ACTOS, PIOGLITAZONE HYDROCHLORIDE
PREVACID, LANSOPRAZOLE
PREVPAC, AMOXICILLIN

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
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
IMIQUIMOD, IMIQUIMOD
KETOCONAZOLE, KETOCONAZOLE
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LORATADINE, LORATADINE (OTC)
MELOXICAM, MELOXICAM
MEPROBAMATE, MEPROBAMATE
METRONIDAZOLE, METRONIDAZOLE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
OXCARBAZEPINE, OXCARBAZEPINE
PHENYTOIN, PHENYTOIN
- * TARO PHARMACEUTICALS INC
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTTRIMAZOLE, CLOTTRIMAZOLE
CLOTTRIMAZOLE, CLOTTRIMAZOLE (OTC)
DESOXIMETASONE, DESOXIMETASONE
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
WARFARIN SODIUM, WARFARIN SODIUM
- * TARO PHARMACEUTICALS USA INC
ACETIC ACID, ACETIC ACID, GLACIAL
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICALS USA INC
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CICLOPIROX, CICLOPIROX
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - CLOBETASOL PROPIONATE (EMOLlient), CLOBETASOL PROPIONATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - DERMABET, BETAMETHASONE VALERATE
 - DESONIDE, DESONIDE
 - DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
 - ECONAZOLE NITRATE, ECONAZOLE NITRATE
 - FLO-PRED, PREDNISOLONE ACETATE
 - FLUOCINOLONE ACTONIDE, FLUOCINOLONE ACTONIDE
 - FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 - FLUOCINONIDE, FLUOCINONIDE
 - FLUOROURACIL, FLUOROURACIL
 - GENTAMICIN SULFATE, GENTAMICIN SULFATE
 - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
 - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 - HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 - HYDROCORTISONE, HYDROCORTISONE
 - KETOZOLE, KETOCONAZOLE
 - LIDOCaine, LIDOCaine
 - LORATADINE, LORATADINE (OTC)
 - MICONazole 3, MICONazole NITRATE (OTC)
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - MUPIROCIN, MUPIROCIN
 - NYSTATIN AND TRIAMCINOLONE ACTONIDE, NYSTATIN
 - NYSTATIN, NYSTATIN
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RISPERIDONE, RISPERIDONE
 - SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)
 - TERCONAZOLE, TERCONAZOLE
 - TERIL, CARBAMAZEPINE
 - TRIAMCINOLONE ACTONIDE, TRIAMCINOLONE ACTONIDE
 - TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
 - U-CORT, HYDROCORTISONE ACETATE
- * TARO PHARMACEUTICALS, INC.
 - DESOXIMETASONE, DESOXIMETASONE

TARO PHARM INDS

- * TARO PHARMACEUTICAL INDUSTRIES LTD
 - AMCINONIDE, AMCINONIDE
 - CARBAMAZEPINE, CARBAMAZEPINE
 - CICLOPIROX, CICLOPIROX
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 - ETODOLAC, ETODOLAC
 - FLUCONAZOLE, FLUCONAZOLE
 - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
 - LAMOTRIGINE, LAMOTRIGINE

TARO PHARMS IRELAND

- * TARO PHARMACEUTICALS IRELAND LTD
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 - STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION

TARO PHARMS NORTH

- * TARO PHARMACEUTICALS NORTH AMERICA INC
 - OVIDE, MALATHION
 - TOPICORT LP, DESOXIMETASONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TARO PHARMACEUTICALS NORTH AMERICA INC
TOPICORT, DESOXIMETASONE

TAYLOR

- * TAYLOR PHARMACEUTICALS
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LORAZEPAM, LORAZEPAM
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
NAPHAZOLINE HYDROCHLORIDE, NAPHAZOLINE HYDROCHLORIDE

TAYLOR PHARMA

- * TAYLOR PHARMACAL CO
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
PROPARACAIN HYDROCHLORIDE, PROPARACAIN HYDROCHLORIDE
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

TEDOR PHARM

- * TEDOR PHARMA INC
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE

TEIKOKU PHARMA USA

- * TEIKOKU PHARMA USA INC
LIDODERM, LIDOCAINE

TERCICA

- * TERCICA INC
INCRELEX, MECASERMIN RECOMBINANT

TEVA

- * TEVA NEUROSCIENCE INC
AZILECT, RASAGILINE MESYLATE
COPAXONE, GLATIRAMER ACETATE
- * 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
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETA-VAL, BETAMETHASONE VALERATE
BICALUTAMIDE, BICALUTAMIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CALCITRIOL, CALCITRIOL
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CAPTOPRIL, CAPTOPRIL
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARVEDILOL, CARVEDILOL
CEFACLOR, CEFACLOR
CEFPROZIL, CEFPROZIL
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME SODIUM, CEFUROXIME SODIUM
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 HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIMETIDINE, CIMETIDINE
CLARITHROMYCIN, CLARITHROMYCIN
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PHARMACEUTICALS USA INC
 - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 - CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 - CLONAZEPAM, CLONAZEPAM
 - CLOTRIMAZOLE, CLOTRIMAZOLE
 - 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 AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 - ENALAPRIL MALEATE, ENALAPRIL MALEATE
 - EPITOL, CARBAMAZEPINE
 - ESTAZOLAM, ESTAZOLAM
 - ETODOLAC, ETODOLAC
 - FAMOTIDINE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FENOFIBRATE (MICRONIZED), FENOFIBRATE
 - FENOFIBRATE, FENOFIBRATE
 - 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
 - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 - 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
 - MESALAMINE, MESALAMINE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - METOLAZONE, METOLAZONE
 - METOPROLOL TARTRATE, METOPROLOL TARTRATE
 - METRONIDAZOLE, METRONIDAZOLE
 - MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 - MIRTAZAPINE, MIRTAZAPINE
 - MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 - MUPIROCIN, MUPIROCIN
 - NABUMETONE, NABUMETONE
 - NAPROXEN SODIUM, NAPROXEN SODIUM
 - NAPROXEN, NAPROXEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PHARMACEUTICALS USA INC
 - NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 - NEOMYCIN SULFATE, NEOMYCIN SULFATE
 - NICARDIPIINE HYDROCHLORIDE, NICARDIPIINE HYDROCHLORIDE
 - NIZATIDINE, NIZATIDINE
 - NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 - NYSTATIN, NYSTATIN
 - OFLOXACIN, OFLOXACIN
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON, ONDANSETRON
 - ORAP, PIMOZIDE
 - OXaprozin, OXaprozin
 - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
 - PENICILLIN-VK, PENICILLIN V POTASSIUM
 - PIROXICAM, PIROXICAM
 - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
 - PRELONE, PREDNISOLONE
 - PURINETHOL, MERCAPTOPURINE
 - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RIBAVIRIN, RIBAVIRIN
 - RISPERIDONE, RISPERIDONE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 - SUCRALFATE, SUCRALFATE
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TOLMETIN SODIUM, TOLMETIN SODIUM
 - TOPIRAMATE, TOPIRAMATE
 - TORSEMIDE, TORSEMIDE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRIACET, TRIAMCINOLONE ACETONIDE
 - TRIMETHOPRIM, TRIMETHOPRIM
 - VENLAFAKINE HYDROCHLORIDE, VENLAFAKINE HYDROCHLORIDE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

TEVA BRANDED PHARM

- * TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
- * TEVA BRANDED PHARMACEUTICAL PRODUCTS R&D INC
 - QVAR 40, BECLOMETHASONE DIPROPIONATE
 - QVAR 80, BECLOMETHASONE DIPROPIONATE

TEVA GLOBAL

- * TEVA GLOBAL RESPIRATORY RESEARCH LLC
 - PROAIR HFA, ALBUTEROL SULFATE
 - PROGLYCEM, DIAZOXIDE

TEVA PARENTERAL

- * TEVA PARENTERAL MEDICINES INC
 - ADENOSINE, ADENOSINE
 - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - ALPROSTADIL, ALPROSTADIL
 - AMIKACIN SULFATE, AMIKACIN SULFATE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - AZITHROMYCYIN, AZITHROMYCYIN
 - BLEOMYCIN SULFATE, BLEOMYCIN SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PARENTERAL MEDICINES INC
BUDESONIDE, BUDESONIDE
CALCITRIOL, CALCITRIOL
CARBOPLATIN, CARBOPLATIN
CEFTRIAXONE, CEFTRIAXONE SODIUM
CIPROFLOXACIN, CIPROFLOXACIN
CISPLATIN, CISPLATIN
CROMOLYN SODIUM, CROMOLYN SODIUM
CYTOSAR-U, CYTARABINE
DACARBAZINE, DACARBAZINE
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
ENALAPRILAT, ENALAPRILAT
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
ETOPOSIDE, ETOPOSIDE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FLUMAZENIL, FLUMAZENIL
FLUOROURACIL, FLUOROURACIL
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
IFOSFAMIDE, IFOSFAMIDE
IFOSFAMIDE/MESNA KIT, IFOSFAMIDE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
LEVOCARNITINE, LEVOCARNITINE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MESNA, MESNA
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
OXALIPLATIN, OXALIPLATIN
OXYTOCIN, OXYTOCIN
PACLITAXEL, PACLITAXEL
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPOFOL, PROPOFOL
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
ZANOSAR, STREPTOZOZINCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ******TEVA PARENTL**

* TEVA PARENTAL MEDICINE INC
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE

TEVA PHARMS

* TEVA PHARMACEUTICALS USA
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
ALPRAZOLAM, ALPRAZOLAM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
ANASTROZOLE, ANASTROZOLE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
AZITHROMYCIN, AZITHROMYCIN
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFDINIR, CEFDINIR
CEFPROZIL, CEFPROZIL
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
CICLOPIROX, CICLOPIROX
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
DOXE PIN HYDROCHLORIDE, DOXE PIN HYDROCHLORIDE
ETHOSUXIMIDE, ETHOSUXIMIDE
FAMCICLOVIR, FAMCICLOVIR
FENTANYL-100, FENTANYL
FENTANYL-25, FENTANYL
FENTANYL-50, FENTANYL
FENTANYL-75, FENTANYL
GABAPENTIN, GABAPENTIN
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL LACTATE
HYDROCORTISONE, HYDROCORTISONE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
LANSOPRAZOLE, LANSOPRAZOLE
LEFLUNOMIDE, LEFLUNOMIDE
LETROZOLE, LETROZOLE
LEVETIRACETAM, LEVETIRACETAM
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MEBENDAZOLE, MEBENDAZOLE
MEGESTROL ACETATE, MEGESTROL ACETATE
MELOXICAM, MELOXICAM
METHAZOLAMIDE, METHAZOLAMIDE
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
NARatriptan, NARatriptan HYDROCHLORIDE
NATEGLINIDE, NATEGLINIDE
OLANZAPINE, OLANZAPINE
ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
OXCARBAZEPINE, OXCARBAZEPINE
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
QUINIDINE SULFATE, QUINIDINE SULFATE
RAMIPRIL, RAMIPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TEVA PHARMACEUTICALS USA
 - RISEDRONATE SODIUM, RISEDRONATE SODIUM
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 - TRANDOLAPRIL, TRANDOLAPRIL
 - URSODIOL, URSDIOL
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - VALPROIC ACID, VALPROIC ACID
 - VANDAZOLE, METRONIDAZOLE
 - ZALEPLON, ZALEPLON

TEVA PHARMS USA

- * TEVA PHARMACEUTICALS USA INC
 - IMIQUIMOD, IMIQUIMOD

TEVA WOMENS

- * TEVA WOMENS HEALTH INC
 - CENESTIN, ESTROGENS, CONJUGATED SYNTHETIC A
 - ENJUVIA, ESTROGENS, CONJUGATED SYNTHETIC B
 - LOSEASONIQUE, ETHINYL ESTRADIOL
 - MIRCETTE, DESOGESTREL
 - PLAN B, LEVONORGESTREL
 - PLAN B, LEVONORGESTREL (OTC)
 - PREFEST, ESTRADIOL
 - SEASONIQUE, ETHINYL ESTRADIOL
 - SYNTHETIC CONJUGATED ESTROGENS A, ESTROGENS, CONJUGATED SYNTHETIC A

THE PHARMA NETWORK

- * THE PHARMA NETWORK LLC
 - BENZONATATE, BENZONATATE

THE PHARMANETWORK

- * THE PHARMANETWORK LLC
 - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE

ATHERAKOS

- * THERAKOS INC
 - UVADEX, METHOXSALEN

ATHERATECHNOLOGIES

- * THERATECHNOLOGIES INC
 - EGRIFTA, TESAMORELIN ACETATE

THERAVANCE INC

- * THERAVANCE INC
 - VIBATIV, TELAVANCIN HYDROCHLORIDE

THREE RIVERS PHARMS

- * THREE RIVERS PHARMACEUTICALS LLC
 - ANASTROZOLE, ANASTROZOLE
 - RIBASPHERE, RIBAVIRIN
 - RIBAVIRIN, RIBAVIRIN
 - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

TIBOTEC

- * TIBOTEC INC
 - EDURANT, RILPIVIRINE HYDROCHLORIDE
 - INTELENCE, ETRAVIRINE
 - PREZISTA, DARUNAVIR ETHANOLATE

TOLMAR

- * TOLMAR INC
 - ATRIDOX, DOXYCYCLINE HYCLATE
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CALCIPOTRIENE, CALCIPOTRIENE
 - CICLOPIROX, CICLOPIROX
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - IMIQUIMOD, IMIQUIMOD

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TOLMAR INC
 - KETOCONAZOLE, KETOCONAZOLE
 - LEVETIRACETAM, LEVETIRACETAM
 - LIDOCaine AND PRilocaine, LIDOCaine
 - METRONIDAZOLE, METRONIDAZOLE
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

TOLMAR THERAP

- * TOLMAR THERAPEUTICS INC
 - ELIGARD, LEUPROLIDE ACETATE

TOPOTARGET

- * TOPOTARGET AS
 - TOTECT, DEXRAZOXANE HYDROCHLORIDE

TORPHARM

- * TORPHARM INC
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

TORRENT PHARM

- * TORRENT PHARMA INC
 - METFORMIN HYDROCHLORIDE, METFORMIN 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
 - FELODIPINE, FELODIPINE

TRIAx PHARMS LLC

- * TRIAX PHARMACEUTICALS LLC
 - LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
 - LOCOID, HYDROCORTISONE BUTYRATE
 - MINOCIN, MINOCYCLINE HYDROCHLORIDE
 - TRETINOIN, TRETINOIN

TRIS PHARMA INC

- * TRIS PHARMA INC
 - HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
 - IBUPROFEN, IBUPROFEN (OTC)
 - LEVETIRACETAM, LEVETIRACETAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** T ****

- * TRIS PHARMA INC
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

UCB INC

- * UCB INC
CO-GESIC, ACETAMINOPHEN
KEPPRA XR, LEVETIRACETAM
KEPPRA, LEVETIRACETAM
LORTAB, ACETAMINOPHEN
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
METADATE ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
MONOKET, ISOSORBIDE MONONITRATE
PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
SEMPREX-D, ACRIVASTINE
THEO-24, THEOPHYLLINE
TUSSIONEX PENNKinetic, CHLORPHENIRAMINE POLISTIREX
UNIRETIC, HYDROCHLOROTHIAZIDE
UNIVASC, MOEXIPRIL HYDROCHLORIDE
VIMPAT, LACOSAMIDE
XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE
ZAROXOLYN, METOLAZONE

UCYCLYD

- * UCYCLYD PHARMA INC
AMMONUL, SODIUM BENZOATE

UDL LABS

- * UDL LABORATORIES
SULFAMYLYON, MAFENIDE ACETATE

ULURU

- * ULURU INC
APHTHASOL, AMLEXANOX

UNICHEM

- * UNICHEM LABORATORIES LTD
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
MELOXICAM, MELOXICAM
TOPIRAMATE, TOPIRAMATE
ZALEPLON, ZALEPLON

UNICHEM LABS LTD

- * UNICHEM LABORATORIES LIMITED
DIVALPROEX SODIUM, DIVALPROEX SODIUM
- * UNICHEM LABORATORIES LTD
LAMOTRIGINE, LAMOTRIGINE

UNICHEM PHARMS (USA)

- * UNICHEM PHARMACEUTICALS (USA) INC
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE

UNIQUE PHARM LABS

- * UNIQUE PHARMACEUTICAL LABORATORIES
ATENOLOL, ATENOLOL
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
FLUCONAZOLE, FLUCONAZOLE

UNITED GUARDIAN

- * UNITED GUARDIAN INC
RENACIDIN, CITRIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** U ******UNITED THERAP**

* UNITED THERAPEUTICS CORP
 REMODULIN, TREPROSTINIL SODIUM
 TYVASO, TREPROSTINIL SODIUM

UPSHER SMITH

* UPSHER SMITH LABORATORIES INC
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 DIVALPROEX SODIUM, DIVALPROEX SODIUM
 DIVIGEL, ESTRADIOL
 FORTICAL, CALCITONIN SALMON RECOMBINANT
 KLOR-CON M10, POTASSIUM CHLORIDE
 KLOR-CON M15, POTASSIUM CHLORIDE
 KLOR-CON M20, POTASSIUM CHLORIDE
 LAMOTRIGINE, LAMOTRIGINE
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 NIACOR, NIACIN
 NYSTATIN, NYSTATIN
 ORVATEN, MIDODRINE HYDROCHLORIDE
 OXANDROLONE, OXANDROLONE
 PACERONE, AMIODARONE HYDROCHLORIDE
 PENTOXIL, PENTOXIFYLLINE
 PREVALITE, CHOLESTYRAMINE
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 SORINE, SOTALOL HYDROCHLORIDE
 TOPIRAMATE, TOPIRAMATE
 ZALEPLON, ZALEPLON

UPSHER-SMITH LABS

* UPSHER-SMITH LABORATORIES INC
 KLOR-CON, POTASSIUM CHLORIDE

US WORLDMEDS

* US WORLDMEDS LLC
 APOKYN, APOMORPHINE HYDROCHLORIDE
 DANTROLENE SODIUM, DANTROLENE SODIUM

USL PHARMA

* USL PHARMA INC
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 BACLOFEN, BACLOFEN
 BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 ESTRADIOL, ESTRADIOL
 FLUOXYMESTERONE, FLUOXYMESTERONE
 JANTOVEN, WARFARIN SODIUM
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

VALEANT

* VALEANT PHARMACEUTICALS INTERNATIONAL
 ANCOPON, FLUCYTOSINE
 BONTRIL PDM, PHENDIMETRAZINE TARTRATE
 BONTRIL, PHENDIMETRAZINE TARTRATE
 CAPITAL AND CODEINE, ACETAMINOPHEN
 D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
 DIASTAT ACUDIAL, DIAZEPAM
 DIASTAT, DIAZEPAM
 MIGRAL, DIHYDROERGOTAMINE MESYLATE
 MOTOFEN, ATROPINE SULFATE
 MYSOLINE, PRIMIDONE
 PHRENILIN FORTE, ACETAMINOPHEN
 PHRENILIN WITH CAFFEINE AND CODEINE, ACETAMINOPHEN
 PHRENILIN, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** V **

VALEANT INTL

- * VALEANT INTERNATIONAL BARBADOS SRL
 - ATIVAN, LORAZEPAM
 - CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
 - CARDIZEM, DILTIAZEM HYDROCHLORIDE
 - DICLOFENAC SODIUM, DICLOFENAC SODIUM
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - ELIDEL, PIMECROLIMUS
 - ISORDIL, ISOSORBIDE DINITRATE
 - NIFEDIPINE, NIFEDIPINE
 - PENTOXIFYLLINE, PENTOXIFYLLINE
 - TIAZAC, DILTIAZEM HYDROCHLORIDE
 - ULTRAM ER, TRAMADOL HYDROCHLORIDE
 - VASERETIC, ENALAPRIL MALEATE
 - VASOTEC, ENALAPRIL MALEATE
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 - WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
 - XERESE, ACYCLOVIR
 - ZOVIRAX, ACYCLOVIR
- * VALEANT INTERNATIONAL SRL
 - APLENZIN, BUPROPION HYDROBROMIDE
 - RETIN-A MICRO, TRETINOIN
 - XENAZINE, TETRABENAZINE

VALEANT PHARM INTL

- * VALEANT PHARMACEUTICALS INTERNATIONAL
 - 8-MOP, METHOXSALEN
 - ANDROID 10, METHYLTESTOSTERONE
 - ANDROID 25, METHYLTESTOSTERONE
 - EFUDEX, FLUOURACIL
 - LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
 - LIMBITROL DS, AMITRIPTYLINE HYDROCHLORIDE
 - LIMBITROL, AMITRIPTYLINE HYDROCHLORIDE
 - MESTINON, PYRIDOSTIGMINE BROMIDE
 - OXSORALEN, METHOXSALEN
 - OXSORALEN-ULTRA, METHOXSALEN
 - TASMAR, TOLCAPONE
 - TENSILON PRESERVATIVE FREE, EDROPHONIUM CHLORIDE
 - TENSILON, EDROPHONIUM CHLORIDE
 - TESTRED, METHYLTESTOSTERONE
 - VIRAZOLE, RIBAVIRIN
 - ZELAPAR, SELEGILINE HYDROCHLORIDE

VALEANT PHARMS

- * VALEANT PHARMACEUTICALS NORTH AMERICA
 - POTIGA, EZOGABINE

VALIDUS PHARMS

- * VALIDUS PHARMACEUTICALS LLC
 - ROCALTROL, CALCITRIOL

VALIDUS PHARMS INC

- * VALIDUS PHARMACEUTICALS INC
 - EQUETRO, CARBAMAZEPINE
 - MARPLAN, ISOCARBOXAZID

VEROSCIENCE

- * VERO SCIENCE LLC
 - CYCLOSET, BROMOCRIPTINE MESYLATE

VERSAPHARM

- * VERSAPHARM INC
 - CICLOPIROX, CICLOPIROX
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
 - DESOXIMETASONE, DESOXIMETASONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

- * VERSAPHARM INC
 - ERYTHROMYCIN, ERYTHROMYCIN
 - ETHOSUXIMIDE, ETHOSUXIMIDE
 - RIFAMPIN, RIFAMPIN

VERSAPHARM INC

- * VERSAPHARM INC
 - RIFAMPIN, RIFAMPIN
- * VERSAPHARM INCORPORATED
 - TRANEXAMIC ACID, TRANEXAMIC ACID

VERTEX PHARMS

- * VERTEX PHARMACEUTICALS INC
 - INCIVEK, TELAPREVIR

VICURON

- * VICURON PHARMACEUTICALS INC
 - ERAXIS, ANIDULAFUNGIN

VIIV HLTHCARE

- * VIIV HEALTHCARE CO
 - COMBIVIR, LAMIVUDINE
 - EPIVIR, LAMIVUDINE
 - EPZICOM, ABACAVIR SULFATE
 - LEXIVA, FOSAMPRENAVIR CALCIUM
 - SCRIPTOR, DELAVIRDINE MESYLATE
 - RETROVIR, ZIDOVUDINE
 - SELZENTRY, MARAVIROC
 - TRIZIVIR, ABACAVIR SULFATE
 - ZIAGEN, ABACAVIR SULFATE

VINTAGE

- * VINTAGE PHARMACEUTICALS LLC
 - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 - ACETIC ACID, ACETIC ACID, GLACIAL
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - ALPRAZOLAM, ALPRAZOLAM
 - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
 - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - CYCLAFEM 1/35, ETHINYL ESTRADIOL
 - CYCLAFEM 7/7/7, ETHINYL ESTRADIOL
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - EMOQUETTE, DESOGESTREL
 - FOLIC ACID, FOLIC ACID
 - GILDESS FE 1.5/30, ETHINYL ESTRADIOL
 - GILDESS FE 1/20, ETHINYL ESTRADIOL
 - GLIMEPIRIDE, GLIMEPIRIDE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - GRISEOFULVIN, GRISEOFULVIN
 - HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
 - HYDROCORTISONE, HYDROCORTISONE
 - LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - LISINOPRIL, LISINOPRIL
 - NYSTATIN, NYSTATIN
 - PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
 - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
 - PREDNISOLONE, PREDNISOLONE
 - PREVIFEM, ETHINYL ESTRADIOL
 - PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE PHOSPHATE
 - PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
 - RISPERIDONE, RISPERIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ****

* VINTAGE PHARMACEUTICALS LLC
 SPIRONOLACTONE, SPIRONOLACTONE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 TRI-PREVIFEM, ETHINYLN ESTRADIOL
 VALPROIC ACID, VALPROIC ACID
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

VINTAGE PHARMS

* VINTAGE PHARMACEUTICALS
 LEVONORGESTREL AND ETHINYLN ESTRADIOL, ETHINYLN ESTRADIOL
 ORSYTHIA, ETHINYLN ESTRADIOL
 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
 CLONAZEPAM, CLONAZEPAM
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DEXAMETHASONE, DEXAMETHASONE
 DIAZEPAM, DIAZEPAM
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 FUROSEMIDE, FUROSEMIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 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
 LEVETIRACETAM, LEVETIRACETAM
 LORAZEPAM, LORAZEPAM
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 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
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
 PREDNISONE, PREDNISONE
 PRIMIDONE, PRIMIDONE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
 SULFASALAZINE, SULFASALAZINE
 TORSEMIDE, TORSEMIDE
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

VIROPHARMA

* VIROPHARMA INC
 VANCOGIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** V ******VISTA PHARMS**

- * VISTA PHARMACEUTICALS INC
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

- * VISTAPHARM INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
LACTULOSE, LACTULOSE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NYSTATIN, NYSTATIN
PHENYTOIN, PHENYTOIN

WARNER CHILCOTT

- * WARNER CHILCOTT BERMUDA LTD
CHOLEDYL SA, OXTRIHYLLINE
- * WARNER CHILCOTT CO LLC
ACTONEL, RISEDRONATE SODIUM
ATELVIA, RISEDRONATE SODIUM
- * WARNER CHILCOTT INC
ERYC, ERYTHROMYCIN
ESTRACE, ESTRADIOL
ESTROSTEP FE, ETHINYL ESTRADIOL
FEMCON FE, ETHINYL ESTRADIOL
FEMHRT, ETHINYL ESTRADIOL
FEMTRACE, ESTRADIOL ACETATE
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
OVCON-35, ETHINYL ESTRADIOL
OVCON-50, ETHINYL ESTRADIOL
SARAFEM, FLUOXETINE HYDROCHLORIDE

WARNER CHILCOTT CO

- * WARNER CHILCOTT CO INC
LO LOESTRIN FE, ETHINYL ESTRADIOL

WARNER CHILCOTT LLC

- * WARNER CHILCOTT CO LLC
ASACOL HD, MESALAMINE
ASACOL, MESALAMINE
ENABLEX, DARIFENACIN HYDROBROMIDE

WATSON LABS

- * WATSON LABORATORIES
ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE, ACETAMINOPHEN
ALENDRONATE SODIUM, ALENDRONATE SODIUM
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANASTROZOLE, ANASTROZOLE
CARBOPLATIN, CARBOPLATIN
CARVEDILOL, CARVEDILOL
FOLIC ACID, FOLIC ACID
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
METFORMIN HYDROCHLORIDE, METFORMIN 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
NORCO, ACETAMINOPHEN
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
OXAPROZIN, OXAPROZIN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC
 - ACARBOSE, ACARBOSE
 - ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 - ACETAZOLAMIDE, ACETAZOLAMIDE
 - ACETOHEXAMIDE, ACETOHEXAMIDE
 - ACYCLOVIR, ACYCLOVIR
 - AFEDITAB CR, NIFEDIPINE
 - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - ALENDRONATE SODIUM, ALENDRONATE SODIUM
 - ALLOPURINOL, ALLOPURINOL
 - ALORA, ESTRADIOL
 - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - AMOXAPINE, AMOXAPINE
 - ANDRODERM, TESTOSTERONE
 - ATENOLOL AND CHLORTHALIDONE, ATENOLOL
 - ATENOLOL, ATENOLOL
 - BACLOFEN, BACLOFEN
 - BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
 - BREVICON 28-DAY, ETHINYL ESTRADIOL
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
 - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
 - BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
 - CABERGOLINE, CABERGOLINE
 - CAPTOPRIL, CAPTOPRIL
 - CARBOPLATIN, CARBOPLATIN
 - CARISOPRODOL, CARISOPRODOL
 - CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
 - CHLORPROPAMIDE, CHLORPROPAMIDE
 - CHLORZOXAZONE, CHLORZOXAZONE
 - CICLOPIROX, CICLOPIROX
 - CIMETIDINE, CIMETIDINE
 - CIMETIDINE, CIMETIDINE (OTC)
 - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 - CLONAZEPAM, CLONAZEPAM
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 - COL-PROBENECID, COLCHICINE
 - CORDRAN, FLURANDRENOLIDE
 - CRINONE, PROGESTERONE
 - CROMOLYN SODIUM, CROMOLYN SODIUM
 - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 - CYCLOSPORINE, CYCLOSPORINE
 - DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
 - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 - DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 - DIAZEPAM, DIAZEPAM
 - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 - DILACOR XR, DILTIAZEM HYDROCHLORIDE
 - DIMENHYDRINATE, DIMENHYDRINATE
 - DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DIPYRIDAMOLE, DIPYRIDAMOLE
 - DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 - DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 - DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 - DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 - DOXEPEPIN HYDROCHLORIDE, DOXEPEPIN HYDROCHLORIDE
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 - DOXYCYCLINE, DOXYCYCLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC
 - DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
 - ENALAPRIL MALEATE, ENALAPRIL MALEATE
 - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 - ERGOLOID MESYLATES, ERGOLOID MESYLATES
 - ESTAZOLAM, ESTAZOLAM
 - ESTRADIOL CYPIONATE, ESTRADIOL CYPIONATE
 - ESTRADIOL VALERATE, ESTRADIOL VALERATE
 - ESTRADIOL, ESTRADIOL
 - ESTRONE, ESTRONE
 - ESTROPIPATE, ESTROPIPATE
 - FAMCICLOVIR, FAMCICLOVIR
 - FAMOTIDINE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FENTANYL CITRATE, FENTANYL CITRATE
 - FENTANYL-100, FENTANYL
 - FENTANYL-25, FENTANYL
 - FENTANYL-50, FENTANYL
 - FENTANYL-75, FENTANYL
 - FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 - FLUTAMIDE, FLUTAMIDE
 - FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 - FUROSEMIDE, FUROSEMIDE
 - GABAPENTIN, GABAPENTIN
 - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
 - GELNIQUE, OXYBUTYNIN CHLORIDE
 - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
 - GEMFIBROZIL, GEMFIBROZIL
 - GLIMEPIRIDE, GLIMEPIRIDE
 - GLIPIZIDE, GLIPIZIDE
 - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
 - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
 - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 - IBUPROFEN, IBUPROFEN
 - IBUPROFEN, IBUPROFEN (OTC)
 - INDAPAMIDE, INDAPAMIDE
 - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - ISONIAZID, ISONIAZID
 - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 - ISRADIPINE, ISRADIPINE
 - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 - LAMOTRIGINE, LAMOTRIGINE
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 - LEVONORGESTREL, LEVONORGESTREL
 - LEVONORGESTREL, LEVONORGESTREL (OTC)
 - LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 - LISINOPRIL, LISINOPRIL
 - LORAZEPAM, LORAZEPAM
 - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
 - LOW-OGESTREL-21, ETHINYL ESTRADIOL
 - LOW-OGESTREL-28, ETHINYL ESTRADIOL
 - LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 - MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 - MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC
MELOXICAM, MELOXICAM
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METHOCARBAMOL, METHOCARBAMOL
METHYCLOTHIAZIDE, METHYCLOTHIAZIDE
METHYLDOPA, METHYLDOPA
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE, METRONIDAZOLE
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
MICROZIDE, HYDROCHLOROTHIAZIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL, MINOXIDIL
MIRTAZAPINE, MIRTAZAPINE
MORPHINE SULFATE, MORPHINE SULFATE
NABUMETONE, NABUMETONE
NAPOXEN SODIUM, NAPOXEN SODIUM
NAPOXEN, NAPOXEN
NATEGLINIDE, NATEGLINIDE
NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NIZATIDINE, NIZATIDINE
NORCO, ACETAMINOPHEN
NORETHIN 1/35E-21, ETHINYL ESTRADIOL
NORETHIN 1/35E-28, ETHINYL ESTRADIOL
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL (7/14), ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL, 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
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
OXAZEPAM, OXAZEPAM
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE AND ASPIRIN, ASPIRIN
OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
PENTAZOCINE AND NALOXONE HYDROCHLORIDES, NALOXONE HYDROCHLORIDE
PENTOXIFYLLINE, PENTOXIFYLLINE
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
PREDNISOLONE, PREDNISOLONE
PREDNISONE, PREDNISONE
PRIMIDONE, PRIMIDONE
PROBENECID, PROBENECID
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
QUASENSE, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC
 - QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 - QUINIDINE SULFATE, QUINIDINE SULFATE
 - RAMIPRIL, RAMIPRIL
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 - RAPAFLO, SILODOSIN
 - RISPERIDONE, RISPERIDONE
 - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - SIMVASTATIN, SIMVASTATIN
 - SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - SULFASALAZINE, SULFASALAZINE
 - SULINDAC, SULINDAC
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TACROLIMUS, TACROLIMUS
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TEMAZEPAM, TEMAZEPAM
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
 - TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
 - TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 - THIOTHIXENE, THIOTHIXENE
 - TOLAZAMIDE, TOLAZAMIDE
 - TOLBUTAMIDE, TOLBUTAMIDE
 - TOPIRAMATE, TOPIRAMATE
 - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
 - TRANDOLAPRIL, TRANDOLAPRIL
 - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
 - TRELSTAR, TRIPTORELIN PAMOATE
 - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - TRIAZOLAM, TRIAZOLAM
 - TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
 - TRIMETHOPRIM, TRIMETHOPRIM
 - TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
 - TRIVORA-28, ETHINYL ESTRADIOL
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - VECURONIUM BROMIDE, VECURONIUM BROMIDE
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 - ZOVIA 1/35E-28, ETHINYL ESTRADIOL
 - ZOVIA 1/50E-28, ETHINYL ESTRADIOL
- * WATSON LABS INC
 - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

WATSON LABS (UTAH)

- * WATSON LABORATORIES INC
 - INFED, IRON DEXTRAN
 - NOR-QD, NORETHINDRONE
 - OXYTROL, OXYBUTYNIN
 - PROGESTERONE, PROGESTERONE

WATSON LABS FLORIDA

- * WATSON LABORATORIES INC FLORIDA
 - ALPRAZOLAM, ALPRAZOLAM
 - BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
 - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
 - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
 - CARTIA XT, DILTIAZEM HYDROCHLORIDE
 - CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
 - CLARITHROMYCIN, CLARITHROMYCIN
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WATSON LABORATORIES INC FLORIDA
FOSINOPRIL SODIUM AND HYDROCHLORTIAZIDE, FOSINOPRIL SODIUM
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GLIMEPIRIDE, GLIMEPIRIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
KETOPROFEN, KETOPROFEN
LEVETIRACETAM, LEVETIRACETAM
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
MIRTAZAPINE, MIRTAZAPINE
OMEPRAZOLE, OMEPRAZOLE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RISPERIDONE, RISPERIDONE
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
TAZTIA XT, DILTIAZEM HYDROCHLORIDE

WATSON LABS INC

- * WATSON LABORATORIES INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
FIORICET W/ CODEINE, ACETAMINOPHEN
FIORINAL W/CODEINE, ASPIRIN
FIORINAL, ASPIRIN
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
URSODIOL, URSODIOL

WATSON PHARMS

- * WATSON PHARMACEUTICALS
TENUATE DOSPAN, DIETHYLPROMION HYDROCHLORIDE
TENUATE, DIETHYLPROMION HYDROCHLORIDE
- * WATSON PHARMACEUTICALS INC
ACTIGALL, URSODIOL
CONDYLOX, PODOFILOX
CORDRAN SP, FLURANDRENOLIDE
CORDRAN, FLURANDRENOLIDE
FIORICET, ACETAMINOPHEN
MONODOX, DOXYCYCLINE

WE PHARMS

- * WE PHARMACEUTICALS INC
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE

WEILL MEDCL COLL

- * WEILL MEDICAL COLLEGE CORNELL UNIV
FLUDEOXYGLUCOSE F 18, FLUDEOXYGLUCOSE F-18

WELLSPRING PHARM

- * WELLSPRING PHARMACEUTICAL CORP
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE
DUVOID, BETHANECHOL CHLORIDE
DYRENIUM, TRIAMTERENE

WEST WARD

- * WEST WARD INC
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
- * WEST WARD PHARMACEUTICAL CORP
AMINOPHYLLINE, AMINOPHYLLINE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
CAPTOPRIL, CAPTOPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WEST WARD PHARMACEUTICAL CORP
CARISOPRODOL, CARISOPRODOL
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
CHLOROTHIAZIDE, CHLOROTHIAZIDE
CORTISONE ACETATE, CORTISONE ACETATE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIGOXIN, DIGOXIN
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
GLYCOPYRROLATE, GLYCOPYRROLATE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROCORTISONE, HYDROCORTISONE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
ISONIAZID, ISONIAZID
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LITHIUM CARBONATE, LITHIUM CARBONATE
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
PREDNISONE, PREDNISONE
PRIMIDONE, PRIMIDONE
PROPYLTIOURACIL, PROPYLTIOURACIL
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
ZALEPLON, ZALEPLON

WEST WARD PHARM CORP

- * WEST WARD PHARMACEUTICAL CORP
PREDNISONE, PREDNISONE

WEST WARD PHARMS

- * WEST WARD PHARMACEUTICALS CORP
RISPERIDONE, RISPERIDONE

WESTWARD

- * WESTWARD PHARMACEUTICAL CORP
NAPROXEN, NAPROXEN
RIFAMPIN AND ISONIAZID, ISONIAZID

WEST-WARD PHARM CORP

- * WEST-WARD PHARMACEUTICAL CORP
CEFOTETAN, CEFOTETAN DISODIUM
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE

WOCKHARDT

- * WOCKHARDT AMERICAS INC
CAPTOPRIL, CAPTOPRIL
FAMOTIDINE, FAMOTIDINE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
- * WOCKHARDT EU OPERATIONS (SWISS) AG
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETIC ACID, ACETIC ACID, GLACIAL
ALBUTEROL SULFATE, ALBUTEROL SULFATE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
BROMFED-DM, BROMPHENIRAMINE MALEATE
CARBAMAZEPINE, CARBAMAZEPINE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WOCKHARDT EU OPERATIONS (SWISS) AG
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CROMOLYN SODIUM, CROMOLYN SODIUM
 - CYCLOSPORINE, CYCLOSPORINE
 - DEXAMETHASONE, DEXAMETHASONE
 - DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
 - DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 - ERYTHROMYCIN, ERYTHROMYCIN
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - FUROSEMIDE, FUROSEMIDE
 - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - LEVETIRACETAM, LEVETIRACETAM
 - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 - LINDANE, LINDANE
 - LITHIUM CITRATE, LITHIUM CITRATE
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - LORATADINE, LORATADINE (OTC)
 - MEGESTROL ACETATE, MEGESTROL ACETATE
 - METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 - NYSTATIN, NYSTATIN
 - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
 - PHENYTOIN, PHENYTOIN
 - 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
 - TRETINOIN, TRETINOIN
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 - VALPROIC ACID, VALPROIC ACID
- * WOCKHARDT LTD
 - ADENOSINE, ADENOSINE
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
 - AZITHROMYCIN, AZITHROMYCIN
 - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
 - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
 - CARVEDILOL, CARVEDILOL
 - CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
 - CEFOTAXIME, CEFOTAXIME SODIUM
 - CEFPROZIL, CEFPROZIL
 - CEFTAZIDIME, CEFTAZIDIME
 - CEFTRIAXONE, CEFTRIAXONE SODIUM
 - CEFUROXIME AXETIL, CEFUROXIME AXETIL
 - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
 - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
 - CLARITHROMYCIN, CLARITHROMYCIN
 - DIVALPROEX SODIUM, DIVALPROEX SODIUM
 - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
 - EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - FAMOTIDINE, FAMOTIDINE (OTC)
 - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
 - FUROSEMIDE, FUROSEMIDE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - LEVETIRACETAM, LEVETIRACETAM
 - LEVOFLOXACIN, LEVOFLOXACIN
 - LISINOPRIL, LISINOPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ****

- * WOCKHARDT LTD
 - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - NIACIN, NIACIN
 - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
 - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
 - RISPERIDONE, RISPERIDONE
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
 - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
 - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
 - TIMOLOL MALEATE, TIMOLOL MALEATE
 - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
 - VENLAFAKINE HYDROCHLORIDE, VENLAFAKINE HYDROCHLORIDE
 - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
 - ZONISAMIDE, ZONISAMIDE

WOCKHARDT USA

- * WOCKHARDT USA INC
 - EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
 - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 - OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
- * WOCKHARDT USA LLC
 - ENALAPRIL MALEATE, ENALAPRIL MALEATE
 - TOPIRAMATE, TOPIRAMATE

WORLD GEN

- * WORLD GEN LLC
 - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

WRASER PHARMS

- * WRASER PHARMACEUTICALS LLC
 - CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

WRASER PHARMS LLC

- * WRASER PHARMACEUTICALS LLC
 - ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN

WYETH CONS

- * WYETH CONSUMER HEALTHCARE
 - 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 PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
 - ADVIL, IBUPROFEN (OTC)
 - ALAVERT, LORATADINE (OTC)
 - AXID AR, NIZATIDINE (OTC)
 - 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)
 - JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 - LORATADINE, LORATADINE (OTC)
 - PEDIATRIC ADVIL, IBUPROFEN (OTC)

WYETH PHARMS

- * WYETH PHARMACEUTICALS INC SUB PFIZER INC
 - LO/OVRAL-28, ETHINYL ESTRADIOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** W ******WYETH PHARMS INC**

- * WYETH PHARMACEUTICALS INC
 - CORDARONE, AMIODARONE HYDROCHLORIDE
 - EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
 - LYBREL, ETHINYL ESTRADIOL
 - PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
 - PREMARIN, ESTROGENS, CONJUGATED
 - PREMPHASE 14/14, ESTROGENS, CONJUGATED
 - PREMPRO, ESTROGENS, CONJUGATED
 - PRISTIQ, DESVENLAFAXINE SUCCINATE
 - PROTONIX IV, PANTOPRAZOLE SODIUM
 - PROTONIX, PANTOPRAZOLE SODIUM
 - RAPAMUNE, SIROLIMUS
 - TORISEL, TEMSIROLIMUS
 - TRECATOR, ETHIONAMIDE
 - TYGACIL, TIGECYCLINE
 - ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
 - ZOSYN, PIPERACILLIN SODIUM

X GEN PHARMS

- * X GEN PHARMACEUTICALS INC
 - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 - AMPHOTERICIN B, AMPHOTERICIN B
 - BACIIM, BACITRACIN
 - BACI-RX, BACITRACIN
 - COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
 - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
 - HYDROCORTISONE ACETATE, HYDROCORTISONE ACETATE
 - HYDRO-RX, HYDROCORTISONE
 - LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
 - NEO-FRADIN, NEOMYCIN SULFATE
 - NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
 - NEOMYCIN SULFATE, NEOMYCIN SULFATE
 - NEO-RX, NEOMYCIN SULFATE
 - NYSTATIN, NYSTATIN
 - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 - POLY-RX, POLYMYXIN B SULFATE
 - STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

XANODYNE PHARM

- * XANODYNE PHARMACEUTICS INC
 - AMICAR, AMINOCAPROIC ACID
 - ORAMORPH SR, MORPHINE SULFATE
 - ZIPSOR, DICLOFENAC POTASSIUM

XANODYNE PHARMS

- * XANODYNE PHARMACEUTICALS INC
 - ROXICODONE, OXYCODONE HYDROCHLORIDE

X-GEN PHARMS

- * X-GEN PHARMACEUTICALS INC
 - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
 - PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

XTTRIUM

- * XTTRIUM LABORATORIES INC
 - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 - DYNA-HEX, CHLORHEXIDINE GLUCONATE (OTC)
 - EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

YUNG SHIN PHARM

- * YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
 - CEFACLOR, CEFACLOR
 - CEPHALEXIN, CEPHALEXIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ******ZAMBON SPA**

* ZAMBON SPA ITALY
MONUROL, FOSFOMYCIN TROMETHAMINE

ZARS PHARM

* ZARS PHARMA INC
SYNERA, LIDOCAINE

ZOGENIX INC

* ZOGENIX INC
SUMAVEL DOSEPRO, SUMATRIPTAN SUCCINATE

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
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
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
ACETAZOLAMIDE, ACETAZOLAMIDE
ALPRAZOLAM, ALPRAZOLAM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
ANASTROZOLE, ANASTROZOLE
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
BICALUTAMIDE, BICALUTAMIDE
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
CARVEDILOL, CARVEDILOL
DIPYRIDAMOLE, DIPYRIDAMOLE
DIVALPROEX SODIUM, DIVALPROEX SODIUM
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
FINASTERIDE, FINASTERIDE
GABAPENTIN, GABAPENTIN
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
INDOMETHACIN, INDOMETHACIN
LAMOTRIGINE, LAMOTRIGINE
LEVETIRACETAM, LEVETIRACETAM
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
RAMIPRIL, RAMIPRIL
RISPERIDONE, RISPERIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** Z ****

- * ZYDUS PHARMACEUTICALS USA INC
 - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
 - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
 - TOPIRAMATE, TOPIRAMATE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

APPENDIX C**UNIFORM TERMS****DOSAGE FORMS**

AEROSOL	LOTION/SHAMPOO
AEROSOL, FOAM	OIL
AEROSOL, METERED	OIL/DROPS
CAPSULE	ointment
CAPSULE, DELAYED REL PELLETS	ointment, augmented
CAPSULE, DELAYED RELEASE	PASTE
CAPSULE, EXTENDED RELEASE	PATCH
CLOTH	PELLET
CONCENTRATE	POWDER
CREAM	POWDER, EXTENDED RELEASE
CREAM, AUGMENTED	POWDER, METERED
ELIXIR	RING
EMULSION	SHAMPOO
ENEMA	SOLUTION
FILM	SOLUTION FOR SLUSH
FILM, EXTENDED RELEASE	SOLUTION, GEL FORMING/DROPS
FOR SOLUTION	SOLUTION, METERED
FOR SOLUTION, TABLET, FOR SOLUTION	SOLUTION/DROPS
FOR SUSPENSION	SPONGE
FOR SUSPENSION, DELAYED RELEASE	SPRAY
FOR SUSPENSION, EXTENDED RELEASE	SPRAY, METERED
GAS	SUPPOSITORY
GEL	SUSPENSION
GEL, AUGMENTED	SUSPENSION, EXTENDED RELEASE
GEL, METERED	SUSPENSION/DROPS
GRANULE	SWAB
GRANULE, DELAYED RELEASE	SYRUP
GUM, CHEWING	SYSTEM, EXTENDED RELEASE
IMPLANT	TABLET
INHALANT	TABLET, CHEWABLE
INJECTABLE	TABLET, COATED PARTICLES
INJECTABLE, LIPID COMPLEX	TABLET, DELAYED RELEASE
INJECTABLE, LIPOSOMAL	TABLET, DELAYED RELEASE, ORALLY DISINTEGRATING
INSERT	TABLET, EFFERVESCENT
INSERT, EXTENDED RELEASE	TABLET, EXTENDED RELEASE
INTRAUTERINE DEVICE	TABLET, FOR SUSPENSION
JELLY	TABLET, ORALLY DISINTEGRATING
LIQUID	TAPE
LOTION	TROCHE/LOZENGE
LOTION, AUGMENTED	

APPENDIX C**UNIFORM TERMS*****ROUTES OF ADMINISTRATION***

BUCCAL	INTRAVITREAL
DENTAL	IRRIGATION
ENDOCERVICAL	IV (INFUSION)
EPIDURAL	IV (INFUSION)-SC
FOR RX COMPOUNDING	IV-SC
IM-IV	N/A
IM-IV-SC	NASAL
IMPLANTATION	OPHTHALMIC
IM-SC	ORAL
INHALATION	ORAL-21
INJECTION	ORAL-28
INTRA-ANAL	OTIC
INTRA-ARTICULAR, INTRAMUSCULAR, INTRAVITREAL	PERFUSION, CARDIAC PERIODONTAL
INTRACRANIAL	RECTAL
INTRAMUSCULAR	SPINAL
INTRAOCULAR	SUBCUTANEOUS
INTRAPERITONEAL	SUBLINGUAL
INTRAPLEURAL	TOPICAL
INTRATHECAL	TRANSDERMAL
INTRATRACHEAL	TRANSMUCOSAL
INTRAUTERINE	URETHRAL
INTRAVENOUS	VAGINAL
INTRAVESICAL	

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
INH	INHALATION
IU	INTERNATIONAL UNITS
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 Drug Price Competition and Patent Term Restoration Act (1984 Amendments) for periods of exclusivity, during which abbreviated new drug applications (ANDAs) and applications described in Section 505(b)(2) of the Federal Food, Drug, and Cosmetic Act (the Act) for those drug products may, in some instances, not be submitted or made effective as described below, and provides patent information concerning the listed drug products. Those drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the Act and those drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A are also included in this *Addendum*. This section is arranged in alphabetical order by active ingredient name followed by the trade name. Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For an explanation of the codes used in the *Addendum*, see the *Patent and Exclusivity Terms* Section. Exclusivity prevents the submission or effective approval of ANDAs or applications described in Section 505(b)(2) of the Act. It does not prevent the submission or approval of a second 505(b)(1) application except in the case of Orphan Drug exclusivity. Applications qualifying for periods of exclusivity are:

- (1) A new drug application approved after September 24, 1984, for a drug product all active ingredients (including any ester or salt of the active ingredient) of which had never been approved in any other new drug application under Section 505 (b) of the Act. No subsequent ANDA or application described in Section 505(b)(2) of the Act for the same drug 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.
- (2) A new drug application approved after September 24, 1984, for a drug product containing an active ingredient (including any ester or salt of that 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, the approval of a subsequent ANDA or an application described in Section 505(b)(2) of the Act may not be made effective for the same drug or use, if for a new indication, before the expiration of three years from the date of approval of the original application. If an applicant has exclusivity for a new application or 505(b)(2) application for the drug product with indications or use, this does not preclude the approval of an ANDA or 505(b)(2) application not covered by the exclusivity.
- (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. The approval of a subsequent ANDA or 505(b)(2) application for a change approved in the supplement may not be

made effective for three years from the date of approval of the original supplement.

The 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 FDA Form 3524a "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 approval on FDA Form 3542 "Patent Information Submitted Upon and After Approval of an NDA or Supplement". Patent information on unapproved applications or on patents beyond the scope of the Act (i.e., process or manufacturing patents) will not be published. FDA form 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; use patents for a particular approved indication or method of using the product; and certain other patents as detailed on FDA Form 3542. This information, as provided by the sponsor on FDA form 3542, will be published as described above.

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 Q1 Act. A guidance for industry on this subject is available at <http://www.fda.gov/downloads/AboutFDA/ReportsManualsForms/Forms/UCM048345.pdf>

Upon approval, patent numbers and expiration dates, in addition to certain other information on appropriate patents claiming drug products that are the subject of approved applications, will be published daily in the Electronic Orange Book Query. 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 Electronic Orange Book, updated daily, should be consulted for the most recent patent and exclusivity information.

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ABACAVIR SULFATE - ZIAGEN</u>								
N020977 001	5034394	Dec	18, 2011	DS	DP			
	5034394*PED	Jun	18, 2012					
	6294540	May	14, 2018	DS	DP	U-65		
	6294540*PED	Nov	14, 2018			U-65		
<u>ABACAVIR SULFATE - ZIAGEN</u>								
N020978 001	5034394	Dec	18, 2011	DS	DP			
	5034394*PED	Jun	18, 2012					
	6294540	May	14, 2018	DS	DP	U-65		
	6294540*PED	Nov	14, 2018			U-65		
	6641843	Feb	04, 2019		DP			
	6641843*PED	Aug	04, 2019					
<u>ABACAVIR SULFATE; LAMIVUDINE - EPZICOM</u>								
N021652 001	5034394	Dec	18, 2011	DS	DP			
	5034394*PED	Jun	18, 2012					
	5905082	May	18, 2016	DS	DP			
	5905082*PED	Nov	18, 2016					
	6294540	May	14, 2018	DS	DP	U-257		
	6294540*PED	Nov	14, 2018					
	6417191	Mar	28, 2016		DP	U-257		
<u>ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE - TRIZIVIR</u>								
N021205 001	5034394	Dec	18, 2011	DS	DP			
	5034394*PED	Jun	18, 2012					
	5905082	May	18, 2016	DS	DP	U-248		
	5905082*PED	Nov	18, 2016					
	6294540	May	14, 2018	DS	DP	U-65		
	6294540*PED	Nov	14, 2018			U-65		
	6417191	Mar	28, 2016		DP	U-248		
<u>ABARELIX - PLENAXIS</u>								
N021320 001	5843901	Dec	01, 2015	DS	DP			
	5968895	Dec	11, 2016		DP			
	6180608	Dec	11, 2016		DP	U-549		
	6423686	Jun	07, 2015	DS				
	6455499	Jun	07, 2015			U-549		
	6699833	Dec	11, 2016		DP			
<u>ABIRATERONE ACETATE - ZYTIGA</u>								
N020379 001	5604213	Feb	18, 2014	DS	DP	U-1126	NCE	Apr 28, 2016
<u>ACETAMINOPHEN - OFIRMEV</u>								
N022450 001	6028222	Aug	05, 2017		DP		NP	Nov 02, 2013
	6992218	Jun	06, 2021		DP			
<u>ACETAMINOPHEN; ASPIRIN; CAFFEINE - EXCEDRIN (MIGRAINE)</u>								
N020802 001	5972916	Jul	14, 2017			U-296		
<u>ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE - ULTRACET</u>								
N021123 001	RE39221	Aug	09, 2011	DS	DP	U-55		
<u>ACETYLCOLINE CHLORIDE - MIOCHOL-E</u>								
N020213 001	6261546	Apr	29, 2019			U-506		
<u>ACYCLOVIR; HYDROCORTISONE - XERESE</u>								
N022436 001	6514980	Jan	24, 2017		DP	U-1006	NC	Jul 31, 2012
	7223387	Feb	28, 2021		DP	U-1006		
	RE39264	Feb	02, 2016		DP	U-1006		
<u>ADAPALENE - DIFFERIN</u>								
N021753 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		
<u>ADAPALENE - DIFFERIN</u>								
N022502 001	7998467	May	31, 2028	DP	U-1078		NDF	Mar 17, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ADAPALENE; BENZOYL PEROXIDE - EPIDUO</u>						
N022320 001	7820186	Nov 23, 2025	DP		NC	Dec 08, 2011
	7964202	Sep 01, 2024	DP	U-1078		
	8071644	Jul 18, 2027	DP	U-1078		
	8080537	Jul 18, 2027		U-1078		
<u>ADEFEOVIR DIPIVOXIL - HEP SERA</u>						
N021449 001	5663159	Sep 02, 2014	DS	DP		
	6451340	Jul 23, 2018	DS	DP	U-470	
<u>ADENOSINE - ADENOSCAN</u>						
N020059 001	5731296	Mar 24, 2015			U-221	
<u>ALATROFLOXACIN MESYLATE - TROVAN PRESERVATIVE FREE</u>						
N020760 001	5763454	Jun 15, 2015			U-282	
	6080756	Jul 05, 2016				
	6194429	Jul 23, 2018				
<u>ALATROFLOXACIN MESYLATE - TROVAN PRESERVATIVE FREE</u>						
N020760 002	5763454	Jun 15, 2015			U-282	
	6080756	Jul 05, 2016				
	6194429	Jul 23, 2018				
<u>ALBUMIN HUMAN - OPTISON</u>						
N020899 001	5529766	Jun 25, 2013			U-505	
	5558094	Feb 28, 2012			U-505	
	5573751	Apr 25, 2012				
	6723303	Apr 20, 2021	DP			
<u>ALBUTEROL SULFATE - ACCUNEB</u>						
N020949 001	6702997	Dec 28, 2021			U-558	
<u>ALBUTEROL SULFATE - ACCUNEB</u>						
N020949 002	6702997	Dec 28, 2021			U-558	
<u>ALBUTEROL SULFATE - PROAIR HFA</u>						
N021457 001	5605674	Feb 25, 2014	DP		NPP	Sep 16, 2011
	7105152	Sep 12, 2023	DP			
	7566445	Jun 04, 2017	DP			
<u>ALBUTEROL SULFATE - PROVENTIL-HFA</u>						
N020503 001	5605674	Feb 25, 2014				
	5766573	Jun 16, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ALBUTEROL SULFATE - VENTOLIN HFA</u>						
N020983 001	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018				
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6510969	Dec 23, 2017	DP			
	6510969*PED	Jun 23, 2018				
	6558651	Dec 19, 2016	DP U-716			
	6558651*PED	Jun 19, 2017				
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	6743413	Jun 01, 2021		U-716		
	6743413*PED	Dec 01, 2021				
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018				
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018				
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018				
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 06, 2018				
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018				
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019				
	7500444	Jan 04, 2025	DP			
	7500444*PED	Jul 04, 2025				
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT</u>						
N020291 001	5603918	Jun 09, 2015				
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT RESPIMAT</u>						
N021747 001	5405084	Apr 11, 2012	DP		NP	Oct 07, 2014
	5472143	Sep 29, 2013	DP			
	5497944	Mar 12, 2013	DP			
	5662271	Sep 02, 2014	DP			
	5911851	Sep 29, 2013	DP			
	5964416	Oct 04, 2016	DP			
	6007676	Sep 29, 2013	DP			
	6149054	Dec 19, 2016	DP			
	6176442	Oct 04, 2016	DP			
	6453795	Dec 05, 2016	DP			
	6503362	Sep 29, 2013	DP			
	6726124	Oct 04, 2016	DP			
	6846413	Aug 28, 2018	DP			
	6977042	Aug 28, 2018	DP			
	6988496	Feb 23, 2020	DP			
	7104470	Oct 04, 2016	DP			
	7246615	May 31, 2016	DP			
	7284474	Aug 26, 2024	DP			
	7396341	Oct 10, 2026	DP			
	7802568	Feb 26, 2019	DP			
	7837235	Mar 13, 2028	DP			
	7896264	May 26, 2025	DP			
	7988001	Aug 04, 2021	DP			
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - DUONEB</u>						
N020950 001	6632842	Dec 28, 2021		U-532		
<u>ALCAFTADINE - LASTACAF</u>						
N022134 001	5468743	Nov 21, 2012	DS DP		NCE	Jul 28, 2015

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ALCOHOL; CHLORHEXIDINE GLUCONATE - AVAGARD</u>						
N021074 001	5897031	Jun 21, 2016				
	6090395	Jun 22, 2015	DP			
	6534069	Jun 22, 2015	DP			
	6623744	Jun 23, 2015		U-1008		
	7081246	Aug 03, 2016	DP			
	7566460	Jun 22, 2015	DP	U-1008		
<u>ALENDRONATE SODIUM - FOSAMAX</u>						
N020560 001	5358941	Dec 02, 2012				
	5358941*PED	Jun 02, 2013				
	5681590	Dec 02, 2012				
	5681590*PED	Jun 02, 2013				
	6090410	Dec 02, 2012				
	6090410*PED	Jun 02, 2013				
	6194004	Dec 02, 2012				
	6194004*PED	Jun 02, 2013				
<u>ALENDRONATE SODIUM - FOSAMAX</u>						
N020560 002	5358941	Dec 02, 2012				
	5358941*PED	Jun 02, 2013				
	5681590	Dec 02, 2012				
	5681590*PED	Jun 02, 2013				
	6090410	Dec 02, 2012				
	6090410*PED	Jun 02, 2013				
<u>ALENDRONATE SODIUM - FOSAMAX</u>						
N020560 003	5358941	Dec 02, 2012				
	5358941*PED	Jun 02, 2013				
	5681590	Dec 02, 2012				
	5681590*PED	Jun 02, 2013				
	6090410	Dec 02, 2012				
	6090410*PED	Jun 02, 2013				
<u>ALENDRONATE SODIUM - FOSAMAX</u>						
N020560 004	5358941	Dec 02, 2012				
	5358941*PED	Jun 02, 2013				
	5681590	Dec 02, 2012				
	5681590*PED	Jun 02, 2013				
	6090410	Dec 02, 2012				
	6090410*PED	Jun 02, 2013				
<u>ALENDRONATE SODIUM - FOSAMAX</u>						
N020560 005	5358941	Dec 02, 2012				
	5358941*PED	Jun 02, 2013				
	5681590	Dec 02, 2012				
	5681590*PED	Jun 02, 2013				
	6090410	Dec 02, 2012				
	6090410*PED	Jun 02, 2013				
<u>ALENDRONATE SODIUM - FOSAMAX</u>						
N021575 001	5462932	May 17, 2014				
	5462932*PED	Nov 17, 2014				
	5994329	Jul 17, 2018			Y	
	5994329*PED	Jan 17, 2019				
	6015801	Jul 17, 2018			Y	
	6015801*PED	Jan 17, 2019				
	6225294	Jul 17, 2018			Y	
	6225294*PED	Jan 17, 2019				
<u>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u>						
N021762 001	5358941	Dec 02, 2012	DP			
	5358941*PED	Jun 02, 2013				
	5681590	Dec 02, 2012	DP			
	5681590*PED	Jun 02, 2013				
	5994329	Jul 17, 2018		U-647	Y	
	5994329*PED	Jan 17, 2019				
	6090410	Dec 02, 2012	DP			
	6090410*PED	Jun 02, 2013	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>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u>							
N021762 002	5358941	Dec	02, 2012	DP			
	5358941*PED	Jun	02, 2013				
	5681590	Dec	02, 2012	DP			
	5681590*PED	Jun	02, 2013				
	6090410	Dec	02, 2012	DP			
	6090410*PED	Jun	02, 2013	DP			
<u>ALFUZOSIN HYDROCHLORIDE - ALFUZOSIN HYDROCHLORIDE</u>							
A079013 001					PC		Jan 14, 2012
<u>ALFUZOSIN HYDROCHLORIDE - ALFUZOSIN HYDROCHLORIDE</u>							
A079014 001					PC		Jan 14, 2012
<u>ALFUZOSIN HYDROCHLORIDE - ALFUZOSIN HYDROCHLORIDE</u>							
A079054 001					PC		Jan 14, 2012
<u>ALFUZOSIN HYDROCHLORIDE - ALFUZOSIN HYDROCHLORIDE</u>							
A079056 001					PC		Jan 14, 2012
<u>ALFUZOSIN HYDROCHLORIDE - ALFUZOSIN HYDROCHLORIDE</u>							
A079057 001					PC		Jan 14, 2012
<u>ALFUZOSIN HYDROCHLORIDE - UROXATRAL</u>							
N021287 001	4661491*PED	Jul	18, 2011		M-97		Dec 15, 2013
	6149940	Aug	22, 2017		PED		Jun 15, 2014
	6149940*PED	Feb	22, 2018				
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>							
N021985 001	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>							
N021985 002	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>							
N022545 001					NC		Aug 26, 2013
					NCE		Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>							
N022545 002					NC		Aug 26, 2013
					NCE		Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>							
N022545 003					NC		Aug 26, 2013
					NCE		Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u>							
N022545 004					NC		Aug 26, 2013
					NCE		Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>							
N200045 001	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
					NC		Dec 21, 2013
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>							
N200045 002	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
					NC		Dec 21, 2013
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>							
N200045 003	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
					NC		Dec 21, 2013
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>							
N200045 004	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
					NC		Dec 21, 2013
<u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u>							
N200045 005	5559111	Jul	21, 2018	DS DP U-3	NCE		Mar 05, 2012
					NC		Dec 21, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N022107 001	5559111	Jul 21, 2018	DS DP U-3		I-600 NCE	Jul 16, 2012 Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N022107 002	5559111	Jul 21, 2018	DS DP U-3		I-600 NCE	Jul 16, 2012 Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N022107 003	5559111	Jul 21, 2018	DS DP U-3		I-600 NCE	Jul 16, 2012 Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKTURNA HCT</u>						
N022107 004	5559111	Jul 21, 2018	DS DP U-3		I-600 NCE	Jul 16, 2012 Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N022217 001	5399578 5559111	Mar 21, 2012 Jul 21, 2018	DS DP U-3		NCE NC	Mar 05, 2012 Sep 16, 2012
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N022217 002	5399578 5559111	Mar 21, 2012 Jul 21, 2018	DS DP U-3		NCE NC	Mar 05, 2012 Sep 16, 2012
<u>ALITRETNINOIN - PANRETIN</u>						
N020886 001	5932622	Aug 03, 2016		U-562		
<u>ALMOTRIPTAN MALATE - AXERT</u>						
N021001 001	5565447 5565447*PED	May 07, 2015 Nov 07, 2015	DS DP U-969			
<u>ALMOTRIPTAN MALATE - AXERT</u>						
N021001 002	5565447 5565447*PED	May 07, 2015 Nov 07, 2015	DS DP U-969			
<u>ALOSETRON HYDROCHLORIDE - LOTRONEX</u>						
N021107 001	5360800 6284770	Jan 13, 2013 Oct 05, 2018	DS DP U-405 U-405			
<u>ALOSETRON HYDROCHLORIDE - LOTRONEX</u>						
N021107 002	5360800 6284770	Jan 13, 2013 Oct 05, 2018	DS DP U-405 U-405			
<u>ALPRAZOLAM - NIRAVAM</u>						
N021726 001	6024981 6221392	Apr 09, 2018 Apr 09, 2018	DP			
<u>ALPRAZOLAM - NIRAVAM</u>						
N021726 002	6024981 6221392	Apr 09, 2018 Apr 09, 2018	DP			
<u>ALPRAZOLAM - NIRAVAM</u>						
N021726 003	6024981 6221392	Apr 09, 2018 Apr 09, 2018	DP			
<u>ALPRAZOLAM - NIRAVAM</u>						
N021726 004	6024981 6221392	Apr 09, 2018 Apr 09, 2018	DP			
<u>ALPROSTADIL - CAVERJECT</u>						
N020379 001	5741523	Apr 21, 2015				
<u>ALPROSTADIL - CAVERJECT</u>						
N020379 002	5741523	Apr 21, 2015				
<u>ALPROSTADIL - CAVERJECT</u>						
N020379 003	5741523	Apr 21, 2015				
<u>ALPROSTADIL - CAVERJECT</u>						
N020379 004	5741523	Apr 21, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ALPROSTADIL - CAVERJECT IMPULSE</u>							
N021212 001	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
<u>ALPROSTADIL - CAVERJECT IMPULSE</u>							
N021212 002	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
<u>ALPROSTADIL - MUSE</u>							
N020700 001	5474535	Dec	12, 2012	U-155			
	5886039	Mar	23, 2016	DP	U-155		
<u>ALPROSTADIL - MUSE</u>							
N020700 002	5474535	Dec	12, 2012	U-155			
	5886039	Mar	23, 2016	DP	U-155		
<u>ALPROSTADIL - MUSE</u>							
N020700 003	5474535	Dec	12, 2012	U-155			
	5886039	Mar	23, 2016	DP	U-155		
<u>ALPROSTADIL - MUSE</u>							
N020700 004	5474535	Dec	12, 2012	U-155			
	5886039	Mar	23, 2016	DP	U-155		
<u>ALVIMOPAN - ENTEREG</u>							
N021775 001	5250542	Mar	29, 2016	DS	DP	U-878	
	5434171	Dec	08, 2013	DS	DP	U-878	NCE
	6469030	Nov	29, 2020			U-879	May 20, 2013
<u>AMBRISENTAN - LETAIRIS</u>							
N022081 001	5703017	Dec	30, 2014	DS			
	5840722	Nov	24, 2015			U-821	NCE
	7109205	Oct	07, 2015	DS	DP		ODE
	7601730	Oct	07, 2015			U-1080	Jun 15, 2014
	RE42462	Oct	07, 2015	DS			
<u>AMBRISENTAN - LETAIRIS</u>							
N022081 002	5703017	Dec	30, 2014	DS			
	5840722	Nov	24, 2015			U-821	NCE
	7109205	Oct	07, 2015	DS	DP		ODE
	7601730	Oct	07, 2015			U-1080	Jun 15, 2014
	RE42462	Oct	07, 2015	DS			
<u>AMIFOSTINE - ETHYOL</u>							
N020221 001	5424471	Jul	31, 2012				
	5591731	Jul	31, 2012				
	5994409	Dec	08, 2017			U-305	
<u>AMIFOSTINE - ETHYOL</u>							
N020221 002	5424471	Jul	31, 2012				
	5591731	Jul	31, 2012				
	5994409	Dec	08, 2017			U-305	
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - LEVULAN</u>							
N020965 001	5079262	Sep	30, 2013	U-289			
	5954703	Oct	31, 2017	U-289			M-82
	6709446	May	01, 2018	U-289			Mar 12, 2013
	7723910	Jun	17, 2019	U-289			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>							
N022325 001	6869939	May	04, 2022	DP			
	7635773	Mar	13, 2029	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>							
N022325 002	6869939	May	04, 2022	DP			
	7635773	Mar	13, 2029	DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>							
N022325 003	6869939	May	04, 2022	DP			
	7635773	Mar	13, 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
<u>AMLEXANOX - APHTHASOL</u>						
N020511 001	5362737	Nov 08, 2011		U-243		
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N022026 001	6828339	Nov 20, 2022	DS			
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N022026 002	6828339	Nov 20, 2022	DS			
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u>						
N022026 003	6828339	Nov 20, 2022	DS			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 001	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 002	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 003	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 004	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 005	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 006	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 007	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018			U-552	
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 008	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018			U-552	
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 009	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018			U-552	
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
N021540 010	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018			U-552	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE</u>						
A078381 005					PC	Jul 02, 2011
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE</u>						
A078381 006					PC	Jul 02, 2011
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>						
N020364 002	6162802	Dec 19, 2017			U-367	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>						
N020364 003	6162802	Dec 19, 2017			U-367	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>						
N020364 004	6162802	Dec 19, 2017			U-367	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>						
N020364 005	6162802	Dec 19, 2017			U-367	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>						
N020364 006	6162802	Dec 19, 2017	DS DP		U-185	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>						
N020364 007	6162802	Dec 19, 2017	DS DP		U-185	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - TRIBENZOR</u>						
N200175 001	5616599	Apr 25, 2016	DS DP U-3		NC	Jul 23, 2013
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - TRIBENZOR</u>						
N200175 002	5616599	Apr 25, 2016	DS DP U-3		NC	Jul 23, 2013
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - TRIBENZOR</u>						
N200175 003	5616599	Apr 25, 2016	DS DP U-3		NC	Jul 23, 2013
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - TRIBENZOR</u>						
N200175 004	5616599	Apr 25, 2016	DS DP U-3		NC	Jul 23, 2013
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - TRIBENZOR</u>						
N200175 005	5616599	Apr 25, 2016	DS DP U-3		NC	Jul 23, 2013
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 001	5399578	Mar 21, 2012	DS DP U-3		NC	Apr 30, 2012
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 002	5399578	Mar 21, 2012	DS DP U-3		NC	Apr 30, 2012
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 003	5399578	Mar 21, 2012	DS DP U-3		NC	Apr 30, 2012
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 004	5399578	Mar 21, 2012	DS DP U-3		NC	Apr 30, 2012
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
<u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
N022314 005	5399578	Mar 21, 2012	DS DP U-3		NC	Apr 30, 2012
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
<u>AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL - AZOR</u>						
N022100 001	5616599	Apr 25, 2016	DS DP U-3		NC	Oct 16, 2012
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL - AZOR</u>						
N022100 002	5616599	Apr 25, 2016	DS DP U-3		NC	Oct 16, 2012
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL - AZOR</u>						
N022100 003	5616599	Apr 25, 2016	DS DP U-3		NC	Oct 16, 2012
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL - AZOR</u>						
N022100 004	5616599	Apr 25, 2016	DS DP U-3		NC	Oct 16, 2012
	5616599*PED	Oct 25, 2016				
<u>AMLODIPINE BESYLATE; TELMISARTAN - TWYNSTA</u>						
N022401 001	5591762	Jan 07, 2014	DS DP U-3		NC	Oct 16, 2012
<u>AMLODIPINE BESYLATE; TELMISARTAN - TWYNSTA</u>						
N022401 002	5591762	Jan 07, 2014	DS DP U-3		NC	Oct 16, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>AMLODIPINE BESYLATE; TELMISARTAN - TWYNSTA</u>						
N022401 003	5591762	Jan 07, 2014	DS DP U-3		NC	Oct 16, 2012
<u>AMLODIPINE BESYLATE; TELMISARTAN - TWYNSTA</u>						
N022401 004	5591762	Jan 07, 2014	DS DP U-3		NC	Oct 16, 2012
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>						
N021990 002	5399578	Mar 21, 2012	DS DP U-3			
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
	6395728	Jul 08, 2019	DP			
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>						
N021990 003	5399578	Mar 21, 2012	DS DP U-3			
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
	6395728	Jul 08, 2019	DP			
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>						
N021990 004	5399578	Mar 21, 2012	DS DP U-3			
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
	6395728	Jul 08, 2019	DP			
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>						
N021990 005	5399578	Mar 21, 2012	DS DP U-3			
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	DP U-3			
	6294197*PED	Dec 18, 2017				
	6395728	Jul 08, 2019	DP			
<u>AMMONIA, N-13 - AMMONIA N 13</u>						
N022119 001					NCE	Aug 23, 2012
					W	Aug 23, 2012
<u>AMOXICILLIN - MOXATAG</u>						
N050813 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			
<u>AMOXICILLIN; CLAVULANATE POTASSIUM - AUGMENTIN XR</u>						
N050785 001	6746692	Apr 04, 2020	DP			
	6783773	Apr 04, 2020	DP			
	6878386	Apr 04, 2020	U-926			
	7217430	Apr 04, 2020	DP U-926			
	7250176	Apr 04, 2020	U-926			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 10</u>						
N011522 007	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 2021				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 12.5</u>						
N011522 012	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 2021				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 15</u>						
N011522 013	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 2021				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 20</u>						
N011522 008	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 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
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 30</u>						
N011522 010	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 2021				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 5</u>						
N011522 009	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 2021				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 7.5</u>						
N011522 011	6384020	Jul 06, 2020				
	6384020*PED	Jan 06, 2021				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 10</u>						
N021303 001	6322819	Oct 21, 2018				
	6322819*PED	Apr 21, 2019				
	6605300	Oct 21, 2018				
	6605300*PED	Apr 21, 2019				
	RE41148	Oct 21, 2018		DP		
	RE41148*PED	Apr 21, 2019				
	RE42096	Oct 21, 2018		DP		
	RE42096*PED	Apr 21, 2019				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 15</u>						
N021303 006	6322819	Oct 21, 2018				
	6322819*PED	Apr 21, 2019				
	6605300	Oct 21, 2018				
	6605300*PED	Apr 21, 2019				
	RE41148	Oct 21, 2018		DP		
	RE41148*PED	Apr 21, 2019				
	RE42096	Oct 21, 2018		DP		
	RE42096*PED	Apr 21, 2019				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 20</u>						
N021303 002	6322819	Oct 21, 2018				
	6322819*PED	Apr 21, 2019				
	6605300	Oct 21, 2018				
	6605300*PED	Apr 21, 2019				
	RE41148	Oct 21, 2018		DP		
	RE41148*PED	Apr 21, 2019				
	RE42096	Oct 21, 2018		DP		
	RE42096*PED	Apr 21, 2019				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 25</u>						
N021303 004	6322819	Oct 21, 2018				
	6322819*PED	Apr 21, 2019				
	6605300	Oct 21, 2018				
	6605300*PED	Apr 21, 2019				
	RE41148	Oct 21, 2018		DP		
	RE41148*PED	Apr 21, 2019				
	RE42096	Oct 21, 2018		DP		
	RE42096*PED	Apr 21, 2019				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 30</u>						
N021303 003	6322819	Oct 21, 2018				
	6322819*PED	Apr 21, 2019				
	6605300	Oct 21, 2018				
	6605300*PED	Apr 21, 2019				
	RE41148	Oct 21, 2018		DP		
	RE41148*PED	Apr 21, 2019				
	RE42096	Oct 21, 2018		DP		
	RE42096*PED	Apr 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
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 5</u>							
N021303 005	6322819	Oct	21, 2018				
	6322819*PED	Apr	21, 2019				
	6605300	Oct	21, 2018				
	6605300*PED	Apr	21, 2019				
	RE41148	Oct	21, 2018		DP		
	RE41148*PED	Apr	21, 2019				
	RE42096	Oct	21, 2018		DP		
	RE42096*PED	Apr	21, 2019				
<u>AMPHOTERICIN B - ABELCET</u>							
N050724 001	5616334	Apr	01, 2014	DS			
	6406713	Jun	18, 2019	DS			
<u>AMPHOTERICIN B - AMBISOME</u>							
N050740 001	5874104	Feb	23, 2016		DP	U-922	
	5965156	Oct	12, 2016		DP	U-922	
<u>AMPRENAVIR - AGENERASE</u>							
N021007 001	5585397	Dec	17, 2013				
	5646180	Jul	08, 2014		U-257		
	5723490	Mar	03, 2015		U-257		
	6730679	Nov	11, 2017		DP		
<u>AMPRENAVIR - AGENERASE</u>							
N021007 002	5585397	Dec	17, 2013				
	5646180	Jul	08, 2014		U-257		
	5723490	Mar	03, 2015		U-257		
	6730679	Nov	11, 2017		DP		
<u>AMPRENAVIR - AGENERASE</u>							
N021039 001	5585397	Dec	17, 2013				
	5646180	Jul	08, 2014		U-257		
	5723490	Mar	03, 2015		U-257		
<u>ANASTROZOLE - ARIMIDEX</u>							
N020541 001						M-61	Dec 05, 2011
						PED	Jun 05, 2012
<u>ANIDULAFUNGIN - ERAXIS</u>							
N021632 001	5965525	Feb	17, 2020	DS	DP	U-540	
	6384013	Mar	19, 2012	DS			
	6743777	Mar	19, 2012		DP	U-540	
	6960564	Apr	12, 2021		DP	U-540	
	7709444	Apr	12, 2021		DP	U-540	
<u>ANIDULAFUNGIN - ERAXIS</u>							
N021632 002	5965525	Feb	17, 2020	DS	DP	U-540	
	6384013	Mar	19, 2012	DS			
	6743777	Mar	19, 2012		DP	U-540	
	6960564	Apr	12, 2021		DP	U-540	
	7709444	Apr	12, 2021		DP	U-540	
<u>APREPITANT - EMEND</u>							
N021549 001	5538982	Jul	23, 2013		U-745		
	5719147	Apr	17, 2015	DS	DP	U-853	
	6048859	Jun	29, 2012			U-745	
	6096742	Jul	01, 2018	DS	DP	U-745	
	6235735	Jun	29, 2012			U-747	
	6235735	Jun	29, 2012			U-746	
	7214692	Sep	18, 2012			U-853	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
APREPITANT - EMEND							
N021549 002	5538982	Jul	23, 2013		U-745	M-82	Mar 19, 2013
	5719147	Apr	17, 2015	DS DP	U-853		
	6048859	Jun	29, 2012		U-745		
	6096742	Jul	01, 2018	DS DP	U-745		
	6235735	Jun	29, 2012		U-747		
	6235735	Jun	29, 2012		U-746		
	7214692	Sep	18, 2012		U-853		
APREPITANT - EMEND							
N021549 003	5538982	Jul	23, 2013		U-745	M-82	Mar 19, 2013
	5719147	Apr	17, 2015	DS DP	U-853		
	6048859	Jun	29, 2012		U-745		
	6096742	Jul	01, 2018	DS DP	U-745		
	6235735	Jun	29, 2012		U-747		
	6235735	Jun	29, 2012		U-746		
	7214692	Sep	18, 2012		U-853		
ARbutamine Hydrochloride - GENESA							
N020420 001	5395970	Mar	07, 2012				
ARFORMOTEROL TARTRATE - BROVANA							
N021912 001	5795564	Apr	03, 2012		U-793		
	6040344	Nov	12, 2016	DS			
	6068833	Apr	03, 2012		U-793		
	6472563	Nov	09, 2021	DS			
	6589508	Apr	03, 2012		U-793		
	6720453	Nov	09, 2021	DS			
	6866839	Apr	03, 2012		U-793		
	7145036	Nov	09, 2021	DS			
ARGATROBAN - ARGATROBAN							
N020883 001	5214052	Jun	30, 2014				
ARGATROBAN - ARGATROBAN IN SODIUM CHLORIDE							
N022434 001	7589106	Sep	26, 2027	DP	U-1163		
	7687516	Sep	26, 2027	DP	U-1164		
ARIPIPRAZOLE - ABILIFY							
N021436 001	5006528	Oct	20, 2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20, 2015			I-616	Nov 19, 2012
						PED	Aug 27, 2011
ARIPIPRAZOLE - ABILIFY							
N021436 002	5006528	Oct	20, 2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20, 2015			I-616	Nov 19, 2012
						PED	Aug 27, 2011
ARIPIPRAZOLE - ABILIFY							
N021436 003	5006528	Oct	20, 2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20, 2015			I-616	Nov 19, 2012
						PED	Aug 27, 2011
ARIPIPRAZOLE - ABILIFY							
N021436 004	5006528	Oct	20, 2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20, 2015			I-616	Nov 19, 2012
						PED	Aug 27, 2011
ARIPIPRAZOLE - ABILIFY							
N021436 005	5006528	Oct	20, 2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20, 2015			I-616	Nov 19, 2012
						PED	Aug 27, 2011
ARIPIPRAZOLE - ABILIFY							
N021436 006	5006528	Oct	20, 2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20, 2015			I-616	Nov 19, 2012
						PED	Aug 27, 2011

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ARIPIPRAZOLE - ABILIFY</u>								
N021713 001	5006528	Oct	20,	2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20,	2015			PED	Aug 27, 2011
	6977257	Apr	24,	2022	DS DP			
	6977257*PED	Oct	24,	2022				
<u>ARIPIPRAZOLE - ABILIFY</u>								
N021729 002	5006528	Oct	20,	2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20,	2015			PED	Aug 27, 2011
<u>ARIPIPRAZOLE - ABILIFY</u>								
N021729 003	5006528	Oct	20,	2014	DS DP	U-761	I-633	Feb 16, 2014
	5006528*PED	Apr	20,	2015			PED	Aug 27, 2011
<u>ARIPIPRAZOLE - ABILIFY</u>								
N021729 004	5006528	Oct	20,	2014	DS DP	U-761	PED	Aug 27, 2011
	5006528*PED	Apr	20,	2015				
<u>ARIPIPRAZOLE - ABILIFY</u>								
N021729 005	5006528	Oct	20,	2014	DS DP	U-761	PED	Aug 27, 2011
	5006528*PED	Apr	20,	2015				
<u>ARIPIPRAZOLE - ABILIFY</u>								
N021866 001	5006528	Oct	20,	2014	DS DP	U-763	I-633	Feb 16, 2014
	5006528*PED	Apr	20,	2015			PED	Aug 27, 2011
	7115587	Jul	21,	2024	DS DP	U-764		
	7115587*PED	Jan	21,	2025				
	7550445	Jul	21,	2024	DP			
	7550445*PED	Jan	21,	2025				
<u>ARMODAFINIL - NUVIGIL</u>								
N021875 001	7132570	Dec	18,	2023	DS DP			
	7132570*PED	Jun	18,	2024				
	7297346	Nov	29,	2023	DP			
	7297346*PED	May	29,	2024				
	RE37516	Oct	06,	2014	DP	U-820		
	RE37516*PED	Apr	06,	2015				
<u>ARMODAFINIL - NUVIGIL</u>								
N021875 002	7132570	Dec	18,	2023	DS DP			
	7132570*PED	Jun	18,	2024				
	7297346	Nov	29,	2023	DP			
	7297346*PED	May	29,	2024				
	RE37516	Oct	06,	2014	DP	U-820		
	RE37516*PED	Apr	06,	2015				
<u>ARMODAFINIL - NUVIGIL</u>								
N021875 003	7132570	Dec	18,	2023	DS DP			
	7132570*PED	Jun	18,	2024				
	7297346	Nov	29,	2023	DP			
	7297346*PED	May	29,	2024				
	RE37516	Oct	06,	2014	DP	U-820		
	RE37516*PED	Apr	06,	2015				
<u>ARMODAFINIL - NUVIGIL</u>								
N021875 004	7132570	Dec	18,	2023	DS DP			
	7132570*PED	Jun	18,	2024				
	7297346	Nov	29,	2023	DP			
	7297346*PED	May	29,	2024				
	RE37516	Oct	06,	2014	DP	U-820		
	RE37516*PED	Apr	06,	2015				
<u>ARMODAFINIL - NUVIGIL</u>								
N021875 005	7132570	Dec	18,	2023	DS DP			
	7132570*PED	Jun	18,	2024				
	7297346	Nov	29,	2023	DP			
	7297346*PED	May	29,	2024				
	RE37516	Oct	06,	2014	DP	U-820		
	RE37516*PED	Apr	06,	2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ARSENIC TRIOXIDE - TRISENOX</u>							
N021248 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		
<u>ARTEMETHER; LUMEFANTRINE - COARTEM</u>							
N022268 001	5677331	Oct	14, 2014	DP	U-977	NCE	Apr 07, 2014
						ODE	Apr 07, 2016
<u>ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E - M.V.I.-12 (WITHOUT VITAMIN K)</u>							
N008809 006						ODE	Sep 09, 2011
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u>							
N021881 001	7169381	Sep	01, 2024	DS	DP		
	7658914	Sep	01, 2024	DS	DP		
<u>ASENAPINE MALEATE - SAPHRIS</u>							
N022117 001	5763476	Jun	09, 2015	DP	U-326	I-629	Sep 03, 2013
	7741358	Apr	06, 2026	DS	DP U-1064	I-628	Sep 03, 2013
						NCE	Aug 13, 2014
<u>ASENAPINE MALEATE - SAPHRIS</u>							
N022117 002	5763476	Jun	09, 2015	DP	U-326	I-629	Sep 03, 2013
	7741358	Apr	06, 2026	DS	DP U-1064	I-628	Sep 03, 2013
						NCE	Aug 13, 2014
<u>ASPIRIN; DIPYRIDAMOLE - AGGRENOX</u>							
N020884 001	6015577	Jan	18, 2017		U-302		
<u>ASPIRIN; PRAVASTATIN SODIUM - PRAVIGARD PAC (COPACKAGED)</u>							
N021387 001	5622985	Apr	22, 2014		U-335		
	5622985*PED	Oct	22, 2014		U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM - PRAVIGARD PAC (COPACKAGED)</u>							
N021387 002	5622985	Apr	22, 2014		U-335		
	5622985*PED	Oct	22, 2014		U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM - PRAVIGARD PAC (COPACKAGED)</u>							
N021387 003	5622985	Apr	22, 2014		U-335		
	5622985*PED	Oct	22, 2014		U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM - PRAVIGARD PAC (COPACKAGED)</u>							
N021387 004	5622985	Apr	22, 2014		U-335		
	5622985*PED	Oct	22, 2014		U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM - PRAVIGARD PAC (COPACKAGED)</u>							
N021387 005	5622985	Apr	22, 2014		U-335		
	5622985*PED	Oct	22, 2014		U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM - PRAVIGARD PAC (COPACKAGED)</u>							
N021387 006	5622985	Apr	22, 2014		U-335		
	5622985*PED	Oct	22, 2014		U-335		
<u>ATAZANAVIR SULFATE - REYATAZ</u>							
N021567 001	5849911	Jun	20, 2017	DS	DP U-167	D-130	Feb 04, 2014
	6087383	Dec	21, 2018	DS	DP	D-116	Sep 30, 2011
<u>ATAZANAVIR SULFATE - REYATAZ</u>							
N021567 002	5849911	Jun	20, 2017	DS	DP U-167	D-130	Feb 04, 2014
	6087383	Dec	21, 2018	DS	DP	D-116	Sep 30, 2011
<u>ATAZANAVIR SULFATE - REYATAZ</u>							
N021567 003	5849911	Jun	20, 2017	DS	DP U-167	D-130	Feb 04, 2014
	6087383	Dec	21, 2018	DS	DP	D-116	Sep 30, 2011

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ATAZANAVIR SULFATE - REYATAZ						
N021567 004	5849911	Jun 20, 2017	DS DP	U-167	D-130	Feb 04, 2014
	6087383	Dec 21, 2018	DS DP		D-116	Sep 30, 2011
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 001	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 002	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 003	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 004	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 005	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 006	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 007	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATOMOXETINE HYDROCHLORIDE - STRATTERA						
N021411 008	5658590	Nov 26, 2016	U-494	M-78	Jul 23, 2011	
	5658590*PED	May 26, 2017	U-494			
ATORVASTATIN CALCIUM - ATORVASTATIN CALCIUM						
A076477 001				PC	May 28, 2012	
ATORVASTATIN CALCIUM - ATORVASTATIN CALCIUM						
A076477 002				PC	May 28, 2012	
ATORVASTATIN CALCIUM - ATORVASTATIN CALCIUM						
A076477 003				PC	May 28, 2012	
ATORVASTATIN CALCIUM - ATORVASTATIN CALCIUM						
A076477 004				PC	May 28, 2012	
ATORVASTATIN CALCIUM - LIPITOR						
N020702 001	5686104	Nov 11, 2014	DP U-213			
	5686104*PED	May 11, 2015	U-213			
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013	DP			
	6126971*PED	Jul 19, 2013				
ATORVASTATIN CALCIUM - LIPITOR						
N020702 002	5686104	Nov 11, 2014	DP U-213			
	5686104*PED	May 11, 2015	U-213			
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013	DP			
	6126971*PED	Jul 19, 2013				
ATORVASTATIN CALCIUM - LIPITOR						
N020702 003	5686104	Nov 11, 2014	DP U-213			
	5686104*PED	May 11, 2015	U-213			
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013	DP			
	6126971*PED	Jul 19, 2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ATORVASTATIN CALCIUM - LIPITOR</u>							
N020702 004	5686104	Nov	11, 2014	DP	U-213		
	5686104*PED	May	11, 2015		U-213		
	5969156	Jul	08, 2016	DS			
	5969156*PED	Jan	08, 2017				
	6126971	Jan	19, 2013	DP			
	6126971*PED	Jul	19, 2013				
<u>ATOVAQUONE - MEPRON</u>							
N020500 001	6649659	Jul	10, 2016	DS	DP	U-69	
	6649659*PED	Jan	10, 2017				
<u>ATOVAQUONE; PROGUANIL HYDROCHLORIDE - MALARONE</u>							
N021078 001	5998449	Nov	25, 2013		U-990		
	5998449*PED	May	25, 2014				
	6166046	Nov	25, 2013		U-406		
	6166046*PED	May	25, 2014				
	6291488	Nov	25, 2013		U-406		
	6291488*PED	May	25, 2014				
<u>ATOVAQUONE; PROGUANIL HYDROCHLORIDE - MALARONE PEDIATRIC</u>							
N021078 002	5998449	Nov	25, 2013		U-990		
	5998449*PED	May	25, 2014				
	6166046	Nov	25, 2013				
	6166046*PED	May	25, 2014				
	6291488	Nov	25, 2013	U-406			
	6291488*PED	May	25, 2014				
<u>AVOBENZONE; ECAMSULE; OCTOCRYLENE - ANTHELIOS SX</u>							
N021502 001	5587150	Dec	24, 2013	DP	U-752		
<u>AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE - ANTHELIOS 40</u>							
N022009 002						NP	Oct 29, 2012
<u>AZELAIC ACID - FINACEA</u>							
N021470 001	6534070	Nov	18, 2018				
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u>							
N022203 001	8071073	Jun	04, 2028	DP		NP	Oct 15, 2011
<u>AZELASTINE HYDROCHLORIDE - ASTEPRO</u>							
N022371 001	8071073	Jun	04, 2028	DP		NP	Aug 31, 2012
<u>AZILSARTAN KAMEDOXOMIL - EDARBI</u>							
N200796 001	5583141	Dec	10, 2013	DS	DP U-3	NCE	Feb 25, 2016
	5736555	Jun	25, 2012	DS	DP U-3		
	5958961	Jun	06, 2014	DS	DP U-3		
	7157584	May	22, 2025	DS			
	7572920	Jan	07, 2025	DP	U-3		
<u>AZILSARTAN KAMEDOXOMIL - EDARBI</u>							
N200796 002	5583141	Dec	10, 2013	DS	DP U-3	NCE	Feb 25, 2016
	5736555	Jun	25, 2012	DS	DP U-3		
	5958961	Jun	06, 2014	DS	DP U-3		
	7157584	May	22, 2025	DS			
	7572920	Jan	07, 2025	DP	U-3		
<u>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</u>							
N202331 001						NCE	Feb 25, 2016
						NC	Dec 20, 2014
<u>AZITHROMYCIN - AZASITE</u>							
N050810 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		
<u>AZITHROMYCIN - ZITHROMAX</u>							
N050693 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
<u>AZITHROMYCIN - ZITHROMAX</u>						
N050710 001	6268489	Jul 31, 2018	DS			
<u>AZITHROMYCIN - ZITHROMAX</u>						
N050710 002	6268489	Jul 31, 2018	DS			
<u>AZITHROMYCIN - ZITHROMAX</u>						
N050711 001	6268489	Jul 31, 2018	DS			
<u>AZITHROMYCIN - ZITHROMAX</u>						
N050730 001	6268489	Jul 31, 2018	DS			
<u>AZITHROMYCIN - ZITHROMAX</u>						
N050733 001	6268489	Jul 31, 2018	DS			
<u>AZITHROMYCIN - ZITHROMAX</u>						
N050784 001	6268489	Jul 31, 2018	DS			
<u>AZITHROMYCIN - ZMAX</u>						
N050797 001	6068859	May 30, 2017	DP			
	6268489	Jul 31, 2018	DS			
	6984403	Feb 14, 2024	DP	U-282		
	7887844	Feb 14, 2024	DP			
<u>AZTREONAM - CAYSTON</u>						
N050814 001	7208141	Dec 20, 2021	DP	U-1031	ODE	Feb 22, 2017
	7214364	Dec 20, 2021	DP			
	7427633	Dec 20, 2021	DP	U-1031		
<u>BACLOFEN - KEMSTRO</u>						
N021589 001	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<u>BACLOFEN - KEMSTRO</u>						
N021589 002	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<u>BALSALAZIDE DISODIUM - COLAZAL</u>						
N020610 001	7452872	Aug 24, 2026	U-141		ODE	Dec 20, 2013
	7452872*PED	Feb 24, 2027			PED	Jun 20, 2014
	7625884	Aug 24, 2026	U-141			
	7625884*PED	Feb 24, 2027				
<u>BECLOMETHASONE DIPROPIONATE - QVAR 40</u>						
N020911 002	5605674	Feb 25, 2014				
	5683677	Nov 04, 2014				
	5776432	Jul 07, 2015				
<u>BECLOMETHASONE DIPROPIONATE - QVAR 80</u>						
N020911 001	5605674	Feb 25, 2014				
	5683677	Nov 04, 2014				
	5776432	Jul 07, 2015				
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N022249 001				I-580	Oct 31, 2011	
				NCE	Mar 20, 2013	
				ODE	Mar 20, 2015	
<u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
N022249 002				I-580	Oct 31, 2011	
				NCE	Mar 20, 2013	
				ODE	Mar 20, 2015	
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
N050819 001	5733886	Mar 31, 2015	DP	U-124		
	6117843	Feb 18, 2012	DP			
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - DUAC</u>						
N050741 001	5466446	Feb 16, 2014	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
BENZYL ALCOHOL - ULESFIA							
N022129 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		
BEPOTASTINE BESILATE - BEPREVE							
N022288 001	6780877	Dec	25, 2017	DS DP			
BESIFLOXACIN HYDROCHLORIDE - BESIVANCE							
N022308 001	5447926	Sep	05, 2012	DS DP	U-80		
	6685958	Jun	29, 2021		DP U-80		
	6699492	Mar	31, 2019		DP U-80		
BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX							
N021852 001	5763426	Jun	09, 2015	DS DP			
	6753013	Jan	27, 2020		DP U-88		
	6753013	Jan	27, 2020		DP U-193		
	RE39706	Jun	09, 2015	DS DP			
BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX SCALP							
N022185 001	6753013	Jan	27, 2020	DP U-88			
	6753013	Jan	27, 2020		DP U-193		
	6787529	Jan	27, 2020		DP U-193		
	6787529	Jan	27, 2020		DP U-88		
	RE39706	Jun	09, 2015	DS DP			
BETAMETHASONE VALERATE - LUXIQ							
N020934 001	6126920	Mar	01, 2016		U-484		
	7078058	May	24, 2017	DP			
BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE - BETOPTIC PILO							
N020619 001	5635172	Jun	03, 2014		U-191		
BEXAROTENE - TARGRETIN							
N021055 001	5780676	Jul	14, 2015		U-509		
	5962731	Oct	05, 2016		U-475		
	6043279	Apr	22, 2012		U-509		
	6320074	Apr	22, 2012	DS	U-509		
	7655699	Apr	22, 2012	DS DP	U-509		
BEXAROTENE - TARGRETIN							
N021056 001	5780676	Jul	14, 2015		U-510		
	5962731	Oct	05, 2016				
	6043279	Apr	22, 2012		U-510		
	6320074	Apr	22, 2012	DS	U-510		
	7655699	Apr	22, 2012	DS DP	U-510		
BICALUTAMIDE - CASODEX							
N020498 001						M-81	Dec 19, 2011
						PED	Jun 19, 2012
BIMATOPROST - LATISSE							
N022369 001	6403649	Sep	21, 2012	DS			
	7351404	May	25, 2024		U-939		
	7388029	Jan	21, 2022		U-938		
	8017655	Nov	27, 2012	DS DP			
	8038988	Aug	25, 2023	DS DP	U-1208		
BIMATOPROST - LUMIGAN							
N021275 001	5688819	Aug	19, 2014		U-446		
	6403649	Sep	21, 2012	DS DP	U-446		
	8017655	Nov	27, 2012	DS DP			
BIMATOPROST - LUMIGAN							
N022184 001	5688819	Aug	19, 2014		U-1081		
	6403649	Sep	21, 2012	DS	U-1081		
	7851504	Jun	13, 2027	DS DP			
	8017655	Nov	27, 2012	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
<u>BISACODYL - HALFLYTELY</u>						
	N021551 003 7291324	Oct 22, 2022	U-837		NP	Jul 16, 2013
<u>BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - HALFLYTELY</u>						
	N021551 002 7291324	Oct 22, 2022	U-837			
<u>BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE - PYLERA</u>						
	N050786 001 6350468	Dec 14, 2018	U-932			
	6350468	Dec 14, 2018	U-956			
<u>BIVALIRUDIN - ANGIOMAX</u>						
	N020873 001 5196404	Aug 13, 2012	DS DP U-1040			
	5196404*PED	Feb 13, 2013				
	7582727	Jul 27, 2028	DP			
	7582727*PED	Jan 27, 2029				
	7598343	Jul 27, 2028	DP			
	7598343*PED	Jan 27, 2029				
<u>BOCEPREVIR - VICTRELIS</u>						
	N202258 001 7012066	Feb 22, 2022	DS DP U-1128		NCE	May 13, 2016
	7772178	Nov 11, 2027	DP U-1128			
<u>BORTEZOMIB - VELCADE</u>						
	N021602 001 5780454	May 03, 2017	DP		ODE	Mar 25, 2012
	6083903	Oct 28, 2014	DP U-515			
	6297217	Oct 28, 2014	U-515			
	6297217	Oct 28, 2014	U-885			
	6297217	Oct 28, 2014	U-884			
	6617317	Oct 28, 2014	DS DP			
	6713446	Jan 25, 2022	DP			
	6747150	Oct 28, 2014	DP			
	6958319	Jan 25, 2022	DP			
	7119080	Oct 28, 2014	DP			
<u>BOSENTAN - TRACLEER</u>						
	N021290 001 5292740	Nov 20, 2015			I-607	Aug 07, 2012
<u>BOSENTAN - TRACLEER</u>						
	N021290 002 5292740	Nov 20, 2015			I-607	Aug 07, 2012
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
	N021262 001 5424078	Jun 13, 2012		Y		
	5424078*PED	Dec 13, 2012				
	6562873	Jul 10, 2021				
	6562873*PED	Jan 10, 2022				
	6627210	Jul 18, 2021	DP			
	6627210*PED	Jan 18, 2022				
	6641834	Jul 28, 2021	DP			
	6641834*PED	Jan 28, 2022				
	6673337	Jul 26, 2021	DP			
	6673337*PED	Jan 26, 2022				
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>						
	N021770 001 5424078	Jun 13, 2012	DP	Y		
	5424078*PED	Dec 13, 2012				
	6562873	Jul 10, 2021	DP			
	6562873*PED	Jan 10, 2022				
	6627210	Jul 18, 2021	DP			
	6627210*PED	Jan 18, 2022				
	6641834	Jul 28, 2021	DP			
	6641834*PED	Jan 28, 2022				
	6673337	Jul 26, 2021	DP			
	6673337*PED	Jan 26, 2022				
<u>BRIMONIDINE TARTRATE - BRIMONIDINE TARTRATE</u>						
	N021764 001 7265117	Aug 19, 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
<u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u>							
N021398 001	7030149	Apr	19, 2022		U-849		
	7320976	Apr	19, 2022		U-849		
	7323463	Jan	19, 2023	DP			
	7642258	Apr	19, 2022	DS DP	U-1024		
<u>BRINZOLAMIDE - AZOPT</u>							
N020816 001	5378703	Apr	01, 2012	DS	DP U-224		
	5378703*PED	Oct	01, 2012				
	5461081	Oct	24, 2012	DP	U-225		
	5461081*PED	Apr	24, 2013				
<u>BROMFENAC SODIUM - BROMDAY</u>							
N021664 002						NP	Oct 16, 2013
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>							
N020866 001	5468755	Nov	21, 2012		U-976		
	5679685	Oct	21, 2014	DP		NP	May 05, 2012
	5716957	Feb	10, 2015		U-976		
	5756513	Nov	21, 2012		U-976		
	5866584	Nov	21, 2012		U-976		
	7888310	Jul	25, 2023		U-976		
<u>BUDESONIDE - ENTOCORT EC</u>							
N021324 001	5643602	Jul	01, 2014		U-655		
	5643602*PED	Jan	01, 2015	DP			
<u>BUDESONIDE - PULMICORT FLEXHALER</u>							
N021949 001	6027714	Jan	09, 2018	DP	U-787		
	6142145	May	08, 2018	DP			
	6287540	Jan	09, 2018	DP			
	7143764	Mar	13, 2018	DP			
<u>BUDESONIDE - PULMICORT FLEXHALER</u>							
N021949 002	6027714	Jan	09, 2018	DP	U-787		
	6142145	May	08, 2018	DP			
	6287540	Jan	09, 2018	DP			
	7143764	Mar	13, 2018	DP			
<u>BUDESONIDE - PULMICORT RESPULES</u>							
N020929 001	6598603	Dec	23, 2018		U-529		
	6598603*PED	Jun	23, 2019				
	6899099	Dec	23, 2018		U-529		
	6899099*PED	Jun	23, 2019				
	7524834	Nov	11, 2018	DP	U-966		
	7524834*PED	May	11, 2019				
<u>BUDESONIDE - PULMICORT RESPULES</u>							
N020929 002	6598603	Dec	23, 2018		U-529		
	6598603*PED	Jun	23, 2019				
	6899099	Dec	23, 2018		U-529		
	6899099*PED	Jun	23, 2019				
	7524834	Nov	11, 2018	DP	U-966		
	7524834*PED	May	11, 2019				
<u>BUDESONIDE - PULMICORT RESPULES</u>							
N020929 003	6598603	Dec	23, 2018		U-529		
	6598603*PED	Jun	23, 2019				
	6899099	Dec	23, 2018		U-529		
	6899099*PED	Jun	23, 2019				
	7524834	Nov	11, 2018	DP	U-966		
	7524834*PED	May	11, 2019				
<u>BUDESONIDE - RHINOCORT</u>							
N020746 001	6291445	Apr	29, 2017			Y	
	6291445*PED	Oct	29, 2017				
	6686346	Apr	29, 2017	DP	U-557	Y	
	6686346*PED	Oct	29, 2017				
	6986904	Apr	29, 2017	DP	U-699	Y	
	6986904*PED	Oct	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
<u>BUDESONIDE - RHINOCORT</u>						
N020746 002	6291445	Apr 29, 2017				
	6291445*PED	Oct 29, 2017				
	6686346	Apr 29, 2017	DP U-557			
	6686346*PED	Oct 29, 2017				
	6986904	Apr 29, 2017	DP U-699			
	6986904*PED	Oct 29, 2017				
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N021929 001	5349945	Nov 12, 2011	DP		I-582	Feb 27, 2012
	5674860	Oct 07, 2014	DP U-1075			
	5972919	Dec 17, 2012	DP U-1075			
	6123924	Sep 26, 2017	DP			
	6641800	Sep 23, 2012	DP			
	7367333	Nov 11, 2018	DP			
	7587988	Apr 10, 2026	DP			
	7759328	Jan 29, 2023	DP U-1073			
	7967011	Aug 11, 2021	DP			
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
N021929 002	5349945	Nov 12, 2011	DP		I-582	Feb 27, 2012
	5674860	Oct 07, 2014	DP U-1075			
	5972919	Dec 17, 2012	DP U-1075			
	6123924	Sep 26, 2017	DP			
	6641800	Sep 23, 2012	DP			
	7367333	Nov 11, 2018	DP			
	7587988	Apr 10, 2026	DP			
	7759328	Jan 29, 2023	DP U-1073			
	7897646	Sep 09, 2018	U-1118			
	7967011	Aug 11, 2021	DP			
<u>BUPIVACAINE - EXPAREL</u>						
N022496 001					NP	Oct 28, 2014
<u>BUPIVACAINE - EXPAREL</u>						
N022496 002					NP	Oct 28, 2014
<u>BUPRENORPHINE - BUTRANS</u>						
N021306 001	6264980	Dec 18, 2015	DP		NDF	Jun 30, 2013
	6344211	Dec 18, 2015	U-1072			
	RE41408	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1072			
<u>BUPRENORPHINE - BUTRANS</u>						
N021306 002	6264980	Dec 18, 2015	DP		NDF	Jun 30, 2013
	6344211	Dec 18, 2015	U-1072			
	RE41408	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1072			
<u>BUPRENORPHINE - BUTRANS</u>						
N021306 003	6264980	Dec 18, 2015	DP		NDF	Jun 30, 2013
	6344211	Dec 18, 2015	U-1072			
	RE41408	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1072			
<u>BUPRENORPHINE; NALOXONE - SUBOXONE</u>						
N022410 001	8017150	Sep 10, 2023	DP		NDF	Aug 30, 2013
<u>BUPRENORPHINE; NALOXONE - SUBOXONE</u>						
N022410 002	8017150	Sep 10, 2023	DP		NDF	Aug 30, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
BUPROPION HYDROBROMIDE - APLENZIN						
N022108 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			
BUPROPION HYDROBROMIDE - APLENZIN						
N022108 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			
BUPROPION HYDROBROMIDE - APLENZIN						
N022108 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			
BUPROPION HYDROCHLORIDE - FORFIVO XL						
N022497 001	7674479	Jun 25, 2027	DP			
BUPROPION HYDROCHLORIDE - WELLBUTRIN SR						
N020358 001	5358970	Aug 12, 2013				
	5427798	Aug 12, 2013				
	5731000	Aug 12, 2013				
	5763493	Aug 12, 2013				
BUPROPION HYDROCHLORIDE - WELLBUTRIN SR						
N020358 002	5358970	Aug 12, 2013				
	5427798	Aug 12, 2013				
	5731000	Aug 12, 2013				
	5763493	Aug 12, 2013				
BUPROPION HYDROCHLORIDE - WELLBUTRIN SR						
N020358 003	5358970	Aug 12, 2013				
	5427798	Aug 12, 2013				
	5731000	Aug 12, 2013				
	5763493	Aug 12, 2013				
BUPROPION HYDROCHLORIDE - WELLBUTRIN SR						
N020358 004	5358970	Aug 12, 2013				
	5427798	Aug 12, 2013				
	5731000	Aug 12, 2013				
	5763493	Aug 12, 2013				
BUPROPION HYDROCHLORIDE - WELLBUTRIN XL						
N021515 001	6096341	Oct 30, 2018				
BUPROPION HYDROCHLORIDE - WELLBUTRIN XL						
N021515 002	6096341	Oct 30, 2018				
BUPROPION HYDROCHLORIDE - ZYBAN						
N020711 002	5358970	Aug 12, 2013				
	5427798	Aug 12, 2013				
	5731000	Aug 12, 2013				
	5763493	Aug 12, 2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>BUPROPION HYDROCHLORIDE - ZYBAN</u>						
N020711 003	5358970	Aug 12, 2013				
	5427798	Aug 12, 2013				
	5731000	Aug 12, 2013				
	5763493	Aug 12, 2013				
<u>BUSULFAN - BUSULFEX</u>						
N020954 001	5430057	Sep 30, 2013		U-263		
	5430057*PED	Mar 30, 2014		U-263		
	5559148	Sep 30, 2013		U-264		
	5559148*PED	Mar 30, 2014		U-264		
<u>BUTOCONAZOLE NITRATE - GYNAZOLE-1</u>						
N019881 001	5993856	Nov 17, 2017	DP	U-457		
<u>CABAZITAXEL - JEVTANA KIT</u>						
N201023 001	5438072	Nov 22, 2013	DP			
	5698582	Jul 03, 2012	DP			
	5847170	Mar 26, 2016	DS	DP		
	6331635	Mar 26, 2016	DS	DP		
	6372780	Mar 26, 2016		U-1067		
	6387946	Mar 26, 2016		U-1067		
	7241907	Dec 10, 2025	DS			
<u>CALCIPOTRIENE - DOVONEX</u>						
N020554 001	5763426	Jun 09, 2015	DS	DP		
	RE39706	Jun 09, 2015	DS	DP		
<u>CALCIPOTRIENE - DOVONEX</u>						
N020611 001	5763426	Jun 09, 2015	DS	DP		
	RE39706	Jun 09, 2015	DS	DP		
<u>CALCIPOTRIENE - SORILUX</u>						
N022563 001					NDF	Oct 06, 2013
<u>CALCITONIN SALMON - MIACALCIN</u>						
N020313 002	5733569	Mar 31, 2015		U-227		
	5759565	Mar 31, 2015				
<u>CALCITONIN SALMON RECOMBINANT - FORTICAL</u>						
N021406 001	6440392	Feb 02, 2021	DP	U-227		
	RE40812	Feb 02, 2021	DP			
<u>CALCITRIOL - CALCIJEX</u>						
N018874 001	6051567	Aug 02, 2019				
	6051567*PED	Feb 02, 2020				
	6265392	Aug 02, 2019				
	6265392*PED	Feb 02, 2020				
	6274169	Aug 02, 2019				
	6274169*PED	Feb 02, 2020				
<u>CALCITRIOL - CALCIJEX</u>						
N018874 002	6051567	Aug 02, 2019				
	6051567*PED	Feb 02, 2020				
	6265392	Aug 02, 2019				
	6265392*PED	Feb 02, 2020				
	6274169	Aug 02, 2019				
	6274169*PED	Feb 02, 2020				
<u>CALCITRIOL - VECTICAL</u>						
N022087 001					NDF	Jan 23, 2012
<u>CALCIUM ACETATE - PHOSLO</u>						
N021160 002	6576665	Apr 03, 2021				
<u>CALCIUM ACETATE - PHOSLO GELCAPS</u>						
N021160 003	6576665	Apr 03, 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
<u>CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE - PEPCID COMPLETE</u>						
N020958 001	5229137	May 16, 2012		U-349		
	5229137*PED	Nov 16, 2012				
	5989588	Sep 30, 2015		U-349		
	5989588*PED	Mar 30, 2016		U-349		
	6814978	Aug 26, 2021	DP			
	6814978*PED	Feb 26, 2022				
<u>CALCIUM CARBONATE; RISEDRONATE SODIUM - ACTONEL WITH CALCIUM (COPACKAGED)</u>						
N021823 001	5583122	Dec 10, 2013	DS DP	U-353		
	5583122*PED	Jun 10, 2014				
	5994329	Jul 17, 2018		U-353		
	5994329*PED	Jan 17, 2019				
	6015801	Jul 17, 2018		U-353		
	6015801*PED	Jan 17, 2019				
	6096342	Nov 21, 2011	DP			
	6096342*PED	May 21, 2012				
	6165513	Jun 10, 2018	DP			
	6165513*PED	Dec 10, 2018				
	6432932	Jul 17, 2018		U-595		
	6432932*PED	Jan 17, 2019				
	6465443	Aug 14, 2018	DP			
	6465443*PED	Feb 14, 2019				
<u>CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - NAVSTEL</u>						
N022193 001	5409904	Apr 25, 2012	DP	U-891	NP	
	5578578	Apr 25, 2012	DP			
	7084130	Nov 29, 2021	DP	U-891		
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
N020838 001	5196444	Jun 04, 2012	DS DP	U-660		
	5196444	Jun 04, 2012	DS DP	U-3		
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
N020838 002	5196444	Jun 04, 2012	DS DP	U-3		
	5196444	Jun 04, 2012	DS DP	U-660		
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
N020838 003	5196444	Jun 04, 2012	DS DP	U-3		
	5196444	Jun 04, 2012	DS DP	U-660		
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
N020838 004	5196444	Jun 04, 2012	DS DP	U-660		
	5196444	Jun 04, 2012	DS DP	U-3		
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	7538133*PED	Oct 18, 2011				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE - ATACAND HCT						
N021093 001	5196444	Jun 04, 2012	DS DP U-3			
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	5721263	Feb 24, 2015	DP U-3			
	5958961	Jun 06, 2014	DP U-3			
	7538133*PED	Oct 18, 2011				
CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE - ATACAND HCT						
N021093 002	5196444	Jun 04, 2012	DS DP U-3			
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	5721263	Feb 24, 2015	DP U-3			
	5958961	Jun 06, 2014	DP U-3			
	7538133*PED	Oct 18, 2011				
CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE - ATACAND HCT						
N021093 003	5196444	Jun 04, 2012	DS DP U-3			
	5196444*PED	Dec 04, 2012				
	5534534	Jul 09, 2013	DP			
	5534534*PED	Jan 09, 2014				
	5705517*PED	Oct 18, 2011				
	5721263	Feb 24, 2015	DP U-3			
	5958961	Jun 06, 2014	DP U-3			
	7538133*PED	Oct 18, 2011				
CAPECITABINE - XELODA						
N020896 001	5472949	Dec 14, 2013		U-271		
CAPECITABINE - XELODA						
N020896 002	5472949	Dec 14, 2013		U-271		
CAPSaicin - QUTENZA						
N022395 001	6239180	Nov 06, 2016	DP		NCE	Nov 16, 2014
					ODE	Nov 16, 2016
CARBAMAZEPINE - CARBAMAZEPINE						
A076697 003					PC	Nov 16, 2011
CARBAMAZEPINE - CARBATROL						
N020712 001	5326570	Jul 23, 2011	U-215			
	5912013	Jun 15, 2016	U-277			
CARBAMAZEPINE - CARBATROL						
N020712 002	5326570	Jul 23, 2011	U-215			
	5912013	Jun 15, 2016	U-277			
CARBAMAZEPINE - CARBATROL						
N020712 003	5326570	Jul 23, 2011	U-215			
	5912013	Jun 15, 2016	U-277			
CARBAMAZEPINE - EQUETRO						
N021710 001	5326570	Jul 23, 2011	DP U-627			
	5912013	Jun 15, 2016	DP			
	6977253	May 19, 2024	U-693			
CARBAMAZEPINE - EQUETRO						
N021710 002	5326570	Jul 23, 2011	DP U-627			
	5912013	Jun 15, 2016	DP			
	6977253	May 19, 2024	U-693			
CARBAMAZEPINE - EQUETRO						
N021710 003	5326570	Jul 23, 2011	DP U-627			
	5912013	Jun 15, 2016	DP			
	6977253	May 19, 2024	U-693			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100						
N021485 002	5446194	Oct 19, 2013	DS			
	6500867	Jun 29, 2020	DP	U-219		
	6797732	Jun 29, 2020	DP			
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 125						
N021485 006	5446194	Oct 19, 2013	DS			
	6500867	Jun 29, 2020	DP	U-219		
	6797732	Jun 29, 2020	DP			
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150						
N021485 003	5446194	Oct 19, 2013	DS			
	6500867	Jun 29, 2020	DP	U-219		
	6797732	Jun 29, 2020	DP			
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 200						
N021485 004	5446194	Oct 19, 2013	DS			
	6500867	Jun 29, 2020	DP	U-219		
	6797732	Jun 29, 2020	DP			
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 50						
N021485 001	5446194	Oct 19, 2013	DS			
	6500867	Jun 29, 2020	DP	U-219		
	6797732	Jun 29, 2020	DP			
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 75						
N021485 005	5446194	Oct 19, 2013	DS			
	6500867	Jun 29, 2020	DP	U-219		
	6797732	Jun 29, 2020	DP			
CARGLUMIC ACID - CARBAGLU						
N022562 001					NCE	Mar 18, 2015
					ODE	Mar 18, 2017
CARVEDILOL - COREG						
N020297 001	RE40000	Jun 07, 2015	U-233			
	RE40000*PED	Dec 07, 2015				
CARVEDILOL - COREG						
N020297 002	RE40000	Jun 07, 2015	U-233			
	RE40000*PED	Dec 07, 2015				
CARVEDILOL - COREG						
N020297 003	RE40000	Jun 07, 2015	U-233			
	RE40000*PED	Dec 07, 2015				
CARVEDILOL - COREG						
N020297 004	RE40000	Jun 07, 2015	U-233			
	RE40000*PED	Dec 07, 2015				
CARVEDILOL PHOSPHATE - COREG CR						
N022012 001	5902821	Feb 07, 2016	U-313			
	5902821	Feb 07, 2016	U-777			
	5902821*PED	Aug 07, 2016				
	6022562	Oct 17, 2015	DP			
	6022562*PED	Apr 17, 2016				
	7268156	Jun 27, 2023	DS	DP U-313		
	7268156	Jun 27, 2023	DS	DP U-3		
	7268156*PED	Dec 27, 2023				
	RE40000	Jun 07, 2015	U-777			
	RE40000*PED	Dec 07, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
CARVEDILOL PHOSPHATE - COREG CR							
N022012 002	5902821	Feb	07, 2016		U-313		
	5902821	Feb	07, 2016		U-777		
	5902821*PED	Aug	07, 2016				
	6022562	Oct	17, 2015	DP			
	6022562*PED	Apr	17, 2016				
	7268156	Jun	27, 2023	DS DP	U-3		
	7268156	Jun	27, 2023	DS DP	U-313		
	7268156*PED	Dec	27, 2023				
	RE40000	Jun	07, 2015		U-777		
	RE40000*PED	Dec	07, 2015				
CARVEDILOL PHOSPHATE - COREG CR							
N022012 003	5902821	Feb	07, 2016		U-777		
	5902821	Feb	07, 2016		U-313		
	5902821*PED	Aug	07, 2016				
	6022562	Oct	17, 2015	DP			
	6022562*PED	Apr	17, 2016				
	7268156	Jun	27, 2023	DS DP	U-313		
	7268156	Jun	27, 2023	DS DP	U-3		
	7268156*PED	Dec	27, 2023				
	RE40000	Jun	07, 2015		U-777		
	RE40000*PED	Dec	07, 2015				
CARVEDILOL PHOSPHATE - COREG CR							
N022012 004	5902821	Feb	07, 2016		U-777		
	5902821	Feb	07, 2016		U-313		
	5902821*PED	Aug	07, 2016				
	6022562	Oct	17, 2015	DP			
	6022562*PED	Apr	17, 2016				
	7268156	Jun	27, 2023	DS DP	U-313		
	7268156	Jun	27, 2023	DS DP	U-3		
	7268156*PED	Dec	27, 2023				
	RE40000	Jun	07, 2015		U-777		
	RE40000*PED	Dec	07, 2015				
CASPOFUNGIN ACETATE - CANCIDAS							
N021227 001	5378804	Mar	16, 2013	DS			
	5378804*PED	Sep	16, 2013				
	5514650	Jan	26, 2015	DP	U-607		
	5514650*PED	Jul	26, 2015				
	5792746	Mar	16, 2013	DS DP	U-607		
	5792746*PED	Sep	16, 2013				
	5952300	Mar	28, 2017	DP			
	5952300*PED	Sep	28, 2017				
	6136783	Mar	28, 2017		U-607		
	6136783*PED	Sep	28, 2017				
CASPOFUNGIN ACETATE - CANCIDAS							
N021227 002	5378804	Mar	16, 2013	DS			
	5378804*PED	Sep	16, 2013				
	5514650	Jan	26, 2015	DP	U-607		
	5514650*PED	Jul	26, 2015				
	5792746	Mar	16, 2013	DS DP	U-607		
	5792746*PED	Sep	16, 2013				
	5952300	Mar	28, 2017	DP			
	5952300*PED	Sep	28, 2017				
	6136783	Mar	28, 2017		U-607		
	6136783*PED	Sep	28, 2017				
CEFDINIR - OMNICEF							
N050739 001	4935507	Dec	04, 2011	DS			
CEFDINIR - OMNICEF							
N050749 001	4935507	Dec	04, 2011	DS			
CEFDINIR - OMNICEF							
N050749 002	4935507	Dec	04, 2011	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
<u>CEFDITOREN PIVOXIL - SPECTRACEF</u>						
N021222 001	5958915	Oct 14, 2016				
<u>CEFDITOREN PIVOXIL - SPECTRACEF</u>						
N021222 002	5958915	Oct 14, 2016	DP			
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N200327 001	6417175	Dec 17, 2018	DS	DP	U-282	
	6906055	Dec 15, 2021	DS	DP		
	7419973	Dec 15, 2021		DP		
<u>CEFTAROLINE FOSAMIL - TEFLARO</u>						
N200327 002	6417175	Dec 17, 2018	DS	DP	U-282	
	6906055	Dec 15, 2021	DS	DP		
	7419973	Dec 15, 2021		DP		
<u>CEFTIBUTEN DIHYDRATE - CEDAX</u>						
N050685 002	5599557	Apr 30, 2013	DP			
<u>CEFTIBUTEN DIHYDRATE - CEDAX</u>						
N050686 001	5599557	Feb 04, 2014	DP	U-282		
	5599557	Feb 04, 2014	DP	U-578		
<u>CEFTIBUTEN DIHYDRATE - CEDAX</u>						
N050686 002	5599557	Feb 04, 2014	DP	U-578		
	5599557	Feb 04, 2014	DP	U-282		
<u>CELECOXIB - CELEBREX</u>						
N020998 001	5466823	Nov 30, 2013	DS			
	5466823*PED	May 30, 2014				
	5563165	Nov 30, 2013	DP			
	5563165*PED	May 30, 2014				
	5760068	Jun 02, 2015		U-672		
	5760068*PED	Dec 02, 2015				
<u>CELECOXIB - CELEBREX</u>						
N020998 002	5466823	Nov 30, 2013	DS			
	5466823*PED	May 30, 2014				
	5563165	Nov 30, 2013	DP			
	5563165*PED	May 30, 2014				
	5760068	Jun 02, 2015		U-672		
	5760068*PED	Dec 02, 2015				
<u>CELECOXIB - CELEBREX</u>						
N020998 003	5466823	Nov 30, 2013	DS			
	5466823*PED	May 30, 2014				
	5563165	Nov 30, 2013	DP			
	5563165*PED	May 30, 2014				
	5760068	Jun 02, 2015		U-672		
	5760068*PED	Dec 02, 2015				
<u>CELECOXIB - CELEBREX</u>						
N020998 004	5466823	Nov 30, 2013	DS			
	5466823*PED	May 30, 2014				
	5563165	Nov 30, 2013	DP			
	5563165*PED	May 30, 2014				
	5760068	Jun 02, 2015		U-672		
	5760068*PED	Dec 02, 2015				
<u>CERIVASTATIN SODIUM - BAYCOL</u>						
N020740 001	5177080	Nov 26, 2011				
<u>CERIVASTATIN SODIUM - BAYCOL</u>						
N020740 002	5177080	Nov 26, 2011				
<u>CERIVASTATIN SODIUM - BAYCOL</u>						
N020740 003	5177080	Nov 26, 2011				
<u>CERIVASTATIN SODIUM - BAYCOL</u>						
N020740 004	5177080	Nov 26, 2011				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>CERIVASTATIN SODIUM - BAYCOL</u>						
	N020740 005 5177080	Nov 26, 2011				
<u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC ALLERGY</u>						
	N021621 003 6455533	Jul 02, 2018	DP U-295			
<u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC ALLERGY</u>						
	N021621 004 6455533	Jul 02, 2018	DP U-295			
<u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC HIVES RELIEF</u>						
	N021621 005 6455533	Jul 02, 2018	DP U-295			
<u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC HIVES RELIEF</u>						
	N021621 006 6455533	Jul 02, 2018	DP U-295			
<u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u>						
	N021150 002 6469009	Jul 13, 2019	DP U-295			
	6489329	Apr 08, 2016	DP			
	7014867	Jun 10, 2022	DP			
	7226614	Jun 10, 2022	U-295			
<u>CETRORELIX - CETROTIDE</u>						
	N021197 001 6319192	Apr 23, 2019	U-426			
	6863891	Feb 22, 2014	U-426			
	7605121	Feb 22, 2014	DP			
<u>CETRORELIX - CETROTIDE</u>						
	N021197 002 6319192	Apr 23, 2019	U-426			
	6863891	Feb 22, 2014	U-426			
	7605121	Feb 22, 2014	DP			
<u>CEVIMELINE HYDROCHLORIDE - EVOXAC</u>						
	N020989 002 5340821	Jul 07, 2013	U-309			
<u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u>						
	N021669 001 7066916	Feb 17, 2024	U-737			
	7427574	Apr 25, 2026	DP			
	7595021	May 12, 2023	DP U-1022			
	7935093	Oct 02, 2027	DP U-1022			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
	N020832 001 5690958	Sep 30, 2016	DP			
	6536975	Nov 10, 2020	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
	N020832 004 5690958	Sep 30, 2016	DP			
	6536975	Nov 10, 2020	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
	N020832 006 5690958	Sep 30, 2016	DP			
	6991394	Jan 31, 2024	DP			
	7182536	Dec 30, 2023	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP FREPP</u>						
	N020832 003 5538353	Aug 25, 2015	DP			
	5690958	Sep 30, 2016	DP			
	5752363	Apr 22, 2017	DP			
	5772346	Apr 22, 2017	DP			
	D386849	Nov 25, 2011	DP			
	D396911	Aug 11, 2012	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP SEPP</u>						
	N021555 001 5690958	Sep 30, 2016	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
	N020832 002 5690958	Sep 30, 2016	DP			
	6729786	Mar 14, 2023	DP			
	6991393	Mar 14, 2023	DP			
	6991394	Jan 31, 2024	DP			
	7182536	Dec 30, 2023	DP			
	7241065	Mar 14, 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
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>							
N020832 005	5690958	Sep	30, 2016	DP			
	6536975	Nov	10, 2020	DP			
	6729786	Mar	14, 2023	DP			
	6991393	Jan	31, 2024	DP			
	7241065	Mar	14, 2023	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>							
N020832 007	5690958	Sep	30, 2016	DP			
	6536975	Nov	10, 2020	DP			
	6729786	Mar	14, 2023	DP			
	6991393	Mar	14, 2023	DP			
	7241065	Mar	14, 2023	DP			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORASCRUB MAXI SWABSTICK</u>							
N021524 003	D468424	Jan	07, 2017				
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - ADVIL ALLERGY SINUS</u>							
N021441 001	7863287	Feb	28, 2027	DP			
<u>CHOLINE FENOFOBRATE - TRILIPIX</u>							
N022224 001	7259186	Jan	07, 2025	DS		NP	Dec 15, 2011
<u>CHOLINE FENOFOBRATE - TRILIPIX</u>							
N022224 002	7259186	Jan	07, 2025	DS		NP	Dec 15, 2011
<u>CHORIOGONADOTROPIN ALFA - OVIDREL</u>							
N021149 001	5767251	Jun	16, 2015				
<u>CHORIOGONADOTROPIN ALFA - OVIDREL</u>							
N021149 002	5767251	Jun	16, 2015	DS			
	6706681	Mar	16, 2021	DP			
<u>CICLESONIDE - ALVESCO</u>							
N021658 002	5482934	Oct	24, 2017	DS	DP U-1002	NCE	Oct 20, 2011
	5605674	Feb	25, 2014	DP			
	5683677	Nov	04, 2014	DP			
	5775321	Jul	07, 2015	DP			
	6006745	Dec	28, 2016	DP			
	6036942	Apr	30, 2013	DP			
	6120752	May	13, 2018	DP			
	6264923	May	13, 2018	DP			
<u>CICLESONIDE - ALVESCO</u>							
N021658 003	5482934	Oct	24, 2017	DS	DP U-1002	NCE	Oct 20, 2011
	5605674	Feb	25, 2014	DP			
	5683677	Nov	04, 2014	DP			
	5775321	Jul	07, 2015	DP			
	6006745	Dec	28, 2016	DP			
	6036942	Apr	30, 2013	DP			
	6120752	May	13, 2018	DP			
	6264923	May	13, 2018	DP			
<u>CICLESONIDE - OMNARIS</u>							
N022004 001	5482934	Oct	24, 2017	DS	DP U-557	NCE	Oct 20, 2011
	6767901	Oct	21, 2020	DP			
	6939559	Apr	21, 2019	DP			
	7235247	Apr	21, 2019	DP			
<u>CICLOPIROX - LOPROX</u>							
N020519 001	7018656	Sep	05, 2018	DP			
	7026337	Nov	21, 2016	U-714			
<u>CICLOPIROX - LOPROX</u>							
N021159 001	7981909	Sep	16, 2017		U-1162		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>CINACALCET HYDROCHLORIDE - SENSIPIAR</u>							
N021688 001	6011068	Mar	08, 2018	DS DP		I-634	Feb 25, 2014
	6031003	Dec	14, 2016		U-559	M-101	Feb 25, 2014
	6211244	Oct	23, 2015	DS DP	U-560	ODE	Feb 25, 2018
	6313146	Dec	14, 2016	DS DP			
	7829595	Sep	22, 2026	DP	U-1098		
<u>CINACALCET HYDROCHLORIDE - SENSIPIAR</u>							
N021688 002	6011068	Mar	08, 2018	DS DP		I-634	Feb 25, 2014
	6031003	Dec	14, 2016		U-559	M-101	Feb 25, 2014
	6211244	Oct	23, 2015	DS DP	U-560	ODE	Feb 25, 2018
	6313146	Dec	14, 2016	DS DP			
	7829595	Sep	22, 2026	DP	U-1098		
<u>CINACALCET HYDROCHLORIDE - SENSIPIAR</u>							
N021688 003	6011068	Mar	08, 2018	DS DP		I-634	Feb 25, 2014
	6031003	Dec	14, 2016		U-559	M-101	Feb 25, 2014
	6211244	Oct	23, 2015	DS DP	U-560	ODE	Feb 25, 2018
	6313146	Dec	14, 2016	DS DP			
	7829595	Sep	22, 2026	DP	U-1098		
<u>CIPROFLOXACIN - CIPRO</u>							
N020780 001	5695784	Dec	09, 2014				
	5695784*PED	Jun	09, 2015				
	6136347	Jan	06, 2013		U-362		
	6136347*PED	Jul	06, 2013				
<u>CIPROFLOXACIN - CIPRO</u>							
N020780 002	5695784	Dec	09, 2014				
	5695784*PED	Jun	09, 2015				
	6136347	Jan	06, 2013		U-362		
	6136347*PED	Jul	06, 2013				
<u>CIPROFLOXACIN HYDROCHLORIDE - CETRAXAL</u>							
N021918 001						NDF	May 01, 2012
<u>CIPROFLOXACIN HYDROCHLORIDE - CIPRO</u>							
N019537 001	5286754*PED	Aug	15, 2011				
<u>CIPROFLOXACIN HYDROCHLORIDE - CIPRO</u>							
N019537 002	5286754*PED	Aug	15, 2011				
<u>CIPROFLOXACIN HYDROCHLORIDE - CIPRO</u>							
N019537 003	5286754*PED	Aug	15, 2011				
<u>CIPROFLOXACIN HYDROCHLORIDE - CIPRO</u>							
N019537 004	5286754*PED	Aug	15, 2011				
<u>CIPROFLOXACIN HYDROCHLORIDE - PROQUIN XR</u>							
N021744 001	5972389	Sep	19, 2016	DP	U-663		
	6340475	Sep	19, 2016	DP	U-663		
	6488962	Jun	20, 2020	DP			
	6635280	Sep	19, 2016	DP	U-663		
<u>CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE - CIPRO HC</u>							
N020805 001	5843930	Jun	06, 2015	DP	U-646		
	5965549	Jun	06, 2015	DP			
<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u>							
N021473 001	7709022	Jun	23, 2021	DP			
	7709022*PED	Dec	23, 2021	DP			
<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</u>							
N021473 002	7709022	Jun	23, 2021	DP			
	7709022*PED	Dec	23, 2021	DP			
<u>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</u>							
N021537 001	6284804	Aug	10, 2020				
	6359016	Aug	10, 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>CISAPRIDE MONOHYDRATE - PROPULSID QUICKSOLV</u>						
N020767 001	5648093	Jul 15, 2014				
<u>CISATRACURIUM BESYLATE - NIMBEX</u>						
N020551 001	5453510	Sep 26, 2012		U-127		
<u>CISATRACURIUM BESYLATE - NIMBEX PRESERVATIVE FREE</u>						
N020551 002	5453510	Sep 26, 2012		U-127		
<u>CISATRACURIUM BESYLATE - NIMBEX PRESERVATIVE FREE</u>						
N020551 003	5453510	Sep 26, 2012		U-127		
<u>CLARITHROMYCIN - BIAXIN XL</u>						
N050775 001	6010718	Apr 11, 2017	DP	U-924		
	6551616	Jul 15, 2017		U-924		
<u>CLEVIDIPIINE BUTYRATE - CLEVIPREX</u>						
N022156 001	5739152	Apr 14, 2015	DP	U-893	NCE	Aug 01, 2013
	5856346	Jan 05, 2016	DS	DP U-893		
<u>CLEVIDIPIINE BUTYRATE - CLEVIPREX</u>						
N022156 002	5739152	Apr 14, 2015	DP	U-893	NCE	Aug 01, 2013
	5856346	Jan 05, 2016	DS	DP U-893		
<u>CLINDAMYCIN PHOSPHATE - CLEOCIN</u>						
N050767 001	6495157	Jul 20, 2020	DP			
<u>CLINDAMYCIN PHOSPHATE - CLINDAGEL</u>						
N050782 001	6387383	Aug 03, 2020	DP	U-818		
<u>CLINDAMYCIN PHOSPHATE - CLINDESSE</u>						
N050793 001	5993856	Nov 17, 2017	DP	U-137		
	6899890	Apr 27, 2023	DP	U-137		
<u>CLINDAMYCIN PHOSPHATE - EVOCLIN</u>						
N050801 001	7141237	Jan 23, 2024	DS	DP		
	7374747	Aug 09, 2026	DS	DP U-921		
<u>CLINDAMYCIN PHOSPHATE; TRETINOIN - VELTIN</u>						
N050803 001	5690923	Nov 25, 2014	DP			
<u>CLINDAMYCIN PHOSPHATE; TRETINOIN - ZIANA</u>						
N050802 001	6387383	Aug 03, 2020	DP	U-916		
	RE41134	Feb 24, 2015	DP	U-1033		
<u>CLOBAZAM - ONFI</u>						
N202067 001					NCE	Oct 21, 2016
					ODE	Oct 21, 2018
<u>CLOBAZAM - ONFI</u>						
N202067 002					NCE	Oct 21, 2016
					ODE	Oct 21, 2018
<u>CLOBAZAM - ONFI</u>						
N202067 003					NCE	Oct 21, 2016
					ODE	Oct 21, 2018
<u>CLOBETASOL PROPIONATE - CLOBEX</u>						
N021535 001	6106848	Sep 22, 2017				
<u>CLOBETASOL PROPIONATE - CLOBEX</u>						
N021644 001	7316810	Jun 17, 2019	DP			
	7700081	Jan 03, 2022		U-1044		
	8066975	Jun 17, 2019	DP			
	8066976	Jun 17, 2019	DP			
<u>CLOBETASOL PROPIONATE - CLOBEX</u>						
N021835 001	5972920	Feb 12, 2018	DP			
	5990100	Mar 24, 2018	DP	U-742		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>CLOBETASOL PROPIONATE - OLUX</u>						
N021142 001	6126920	Mar 01, 2016		U-484		
<u>CLOBETASOL PROPIONATE - OLUX E</u>						
N022013 001	6730288	Sep 08, 2019	DP			
	7029659	Sep 08, 2019	DP			
<u>CLOFARABINE - CLOLAR</u>						
N021673 001	5661136	Jan 14, 2018		U-626	ODE	Dec 28, 2011
	5661136*PED	Jul 14, 2018			PED	Jun 28, 2012
<u>CLONIDINE HYDROCHLORIDE - JENLOGA</u>						
N022331 001	5869100	Oct 13, 2013	DP		NP	Sep 29, 2012
<u>CLONIDINE HYDROCHLORIDE - JENLOGA</u>						
N022331 002	5869100	Oct 13, 2013	DP			
<u>CLONIDINE HYDROCHLORIDE - KAPVAY</u>						
N022331 003	5869100	Oct 13, 2013	DP		NP	Sep 28, 2013
<u>CLONIDINE HYDROCHLORIDE - KAPVAY</u>						
N022331 004	5869100	Oct 13, 2013	DP		NP	Sep 28, 2013
<u>CLOPIDOGREL BISULFATE - PLAVIX</u>						
N020839 001	4847265	Nov 17, 2011	DS DP		M-61	May 06, 2014
	4847265*PED	May 17, 2012			PED	Nov 06, 2014
	5576328	Jan 31, 2014		U-432	Y	
	5576328*PED	Jul 31, 2014				
	6429210	Jun 10, 2019	DS DP			
	6429210*PED	Dec 10, 2019				
	6504030	Jun 10, 2019	DS			
	6504030*PED	Dec 10, 2019				
<u>CLOPIDOGREL BISULFATE - PLAVIX</u>						
N020839 002	4847265	Nov 17, 2011	DS DP		M-61	May 06, 2014
	4847265*PED	May 17, 2012			PED	Nov 06, 2014
	6429210	Jun 10, 2019	DS DP			
	6429210*PED	Dec 10, 2019				
	6504030	Jun 10, 2019	DS			
	6504030*PED	Dec 10, 2019				
<u>CLOZAPINE - FAZACLO ODT</u>						
N021590 001	6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<u>CLOZAPINE - FAZACLO ODT</u>						
N021590 002	6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<u>CLOZAPINE - FAZACLO ODT</u>						
N021590 003	6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<u>CLOZAPINE - FAZACLO ODT</u>						
N021590 004	6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<u>CLOZAPINE - FAZACLO ODT</u>						
N021590 005	6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<u>CLOZAPINE - FAZACLO ODT</u>						
N021590 006	6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 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
<u>COLCHICINE - COLCRYS</u>						
N022352 001	7601758	Feb 10, 2029	U-1007	I-603	Jul 30, 2012	
	7619004	Dec 03, 2028	U-1020	ODE	Jul 29, 2016	
	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	Feb 17, 2029	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			
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N021141 001	5607669	Jun 10, 2014	U-323			
	5607669*PED	Dec 10, 2014				
	5679717	Apr 29, 2014	U-323			
	5679717*PED	Oct 29, 2014				
	5693675	Dec 02, 2014				
	5693675*PED	Jun 02, 2015				
	5917007	Apr 29, 2014	U-323			
	5917007*PED	Oct 29, 2014				
	5919832	Jun 10, 2014				
	5919832*PED	Dec 10, 2014				
	6066678	Jun 10, 2014	U-323			
	6066678*PED	Dec 10, 2014				
	6433026	Jun 10, 2014				
	6433026*PED	Dec 10, 2014				
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N021176 001	5607669	Jun 10, 2014	U-323	I-608	Oct 02, 2012	
	5607669*PED	Dec 10, 2014		PED	Apr 02, 2013	
	5679717	Apr 29, 2014	U-323	PED	Jul 18, 2011	
	5679717*PED	Oct 29, 2014				
	5693675	Dec 02, 2014	DS			
	5693675*PED	Jun 02, 2015				
	5917007	Apr 29, 2014	DS	U-323		
	5917007*PED	Oct 29, 2014				
	5919832	Apr 29, 2014	DS			
	5919832*PED	Oct 29, 2014				
	6066678	Apr 29, 2014	DS	U-323		
	6066678*PED	Oct 29, 2014				
	6433026	Apr 29, 2014	DS			
	6433026*PED	Oct 29, 2014				
	6784254	Apr 29, 2014	DS	DP		
	6784254*PED	Oct 29, 2014				
	7101960	Apr 29, 2014	DS	DP	U-757	
	7101960*PED	Oct 29, 2014				
	7229613	Apr 17, 2022		U-851		
	7229613*PED	Oct 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
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>							
N022362 001	5607669	Jun	10, 2014	U-757		I-608	Oct 02, 2012
	5607669*PED	Dec	10, 2014			PED	Apr 02, 2013
	5679717	Apr	29, 2014	U-757		PED	Jul 18, 2011
	5679717*PED	Oct	29, 2014				
	5693675	Dec	02, 2014	DS			
	5693675*PED	Jun	02, 2015				
	5917007	Apr	29, 2014	DS	U-757		
	5917007*PED	Oct	29, 2014				
	5919832	Apr	29, 2014	DS			
	5919832*PED	Oct	29, 2014				
	6066678	Apr	29, 2014	DS	U-757		
	6066678*PED	Oct	29, 2014				
	6433026	Apr	29, 2014	DS			
	6433026*PED	Oct	29, 2014				
	6784254	Apr	29, 2014	DS DP			
	6784254*PED	Oct	29, 2014				
	7101960	Apr	29, 2014	DS DP	U-757		
	7101960*PED	Oct	29, 2014				
	7229613	Apr	17, 2022		U-493		
	7229613*PED	Oct	17, 2022				
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>							
N022362 002	5607669	Jun	10, 2014	U-757		I-608	Oct 02, 2012
	5607669*PED	Dec	10, 2014			PED	Apr 02, 2013
	5679717	Apr	29, 2014	U-757		PED	Jul 18, 2011
	5679717*PED	Oct	29, 2014				
	5693675	Dec	02, 2014	DS			
	5693675*PED	Jun	02, 2015				
	5917007	Apr	29, 2014	DS	U-757		
	5917007*PED	Oct	29, 2014				
	5919832	Apr	29, 2014	DS			
	5919832*PED	Oct	29, 2014				
	6066678	Apr	29, 2014	DS	U-757		
	6066678*PED	Oct	29, 2014				
	6433026	Apr	29, 2014	DS			
	6433026*PED	Oct	29, 2014				
	6784254	Apr	29, 2014	DS DP			
	6784254*PED	Oct	29, 2014				
	7101960	Apr	29, 2014	DS DP	U-757		
	7101960*PED	Oct	29, 2014				
	7229613	Apr	17, 2022		U-493		
	7229613*PED	Oct	17, 2022				
<u>COlestipol Hydrochloride - COlestid</u>							
N020222 001	5490987	Feb	13, 2013	DP			
<u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u>							
N021697 001	5723606	Dec	15, 2019	DS DP U-868			
	5723606	Dec	15, 2019	DS DP U-698			
<u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER</u>							
N021697 002	5723606	Dec	15, 2019	DS DP U-698			
	5723606	Dec	15, 2019	DS DP U-868			
<u>CORTICOTROPIN - H.P. ACTHAR GEL</u>							
N008372 008					ODE	Oct 15, 2017	
<u>CRIZOTINIB - XALKORI</u>							
N202570 001	7230098	Mar	01, 2025	DS		NCE	Aug 26, 2016
	7825137	May	12, 2027		U-1179	ODE	Aug 26, 2018
	7858643	Oct	08, 2029	DS DP			
<u>CRIZOTINIB - XALKORI</u>							
N202570 002	7230098	Mar	01, 2025	DS		NCE	Aug 26, 2016
	7825137	May	12, 2027		U-1179	ODE	Aug 26, 2018
	7858643	Oct	08, 2029	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
<u>CYANOCOBALAMIN - NASCOBAL</u>							
N021642 001	7229636	Jun	11, 2024	DP	U-817		
	7404489	Mar	12, 2024	DP			
	7879349	Jun	01, 2024	DP	U-1152		
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>							
N021777 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		
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>							
N021777 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		
<u>CYCLOSPORINE - NEORAL</u>							
N050715 001	5342625	Aug	30, 2011	DP			
	5741512	Aug	30, 2011	DP	U-906		
	5985321	Sep	26, 2014	DP			
	6258808	Jun	25, 2012	DP			
	6262022	Jun	25, 2012	DP			
<u>CYCLOSPORINE - NEORAL</u>							
N050715 002	5342625	Aug	30, 2011	DP			
	5741512	Aug	30, 2011	DP	U-906		
	5985321	Sep	26, 2014	DP			
	6258808	Jun	25, 2012	DP			
	6262022	Jun	25, 2012	DP			
<u>CYCLOSPORINE - NEORAL</u>							
N050715 003	5342625	Aug	30, 2011	DP			
	5741512	Aug	30, 2011	DP	U-906		
	5985321	Sep	26, 2014	DP			
	6258808	Jun	25, 2012	DP			
	6262022	Jun	25, 2012	DP			
<u>CYCLOSPORINE - NEORAL</u>							
N050716 001	5342625	Aug	30, 2011	DP			
	5741512	Aug	30, 2011	DP	U-906		
	5985321	Sep	26, 2014	DP			
	6258808	Jun	25, 2012	DP			
	6262022	Jun	25, 2012	DP			
<u>CYCLOSPORINE - RESTASIS</u>							
N050790 001	5474979	May	17, 2014	DP			
<u>CYTARABINE - DEPOCYT</u>							
N021041 001	5455044	May	14, 2013		U-806		
	5723147	Mar	03, 2015	DP	U-806		
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>							
N022512 001	6087380	Feb	18, 2018	DS	DP	U-1089	
	7866474	Aug	31, 2027		DP		
	7932273	Sep	07, 2025	DS	DP		
<u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u>							
N022512 002	6087380	Feb	18, 2018	DS	DP	U-1089	
	7866474	Aug	31, 2027		DP		
	7932273	Sep	07, 2025	DS	DP		
<u>DALFAMPRIDINE - AMPYRA</u>							
N022250 001	5370879	Dec	06, 2011	DP			
	5540938	Jul	30, 2013		U-1030		
	8007826	May	26, 2027		U-1030		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DAPSONE - ACZONE</u>							
N021794 001	5863560	Sep	11, 2016	DP			
	6060085	Sep	11, 2016		U-124		
	6620435	Sep	11, 2016	DP			
<u>DAPTOMYCIN - CUBICIN</u>							
N021572 001	6468967	Sep	24, 2019		U-282		
	6852689	Sep	24, 2019		U-282		
	8058238	Nov	28, 2020	DS DP			
<u>DAPTOMYCIN - CUBICIN</u>							
N021572 002	6468967	Sep	24, 2019		U-282		
	6852689	Sep	24, 2019		U-282		
	8003673	Sep	04, 2028		U-1180		
	8058238	Nov	28, 2020	DS DP			
	RE39071	Jun	15, 2016	DS DP	U-728		
<u>DARIFENACIN HYDROBROMIDE - ENABLEX</u>							
N021513 001	5096890	Mar	13, 2015	DS	DP U-631		
	6106864	Aug	21, 2016	DP	U-630		
<u>DARIFENACIN HYDROBROMIDE - ENABLEX</u>							
N021513 002	5096890	Mar	13, 2015	DS	DP U-631		
	6106864	Aug	21, 2016	DP	U-630		
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>							
N021976 001	5843946	Dec	01, 2015	DP	U-744	D-129	Dec 13, 2013
	5843946	Dec	01, 2015	DP	U-903	D-119	Dec 18, 2011
	5843946	Dec	01, 2015	DP	U-935	I-578	Oct 21, 2011
	5843946*PED	Jun	01, 2016			NPP	Dec 16, 2014
	6037157	Jun	26, 2016		U-935	PED	Jun 16, 2015
	6037157*PED	Dec	26, 2016			PED	Dec 23, 2011
	6248775	Aug	13, 2014	DS		PED	Apr 21, 2012
	6248775*PED	Feb	13, 2015			PED	Jun 13, 2014
	6335460	Aug	25, 2012	DS DP	U-935	PED	Jun 18, 2012
	6335460	Aug	25, 2012	DS DP	U-903		
	6335460	Aug	25, 2012	DS DP	U-744		
	6335460*PED	Feb	25, 2013				
	6703403	Jun	26, 2016		U-935		
	6703403*PED	Dec	26, 2016				
	7470506	Jun	23, 2019		U-935		
	7470506*PED	Dec	23, 2019				
	7700645	Dec	26, 2026	DS DP			
	7700645*PED	Jun	26, 2027				
	RE42889	Oct	19, 2016		DP		
	RE42889*PED	Apr	19, 2017				
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>							
N021976 002	5843946	Dec	01, 2015	DP	U-935	D-129	Dec 13, 2013
	5843946	Dec	01, 2015	DP	U-744	D-119	Dec 18, 2011
	5843946	Dec	01, 2015	DP	U-903	I-578	Oct 21, 2011
	5843946*PED	Jun	01, 2016			NPP	Dec 16, 2014
	6037157	Jun	26, 2016		U-935	PED	Jun 16, 2015
	6037157*PED	Dec	26, 2016			PED	Dec 23, 2011
	6248775	Aug	13, 2014	DS		PED	Apr 21, 2012
	6248775*PED	Feb	13, 2015			PED	Jun 13, 2014
	6335460	Aug	25, 2012	DS DP	U-744	PED	Jun 18, 2012
	6335460	Aug	25, 2012	DS DP	U-903		
	6335460	Aug	25, 2012	DS DP	U-935		
	6335460*PED	Feb	25, 2013				
	6703403	Jun	26, 2016		U-935		
	6703403*PED	Dec	26, 2016				
	7470506	Jun	23, 2019		U-935		
	7470506*PED	Dec	23, 2019				
	7700645	Dec	26, 2026	DS DP			
	7700645*PED	Jun	26, 2027				
	RE42889	Oct	19, 2016		DP		
	RE42889*PED	Apr	19, 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
DARUNAVIR ETHANOLATE - PREZISTA								
N021976 003	5843946	Dec	01, 2015	DP	U-744		D-129	Dec 13, 2013
	5843946	Dec	01, 2015	DP	U-903		D-119	Dec 18, 2011
	5843946	Dec	01, 2015	DP	U-935		D-118	Oct 21, 2011
	5843946*PED	Jun	01, 2016				I-578	Oct 21, 2011
	6037157	Jun	26, 2016		U-935		NPP	Dec 16, 2014
	6037157*PED	Dec	26, 2016				PED	Jun 16, 2015
	6248775	Aug	13, 2014	DS			PED	Dec 23, 2011
	6248775*PED	Feb	13, 2015				PED	Apr 21, 2012
	6335460	Aug	25, 2012	DS	DP U-935		PED	Jun 18, 2012
	6335460	Aug	25, 2012	DS	DP U-903		PED	Jun 13, 2014
	6335460*PED	Feb	25, 2013				PED	Apr 21, 2012
	6703403	Jun	26, 2016		U-935		PED	
	6703403*PED	Dec	26, 2016				PED	
	7470506	Jun	23, 2019		U-935		PED	
	7470506*PED	Dec	23, 2019				PED	
	7700645	Dec	26, 2026	DS	DP			
	7700645*PED	Jun	26, 2027					
	RE42889	Oct	19, 2016		DP			
	RE42889*PED	Apr	19, 2017					
DARUNAVIR ETHANOLATE - PREZISTA								
N021976 004	5843946	Dec	01, 2015	DP	U-935		D-129	Dec 13, 2013
	5843946*PED	Jun	01, 2016				D-119	Dec 18, 2011
	6037157	Jun	26, 2016		U-935		NPP	Dec 16, 2014
	6037157*PED	Dec	26, 2016				NS	Dec 18, 2011
	6248775	Aug	13, 2014	DS			PED	Jun 16, 2015
	6248775*PED	Feb	13, 2015				PED	Dec 23, 2011
	6335460	Aug	25, 2012	DS	DP U-935		PED	Jun 18, 2012
	6335460*PED	Feb	25, 2013				PED	Jun 13, 2014
	6703403	Jun	26, 2016		U-935		PED	Jun 18, 2012
	6703403*PED	Dec	26, 2016				PED	
	7470506	Jun	23, 2019		U-935		PED	
	7470506*PED	Dec	23, 2019				PED	
	7700645	Dec	26, 2026	DS	DP			
	7700645*PED	Jun	26, 2027					
	RE42889	Oct	19, 2016		DP			
	RE42889*PED	Apr	19, 2017					
DARUNAVIR ETHANOLATE - PREZISTA								
N021976 005	5843946	Dec	01, 2015	DP	U-935		D-129	Dec 13, 2013
	5843946*PED	Jun	01, 2016				D-119	Dec 18, 2011
	6037157	Jun	26, 2016		U-935		NPP	Dec 16, 2014
	6037157*PED	Dec	26, 2016				NS	Dec 18, 2011
	6248775	Aug	13, 2014	DS			PED	Jun 16, 2015
	6248775*PED	Feb	13, 2015				PED	Dec 23, 2011
	6335460	Aug	25, 2012	DS	DP U-935		PED	Jun 18, 2012
	6335460*PED	Feb	25, 2013				PED	Jun 13, 2014
	6703403	Jun	26, 2016		U-935		PED	Jun 18, 2012
	6703403*PED	Dec	26, 2016				PED	
	7470506	Jun	23, 2019		U-935		PED	
	7470506*PED	Dec	23, 2019				PED	
	7700645	Dec	26, 2026	DS	DP			
	7700645*PED	Jun	26, 2027					
	RE42889	Oct	19, 2016		DP			
	RE42889*PED	Apr	19, 2017					
DARUNAVIR ETHANOLATE - PREZISTA								
N202895 001							NDF	Dec 16, 2014
							PED	Jun 16, 2015
DASATINIB - SPRYCEL								
N021986 001	6596746	Jun	28, 2020	DS	DP U-780		D-120	May 21, 2012
	6596746	Jun	28, 2020	DS	DP U-748		M-94	Oct 28, 2013
	7125875	Apr	13, 2020		U-780		ODE	Jun 28, 2013
	7125875	Apr	13, 2020		U-779		ODE	Jun 28, 2013
	7153856	Apr	28, 2020		U-780			
	7491725	Mar	28, 2026	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
DASATINIB - SPRYCEL										
N021986 002	6596746	Jun	28,	2020	DS	DP	U-780		D-120	May 21, 2012
	6596746	Jun	28,	2020	DS	DP	U-748		M-94	Oct 28, 2013
	7125875	Apr	13,	2020			U-780		ODE	Jun 28, 2013
	7125875	Apr	13,	2020			U-779		ODE	Jun 28, 2013
	7153856	Apr	28,	2020			U-780		ODE	Jun 28, 2013
	7491725	Mar	28,	2026	DS	DP				
DASATINIB - SPRYCEL										
N021986 003	6596746	Jun	28,	2020	DS	DP	U-780		D-120	May 21, 2012
	6596746	Jun	28,	2020	DS	DP	U-748		M-94	Oct 28, 2013
	7125875	Apr	13,	2020			U-780		ODE	Jun 28, 2013
	7125875	Apr	13,	2020			U-779		ODE	Jun 28, 2013
	7153856	Apr	28,	2020			U-780		ODE	Jun 28, 2013
	7491725	Mar	28,	2026	DS	DP				
DASATINIB - SPRYCEL										
N021986 004	6596746	Jun	28,	2020	DS	DP	U-780		D-120	May 21, 2012
	6596746	Jun	28,	2020	DS	DP	U-748		M-94	Oct 28, 2013
	7125875	Apr	13,	2020			U-779		ODE	Jun 28, 2013
	7125875	Apr	13,	2020			U-780		ODE	Jun 28, 2013
	7153856	Apr	28,	2020			U-780		ODE	Jun 28, 2013
	7491725	Mar	28,	2026	DS	DP				
DASATINIB - SPRYCEL										
N021986 005	6596746	Jun	28,	2020	DS	DP	U-748		D-120	May 21, 2012
	6596746	Jun	28,	2020	DS	DP	U-780		M-94	Oct 28, 2013
	7125875	Apr	13,	2020			U-780		ODE	Jun 28, 2013
	7125875	Apr	13,	2020			U-779		ODE	Jun 28, 2013
	7153856	Apr	28,	2020			U-780		ODE	Jun 28, 2013
	7491725	Mar	28,	2026	DS	DP				
DASATINIB - SPRYCEL										
N021986 006	6596746	Jun	28,	2020	DS	DP	U-748		D-120	May 21, 2012
	6596746	Jun	28,	2020	DS	DP	U-780		M-94	Oct 28, 2013
	7125875	Apr	13,	2020			U-780		ODE	Jun 28, 2013
	7125875	Apr	13,	2020			U-779		ODE	Jun 28, 2013
	7153856	Apr	28,	2020			U-780		ODE	Jun 28, 2013
	7491725	Mar	28,	2026	DS	DP				
DECITABINE - DACOGEN										
N021790 001									D-123	Mar 11, 2013
									ODE	May 02, 2013
DEFERASIROX - EXJADE										
N021882 001	6465504	Apr	05,	2019	DS	DP			ODE	Nov 02, 2012
	6596750	Jun	24,	2017	DS		U-735			
DEFERASIROX - EXJADE										
N021882 002	6465504	Apr	05,	2019	DS	DP			ODE	Nov 02, 2012
	6596750	Jun	24,	2017	DS		U-735			
DEFERASIROX - EXJADE										
N021882 003	6465504	Apr	05,	2019	DS	DP			ODE	Nov 02, 2012
	6596750	Jun	24,	2017	DS		U-735			
DEFERIPRONE - FERRIPROX										
N021825 001									NCE	Oct 14, 2016
									ODE	Oct 14, 2018
DEGARELIX ACETATE - FIRMAGON										
N022201 001	5925730	Apr	11,	2017	DS	DP	U-943		NCE	Dec 24, 2013
DEGARELIX ACETATE - FIRMAGON										
N022201 002	5925730	Apr	11,	2017	DS	DP	U-943		NCE	Dec 24, 2013
DELAVIRDINE MESYLATE - RESCRIPTOR										
N020705 001	5563142	Oct	08,	2013						

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DELAVIRDINE MESYLATE - RESRIPTOR</u>						
N020705 002	5563142	Oct 08, 2013				
	6177101	Jun 07, 2019				
<u>DESFLURANE - SUPRANE</u>						
N020118 001	5617906	Apr 08, 2014	DP			
	5617906*PED	Oct 08, 2014				
<u>DESIRUDIN RECOMBINANT - IPRIVASK</u>						
N021271 001	5733874	Mar 31, 2015				
<u>DESLORATADINE - CLARINEX</u>						
N021165 001	6100274	Jul 07, 2019				
	6100274*PED	Jan 07, 2020				
	7211582	Dec 30, 2014	U-809			
	7211582*PED	Jun 30, 2015				
	7214683	Dec 30, 2014	DP			
	7214683*PED	Jun 30, 2015				
	7214684	Dec 30, 2014	U-138			
	7214684*PED	Jun 30, 2015				
	7405223	Jul 07, 2019	U-886			
	7405223*PED	Jan 07, 2020				
<u>DESLORATADINE - CLARINEX</u>						
N021300 001	6514520	Jun 01, 2018	DP			
	6514520*PED	Dec 01, 2018				
	7211582	Dec 30, 2014	U-809			
	7211582*PED	Jun 30, 2015				
	7214683	Dec 30, 2014	DP			
	7214683*PED	Jun 30, 2015				
	7214684	Dec 30, 2014	U-138			
	7214684*PED	Jun 30, 2015				
<u>DESLORATADINE - CLARINEX</u>						
N021312 001	5607697	Jun 07, 2015	DP			
	5607697*PED	Dec 07, 2015				
	6100274	Jul 07, 2019	DP			
	6100274*PED	Jan 07, 2020				
	7211582	Dec 30, 2014	U-809			
	7211582*PED	Jun 30, 2015				
	7214683	Dec 30, 2014	DP			
	7214683*PED	Jun 30, 2015				
	7214684	Dec 30, 2014	U-138			
	7214684*PED	Jun 30, 2015				
	7618649	Dec 19, 2020	DP U-1017			
	7618649*PED	Jun 19, 2021				
<u>DESLORATADINE - CLARINEX</u>						
N021312 002	5607697	Jun 07, 2015	DP			
	5607697*PED	Dec 07, 2015				
	6100274	Jul 07, 2019	DP			
	6100274*PED	Jan 07, 2020				
	7211582	Dec 30, 2014	U-809			
	7211582*PED	Jun 30, 2015				
	7214683	Dec 30, 2014	DP			
	7214683*PED	Jun 30, 2015				
	7214684	Dec 30, 2014	U-138			
	7214684*PED	Jun 30, 2015				
	7618649	Dec 19, 2020	DP U-1017			
	7618649*PED	Jun 19, 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
DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 24 HOUR						
N021605 001	6100274	Jul 07, 2019	DP			
	6100274*PED	Jan 07, 2020				
	6979463	Mar 28, 2022	DP			
	7214683	Dec 30, 2014	DP			
	7214683*PED	Jun 30, 2015				
	7214684	Dec 30, 2014	U-138			
	7214684*PED	Jun 30, 2015				
	7618649	Dec 19, 2020	DP U-1017			
	7618649*PED	Jun 19, 2021				
	7820199	Mar 28, 2022	DP			
	7820199*PED	Sep 28, 2022				
DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR						
N021313 001	6100274	Jul 07, 2019	DP			
	6100274*PED	Jan 07, 2020				
	6709676	Feb 18, 2021	DP U-707			
	7214683	Dec 30, 2014	DP			
	7214683*PED	Jun 30, 2015				
	7214684	Dec 30, 2014	U-138			
	7214684*PED	Jun 30, 2015				
	7618649	Dec 19, 2020	DP U-1017			
	7618649*PED	Jun 19, 2021				
DESMOPRESSIN ACETATE - DDAVP						
N017922 001	5500413	Jun 29, 2013				
	5674850	Dec 23, 2013				
	5763407	Jun 29, 2013				
DESMOPRESSIN ACETATE - DDAVP						
N017922 002	5500413	Jun 29, 2013				
	5674850	Dec 23, 2013				
	5763407	Jun 29, 2013				
DESMOPRESSIN ACETATE - DDAVP						
N018938 001	5500413	Jun 29, 2013				
	5763407	Jun 29, 2013				
DESMOPRESSIN ACETATE - DDAVP						
N018938 002	5500413	Jun 29, 2013				
	5763407	Jun 29, 2013				
DESMOPRESSIN ACETATE - DDAVP						
N019955 001	5500413	Jun 29, 2013				
	5674850	Dec 23, 2013				
	5763407	Jun 29, 2013				
	7022340	Apr 30, 2023	DP			
DESMOPRESSIN ACETATE - DDAVP						
N019955 002	5500413	Jun 29, 2013				
	5674850	Dec 23, 2013				
	5763407	Jun 29, 2013				
	7022340	Apr 30, 2023	DP			
DESMOPRESSIN ACETATE - DDAVP (NEEDS NO REFRIGERATION)						
N017922 003	5482931	Jun 29, 2013				
	5500413	Jun 29, 2013				
	5674850	Dec 23, 2013				
	5763407	Jun 29, 2013				
DESMOPRESSIN ACETATE - DESMOPRESSIN ACETATE						
N021795 001	7022340	Apr 30, 2023	DP			
DESMOPRESSIN ACETATE - DESMOPRESSIN ACETATE						
N021795 002	7022340	Apr 30, 2023	DP			
DESONIDE - DESONATE						
N021844 001	6387383	Aug 03, 2020	DS DP U-783			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DESONIDE - VERDESO</u>								
N021978 001	6730288	Sep	08, 2019	DP				
	7029659	Sep	08, 2019	DP				
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>								
N021992 001	6673838	Feb	11, 2022	DS	DP	U-860	NCE	Mar 01, 2013
<u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u>								
N021992 002	6673838	Feb	11, 2022	DS	DP	U-860	NCE	Mar 01, 2013
<u>DEXAMETHASONE - OZURDEX</u>								
N022315 001	6726918	Oct	20, 2020	DP	U-1205		NDF	Jun 17, 2012
	6726918	Oct	20, 2020	DP	U-1204		ODE	Sep 24, 2017
	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	DP	U-1205			
	8063031	Oct	20, 2020	DP				
<u>DEXAMETHASONE; TOBRAMYCIN - TOBRADEX ST</u>								
N050818 001	7795316	Aug	03, 2028	DP	U-1082			
<u>DEXLANSOPRAZOLE - DEXILANT</u>								
N022287 001	6462058	Jun	15, 2020	DS	DP	U-949	NP	Jan 30, 2012
	6462058	Jun	15, 2020	DS	DP	U-951	PED	Jul 30, 2012
	6462058	Jun	15, 2020	DS	DP	U-950		
	6462058*PED	Dec	15, 2020					
	6664276	Jun	15, 2020	DS	DP	U-951		
	6664276	Jun	15, 2020	DS	DP	U-949		
	6664276	Jun	15, 2020	DS	DP	U-950		
	6664276*PED	Dec	15, 2020					
	6939971	Jun	15, 2020		U-949			
	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					
	7790755	Aug	02, 2026		DP			
	7790755*PED	Feb	02, 2027					
<u>DEXLANSOPRAZOLE - DEXILANT</u>								
N022287 002	6462058	Jun	15, 2020	DS	DP	U-951	NP	Jan 30, 2012
	6462058	Jun	15, 2020	DS	DP	U-950	PED	Jul 30, 2012
	6462058	Jun	15, 2020	DS	DP	U-949		
	6462058*PED	Dec	15, 2020					
	6664276	Jun	15, 2020	DS	DP	U-951		
	6664276	Jun	15, 2020	DS	DP	U-950		
	6664276	Jun	15, 2020	DS	DP	U-949		
	6664276*PED	Dec	15, 2020					
	6939971	Jun	15, 2020		U-949			
	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					
	7790755	Aug	02, 2026		DP			
	7790755*PED	Feb	02, 2027					
<u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u>								
N021038 001	4910214	Jul	15, 2013	DS	DP	U-421	I-577	Oct 17, 2011
	5344840	Sep	06, 2011			U-912		
	6716867	Mar	31, 2019			U-572		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN</u>								
N021278 001	5908850	Dec	04, 2015		U-422			
	6355656	Dec	04, 2015					
	6528530	Dec	04, 2015	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
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN						
N021278 002	5908850	Dec 04, 2015		U-422		
	6355656	Dec 04, 2015				
	6528530	Dec 04, 2015	DS DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN						
N021278 003	5908850	Dec 04, 2015		U-422		
	6355656	Dec 04, 2015				
	6528530	Dec 04, 2015	DS DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
N021802 001	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	U-678		M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP U-676			
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP U-677			
	6730325	Nov 01, 2019	DP U-676			
	7431944	Dec 04, 2015	DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
N021802 002	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	U-678		M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP U-676			
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP U-677			
	6730325	Nov 01, 2019	DP U-676			
	7431944	Dec 04, 2015	DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
N021802 003	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	U-678		M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP U-676			
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP U-677			
	6730325	Nov 01, 2019	DP U-676			
	7431944	Dec 04, 2015	DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
N021802 004	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	U-678		M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP U-676			
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP U-677			
	6730325	Nov 01, 2019	DP U-676			
	7431944	Dec 04, 2015	DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
N021802 005	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	U-678		M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP U-676			
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP U-677			
	6730325	Nov 01, 2019	DP U-676			
	7431944	Dec 04, 2015	DP			
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
N021802 006	5837284	Dec 04, 2015	DP U-677		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	U-678		M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP U-676			
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP U-677			
	6730325	Nov 01, 2019	DP U-676			
	7431944	Dec 04, 2015	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>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>						
N021802 007	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	DP	U-678	M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP	U-676		
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP	U-677		
	6730325	Nov 01, 2019	DP	U-676		
	7431944	Dec 04, 2015	DP			
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>						
N021802 008	5837284	Dec 04, 2015	DP		D-121	Oct 23, 2012
	5908850	Dec 04, 2015	DP	U-678	M-80	Oct 17, 2011
	6228398	Nov 01, 2019	DP	U-676		
	6355656	Dec 04, 2015	DP			
	6528530	Dec 04, 2015	DP			
	6635284	Dec 04, 2015	DP	U-677		
	6730325	Nov 01, 2019	DP	U-676		
	7431944	Dec 04, 2015	DP			
<u>DEXRAZOXANE HYDROCHLORIDE - TOTECT</u>						
N022025 001	6727253	Mar 13, 2020	DP	U-829	ODE	Sep 06, 2014
<u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u>						
N021620 001	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP	U-685		
	7838032	Apr 28, 2020	DP			
<u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u>						
N021620 002	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP	U-685		
	7838032	Apr 28, 2020	DP			
<u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u>						
N021879 001	5206248	Mar 27, 2012	DP	U-1093	NC	Oct 29, 2013
	7659282	Aug 13, 2026	DP	U-1093		
	RE38115	Jan 26, 2016	DP			
<u>DEXTROMETHORPHAN POLISTIREX - DELSYM</u>						
N018658 001	5980882	Apr 16, 2017	DP			
<u>DIAZEPAM - DIASTAT</u>						
N020648 001	5462740	Sep 17, 2013	DP			
<u>DIAZEPAM - DIASTAT</u>						
N020648 002	5462740	Sep 17, 2013	DP			
<u>DIAZEPAM - DIASTAT</u>						
N020648 003	5462740	Sep 17, 2013	DP			
<u>DIAZEPAM - DIASTAT</u>						
N020648 004	5462740	Sep 17, 2013	DP			
<u>DIAZEPAM - DIASTAT</u>						
N020648 005	5462740	Sep 17, 2013	DP			
<u>DIAZEPAM - DIASTAT ACUDIAL</u>						
N020648 006	5462740	Sep 17, 2013	DP			
<u>DIAZEPAM - DIASTAT ACUDIAL</u>						
N020648 007	5462740	Sep 17, 2013	DP			
<u>DICLOFENAC EPOLAMINE - FLECTOR</u>						
N021234 001	5607690	Apr 13, 2014	DP			
<u>DICLOFENAC POTASSIUM - CAMBIA</u>						
N022165 001	6974595	May 15, 2017	DS	U-436	NDF	Jun 17, 2012
	7482377	May 15, 2017	DP	U-436		
	7759394	Jun 16, 2026	DS	U-436		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
DICLOFENAC POTASSIUM - ZIPSOR						
N022202 001	6365180	Jul 15, 2019	DP U-980		NDF	Jun 16, 2012
	7662858	Feb 24, 2029	U-1035			
	7884095	Feb 24, 2029	U-1111			
	7939518	Feb 24, 2029	U-980			
DICLOFENAC SODIUM - DICLOFENAC SODIUM						
N020809 001	5603929	Nov 16, 2014	U-239			
	5653972	Nov 16, 2014	U-239			
DICLOFENAC SODIUM - PENNSAID						
N020947 001					NDF	Nov 04, 2012
DICLOFENAC SODIUM - SOLARAZE						
N021005 001	5639738	Jun 17, 2014	U-402			
	5792753	Aug 11, 2015				
	5852002	Jun 17, 2014	U-402			
	5914322	Aug 11, 2015				
	5929048	Jun 17, 2014	U-402			
	5985850	Aug 11, 2015	DP			
DICLOFENAC SODIUM; MISOPROSTOL - ARTHROTEC						
N020607 001	5601843	Feb 11, 2014				
DICLOFENAC SODIUM; MISOPROSTOL - ARTHROTEC						
N020607 002	5601843	Feb 11, 2014				
DIDANOSINE - VIDEX						
N020154 002	5880106	Jul 22, 2011				
	5880106*PED	Jan 22, 2012				
DIDANOSINE - VIDEX						
N020154 003	5880106	Jul 22, 2011				
	5880106*PED	Jan 22, 2012				
DIDANOSINE - VIDEX						
N020154 004	5880106	Jul 22, 2011				
	5880106*PED	Jan 22, 2012				
DIDANOSINE - VIDEX						
N020154 005	5880106	Jul 22, 2011				
	5880106*PED	Jan 22, 2012				
DIDANOSINE - VIDEX						
N020154 006	5880106	Jul 22, 2011				
	5880106*PED	Jan 22, 2012				
DIENOGEST; ESTRADIOL VALERATE - NATAZIA						
N022252 001	6133251	Oct 25, 2016	DP U-828	Y	NP	May 06, 2013
	6133251	Oct 25, 2016	DP U-112	Y		
	6133251	Oct 25, 2016	DP U-1	Y		
	6884793	Oct 25, 2016	DP	Y		
	8071577	May 13, 2026	DP U-1			
DIFLUPREDNATE - DUREZOL						
N022212 001	6114319	May 18, 2019	DP		NCE	Jun 23, 2013
DILTIAZEM HYDROCHLORIDE - CARDIZEM CD						
N020062 001	5364620	Nov 14, 2011	U-3			
	5439689	Aug 08, 2012	U-107			
DILTIAZEM HYDROCHLORIDE - CARDIZEM CD						
N020062 002	5364620	Nov 14, 2011	U-3			
	5439689	Aug 08, 2012	U-107			
DILTIAZEM HYDROCHLORIDE - CARDIZEM CD						
N020062 003	5364620	Nov 14, 2011	U-3			
	5439689	Aug 08, 2012	U-107			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DILTIAZEM HYDROCHLORIDE - CARDIZEM CD</u>						
N020062 004	5364620	Nov 14, 2011		U-3		
	5439689	Aug 08, 2012		U-107		
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM CD</u>						
N020062 005	5439689	Aug 08, 2012		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N021392 001	5529791	Jun 25, 2013		DP		
	6923984	Feb 25, 2021				
	7108866	Dec 17, 2019		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N021392 002	5529791	Jun 25, 2013		DP		
	6923984	Feb 25, 2021				
	7108866	Dec 17, 2019		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N021392 003	5529791	Jun 25, 2013		DP		
	6923984	Feb 25, 2021				
	7108866	Dec 17, 2019		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N021392 004	5529791	Jun 25, 2013		DP		
	6923984	Feb 25, 2021				
	7108866	Dec 17, 2019		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N021392 005	5529791	Jun 25, 2013		DP		
	6923984	Feb 25, 2021				
	7108866	Dec 17, 2019		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>						
N021392 006	5529791	Jun 25, 2013		DP		
	6923984	Feb 25, 2021				
	7108866	Dec 17, 2019		DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - DILACOR XR</u>						
N020092 001	5422123	Jun 06, 2012				
<u>DILTIAZEM HYDROCHLORIDE - DILACOR XR</u>						
N020092 002	5422123	Jun 06, 2012				
<u>DILTIAZEM HYDROCHLORIDE - DILACOR XR</u>						
N020092 003	5422123	Jun 06, 2012				
<u>DILTIAZEM HYDROCHLORIDE - DILTIAZEM HYDROCHLORIDE</u>						
N020939 001	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - DILTIAZEM HYDROCHLORIDE</u>						
N020939 002	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - DILTIAZEM HYDROCHLORIDE</u>						
N020939 003	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - DILTIAZEM HYDROCHLORIDE</u>						
N020939 004	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - TIAZAC</u>						
N020401 001	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - TIAZAC</u>						
N020401 002	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - TIAZAC</u>						
N020401 003	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - TIAZAC</u>						
N020401 004	5529791	Jun 25, 2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DILTIAZEM HYDROCHLORIDE - TIAZAC</u>						
N020401 005	5529791	Jun 25, 2013				
<u>DILTIAZEM HYDROCHLORIDE - TIAZAC</u>						
N020401 006	5529791	Jun 25, 2013				
<u>DIMYRISTOYL LECITHIN; PERFLEXANE - IMAGENT</u>						
N021191 001	5605673	Feb 25, 2014				
	5626833	May 16, 2014		U-458		
	5639443	Jun 17, 2014				
	5695741	Dec 09, 2014		U-458		
	5720938	Feb 24, 2015				
	5798091	Aug 25, 2015		U-458		
	6280704	Jul 30, 2013				
	6280705	Jul 30, 2013				
	6287539	Jul 30, 2013				
<u>DINOPROSTONE - CERVIDIL</u>						
N020411 001	5269321	Jul 14, 2012	DP	U-110		
<u>DIVALPROEX SODIUM - DEPAKOTE</u>						
N019680 001					PED	Sep 24, 2011
<u>DIVALPROEX SODIUM - DEPAKOTE ER</u>						
N021168 001	6419953	Dec 18, 2018			PED	Sep 24, 2011
	6419953*PED	Jun 18, 2019				
	6511678	Dec 18, 2018				
	6511678*PED	Jun 18, 2019				
	6528090	Dec 18, 2018	DP			
	6528090*PED	Jun 18, 2019				
	6528091	Dec 18, 2018		U-106		
	6528091*PED	Jun 18, 2019				
	6713086	Dec 18, 2018	DP	U-579		
	6713086*PED	Jun 18, 2019				
	6720004	Dec 18, 2018	DP			
	6720004*PED	Jun 18, 2019				
<u>DIVALPROEX SODIUM - DEPAKOTE ER</u>						
N021168 002	6511678	Dec 18, 2018			PED	Sep 24, 2011
	6511678*PED	Jun 18, 2019				
	6528090	Dec 18, 2018	DP			
	6528090*PED	Jun 18, 2019				
	6713086	Dec 18, 2018	DP	U-579		
	6713086*PED	Jun 18, 2019				
	6720004	Dec 18, 2018	DP			
	6720004*PED	Jun 18, 2019				
<u>DOCETAXEL - TAXOTERE</u>						
N020449 001	5438072	Nov 22, 2013	DP		M-61	May 13, 2013
	5438072*PED	May 22, 2014			PED	Nov 13, 2013
	5698582	Jul 03, 2012	DP			
	5698582*PED	Jan 03, 2013				
	5714512	Jul 03, 2012	DP			
	5714512*PED	Jan 03, 2013				
	5750561	Jul 03, 2012	DP			
	5750561*PED	Jan 03, 2013				
<u>DOCETAXEL - TAXOTERE</u>						
N020449 003	5698582	Jul 03, 2012	DP		M-61	May 13, 2013
	5698582*PED	Jan 03, 2013			PED	Nov 13, 2013
	5714512	Jul 03, 2012	DP			
	5714512*PED	Jan 03, 2013				
	5750561	Jul 03, 2012	DP			
	5750561*PED	Jan 03, 2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DOCETAXEL - TAXOTERE</u>							
N020449 004	5698582	Jul	03, 2012	DP		M-61	May 13, 2013
	5698582*PED	Jan	03, 2013			PED	Nov 13, 2013
	5714512	Jul	03, 2012	DP			
	5714512*PED	Jan	03, 2013				
	5750561	Jul	03, 2012	DP			
	5750561*PED	Jan	03, 2013				
<u>DOCOSANOL - ABREVA</u>							
N020941 001	4874794	Apr	28, 2014	U-815			
	5534554	Dec	13, 2013	DP U-815			
<u>DOFETILIDE - TIKOSYN</u>							
N020931 001	4959366	Sep	25, 2012	DS DP	U-652		
	6124363	Oct	09, 2018				
<u>DOFETILIDE - TIKOSYN</u>							
N020931 002	4959366	Sep	25, 2012	DS DP	U-652		
	6124363	Oct	09, 2018				
<u>DOFETILIDE - TIKOSYN</u>							
N020931 003	4959366	Sep	25, 2012	DS DP	U-652		
	6124363	Oct	09, 2018				
<u>DOLASETRON MESYLATE - ANZEMET</u>							
N020623 001	4906755	Jul	02, 2011				
<u>DOLASETRON MESYLATE - ANZEMET</u>							
N020623 002	4906755	Jul	02, 2011				
<u>DOLASETRON MESYLATE - ANZEMET</u>							
N020624 001	4906755	Jul	02, 2011				
<u>DOLASETRON MESYLATE - ANZEMET</u>							
N020624 002	4906755	Jul	02, 2011				
<u>DOLASETRON MESYLATE - ANZEMET</u>							
N020624 003	4906755	Jul	02, 2011				
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>							
N020690 001	5985864	Dec	30, 2016				
	6140321	Dec	30, 2016				
	6245911	Dec	01, 2018				
	6372760	Mar	31, 2019		Y		
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>							
N020690 002	5985864	Dec	30, 2016				
	6140321	Dec	30, 2016				
	6245911	Dec	01, 2018				
	6372760	Mar	31, 2019		Y		
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>							
N022568 001						NP	Jul 23, 2013
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u>							
N021720 001	7727548	Jun	23, 2022	DP	U-1062		
	7727552	Mar	26, 2018	DP			
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u>							
N021720 002	7727548	Jun	23, 2022	DP	U-1062		
	7727552	Mar	26, 2018	DP			
<u>DORIPENEM - DORIBAX</u>							
N022106 001	5317016	Jun	05, 2015	DS DP	U-282	NCE	Oct 12, 2012
<u>DORIPENEM - DORIBAX</u>							
N022106 002	5317016	Jun	05, 2015	DS DP	U-282	NCE	Oct 12, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>DOXEPEPIN HYDROCHLORIDE - SILENOR</u>							
N022036 001	5502047	Mar	26, 2013		U-620		NP
	5585115	Jan	09, 2015	DP			
	5725884	Jan	09, 2015	DP			
	5866166	Jan	09, 2015	DP			
	5948438	Jan	09, 2015	DP			
	6103219	Jan	09, 2015	DP			
	6211229	Feb	17, 2020		U-620		
	6217909	Jan	09, 2015	DP			
	7915307	Aug	24, 2027		U-620		
<u>DOXEPEPIN HYDROCHLORIDE - SILENOR</u>							
N022036 002	5502047	Mar	26, 2013		U-620		NP
	5585115	Jan	09, 2015	DP			
	5725884	Jan	09, 2015	DP			
	5866166	Jan	09, 2015	DP			
	5948438	Jan	09, 2015	DP			
	6103219	Jan	09, 2015	DP			
	6211229	Feb	17, 2020		U-620		
	6217909	Jan	09, 2015	DP			
	7915307	Aug	24, 2027		U-620		
<u>DOXERCALCIFEROL - HECTOROL</u>							
N020862 001	5602116	Feb	11, 2014		U-278		
	5602116	Feb	11, 2014		U-987		
	6903083	Jul	18, 2021	DS DP		Y	
<u>DOXERCALCIFEROL - HECTOROL</u>							
N020862 002	5602116	Feb	11, 2014		U-278		
	5602116	Feb	11, 2014		U-987		
	6903083	Jul	18, 2021	DS DP		Y	
<u>DOXERCALCIFEROL - HECTOROL</u>							
N020862 003	5602116	Feb	11, 2014		U-987		
<u>DOXERCALCIFEROL - HECTOROL</u>							
N021027 001	5602116	Feb	11, 2014		U-321		
	6903083	Jul	18, 2021	DS DP		Y	
	7148211	Sep	14, 2023	DP			
<u>DOXERCALCIFEROL - HECTOROL</u>							
N021027 002	5602116	Feb	11, 2014		U-321		
	7148211	Sep	14, 2023	DP			
<u>DOXORUBICIN HYDROCHLORIDE - DOXIL</u>							
N050718 001						ODE	May 17, 2014
<u>DOXORUBICIN HYDROCHLORIDE - DOXIL</u>							
N050718 002						ODE	May 17, 2014
<u>DOXYCYCLINE - ORACEA</u>							
N050805 001	5789395	Aug	30, 2016		U-925		
	5919775	Aug	30, 2016		U-925		
	7211267	Apr	05, 2022		U-925		
	7232572	Apr	05, 2022		U-925		
	7749532	Dec	19, 2027	DP	U-1063		
<u>DOXYCYCLINE HYCLATE - DORYX</u>							
N050795 001	6958161	Dec	15, 2022	DP	U-918		
<u>DOXYCYCLINE HYCLATE - DORYX</u>							
N050795 002	6958161	Dec	15, 2022	DP	U-918		
<u>DOXYCYCLINE HYCLATE - DORYX</u>							
N050795 003	6958161	Dec	15, 2022	DP	U-918		
<u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u>							
N022425 001	5223510	Jul	26, 2012	DS	DP U-992	NCE	Jul 01, 2014
	7323493	Jun	19, 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>DROSPIRENONE; ESTRADIOL - ANGELIQ</u>						
	N021355 002	6933395	Aug 11, 2017	DS		
<u>DROSPIRENONE; ETHINYL ESTRADIOL - YASMIN</u>						
	N021098 001	5569652	Oct 29, 2013	U-1		
		6787531	Aug 31, 2020	DP		
		6933395	Aug 11, 2017	DS		
<u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u>						
	N021676 001	5569652	Oct 29, 2013	U-1		
		5798338	Jul 10, 2015	DP		
		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		
		RE37564	Jun 30, 2014	DP		
		RE37838	Jun 30, 2014	DP		
		RE38253	Jun 30, 2014	DP		
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - BEYAZ</u>						
	N022532 001	5798338	Jul 10, 2015	DP	NC	Sep 24, 2013
		6441168	Apr 17, 2020	DS	NP	Sep 24, 2013
		6958326	Dec 30, 2021	DP		
		7163931	Mar 03, 2022	U-1		
		RE37564	Jun 30, 2014	DP		
		RE37838	Jun 30, 2014	DP		
		RE38253	Jun 30, 2014	DP		
<u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM - SAFYRAL</u>						
	N022574 001	5798338	Jul 10, 2015	DP	NP	Dec 16, 2013
		6441168	Apr 17, 2020	DS		
		6958326	Dec 20, 2021	DP		
		7163931	Mar 03, 2022	U-1		
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
	N021427 001	5023269	Jun 11, 2013	DS DP U-605	I-632	Nov 04, 2013
		5023269	Jun 11, 2013	DS DP U-1094	I-617	Nov 19, 2012
		5023269	Jun 11, 2013	DS DP U-799		
		5023269	Jun 11, 2013	DS DP U-398		
		5023269	Jun 11, 2013	DS DP U-839		
		5023269	Jun 11, 2013	DS DP U-797		
		5023269	Jun 11, 2013	DS DP U-795		
		5023269	Jun 11, 2013	DS DP U-882		
		5023269	Jun 11, 2013	DS DP U-796		
		5508276	Jul 18, 2014	DP		
		6596756	Sep 10, 2019	U-882		
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
	N021427 002	5023269	Jun 11, 2013	DS DP U-1094	I-632	Nov 04, 2013
		5023269	Jun 11, 2013	DS DP U-605	I-617	Nov 19, 2012
		5023269	Jun 11, 2013	DS DP U-882		
		5023269	Jun 11, 2013	DS DP U-839		
		5023269	Jun 11, 2013	DS DP U-799		
		5023269	Jun 11, 2013	DS DP U-795		
		5023269	Jun 11, 2013	DS DP U-796		
		5023269	Jun 11, 2013	DS DP U-797		
		5023269	Jun 11, 2013	DS DP U-398		
		5508276	Jul 18, 2014	DP		
		6596756	Sep 10, 2019	U-882		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE			PATENT DELIST REQUESTED			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>									
N021427 004	5023269	Jun	11,	2013	DS	DP	U-799	I-632	Nov 04, 2013
	5023269	Jun	11,	2013	DS	DP	U-605	I-617	Nov 19, 2012
	5023269	Jun	11,	2013	DS	DP	U-1094		
	5023269	Jun	11,	2013	DS	DP	U-398		
	5023269	Jun	11,	2013	DS	DP	U-882		
	5023269	Jun	11,	2013	DS	DP	U-795		
	5023269	Jun	11,	2013	DS	DP	U-796		
	5023269	Jun	11,	2013	DS	DP	U-797		
	5023269	Jun	11,	2013	DS	DP	U-839		
	5508276	Jul	18,	2014		DP			
	6596756	Sep	10,	2019			U-882		
<u>DUTASTERIDE - AVODART</u>									
N021319 001	5565467	Nov	20,	2015	DS	DP			
	5846976	Sep	17,	2013			U-476		
	5998427	Sep	17,	2013	DS	DP	U-477		
<u>DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE - JALYN</u>									
N022460 001	5565467	Nov	20,	2015	DS	DP			
	5846976	Sep	17,	2013			U-1065		
	5998427	Sep	17,	2013	DS	DP	U-1066		
<u>EFAVIRENZ - SUSTIVA</u>									
N020972 001	5519021	May	21,	2013	DS	DP			
	5663169	Sep	02,	2014			U-257		
	5811423	Aug	07,	2012	DS	DP	U-256		
	6238695	Apr	06,	2019		DP			
	6555133	Apr	06,	2019			U-248		
	6639071	Feb	14,	2018	DS				
	6939964	Jan	20,	2018	DS				
<u>EFAVIRENZ - SUSTIVA</u>									
N020972 002	5519021	May	21,	2013	DS	DP			
	5663169	Sep	02,	2014			U-257		
	5811423	Aug	07,	2012	DS	DP	U-256		
	6238695	Apr	06,	2019		DP			
	6555133	Apr	06,	2019			U-248		
	6639071	Feb	14,	2018	DS				
	6939964	Jan	20,	2018	DS				
<u>EFAVIRENZ - SUSTIVA</u>									
N021360 001	5519021	May	21,	2013					
	5663169	Sep	02,	2014			U-256		
	5811423	Aug	07,	2012	DS				
	6639071	Feb	14,	2018	DS				
	6939964	Jan	20,	2018	DS				
<u>EFAVIRENZ - SUSTIVA</u>									
N021360 002	5519021	May	21,	2013	DS	DP			
	5663169	Sep	02,	2014			U-248		
	5811423	Aug	07,	2012			U-256		
	6639071	Feb	14,	2018	DS				
	6939964	Jan	20,	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
EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA							
N021937 001	5519021	May	21, 2013	DS DP			
	5663169	Sep	02, 2014		U-750		
	5811423	Aug	07, 2012		U-750		
	5814639	Sep	29, 2015	DS DP			
	5814639*PED	Mar	29, 2016				
	5914331	Jul	02, 2017	DS			
	5914331*PED	Jan	02, 2018				
	5922695	Jul	25, 2017	DS	U-750		
	5922695*PED	Jan	25, 2018				
	5935946	Jul	25, 2017	DS DP	U-750		
	5935946*PED	Jan	25, 2018				
	5977089	Jul	25, 2017	DS DP	U-750		
	5977089*PED	Jan	25, 2018				
	6043230	Jul	25, 2017		U-750		
	6043230*PED	Jan	25, 2018				
	6639071	Feb	14, 2018	DS			
	6642245	Nov	04, 2020		U-750		
	6642245*PED	May	04, 2021				
	6703396	Mar	09, 2021	DS DP			
	6703396*PED	Sep	09, 2021				
	6939964	Jan	20, 2018	DS			
EFLORNITHINE HYDROCHLORIDE - VANIQA							
N021145 001	5648394	Jul	15, 2014		U-334		
ELETRIPTAN HYDROBROMIDE - RELPAX							
N021016 001	5545644	Dec	26, 2016	DS DP	U-876		
	6110940	Aug	29, 2017				
ELETRIPTAN HYDROBROMIDE - RELPAX							
N021016 002	5545644	Dec	26, 2016	DS DP	U-876		
	6110940	Aug	29, 2017				
ELTROMBOPAG OLAMINE - PROMACTA							
N022291 001	6280959	Oct	30, 2018	DS DP	U-930	NCE	Nov 20, 2013
	7160870	Dec	08, 2021	DS DP	U-930	ODE	Nov 20, 2015
	7332481	May	24, 2021		U-930		
	7452874	May	24, 2021	DS DP			
	7473686	May	24, 2021	DS DP	U-930		
	7547719	Jul	13, 2025	DS DP	U-930		
	7790704	May	24, 2021		U-930		
	7795293	May	21, 2023		U-930		
	8052993	Aug	01, 2027	DP	U-930		
ELTROMBOPAG OLAMINE - PROMACTA							
N022291 002	6280959	Oct	30, 2018	DS DP	U-930	NCE	Nov 20, 2013
	7160870	Dec	08, 2021	DS DP	U-930	ODE	Nov 20, 2015
	7332481	May	24, 2021		U-930		
	7452874	May	24, 2021	DS DP			
	7473686	May	24, 2021	DS DP	U-930		
	7547719	Jul	13, 2025	DS DP	U-930		
	7790704	May	24, 2021		U-930		
	7795293	May	21, 2023		U-930		
	8052994	Aug	01, 2027	DP	U-930		
ELTROMBOPAG OLAMINE - PROMACTA							
N022291 003	6280959	Oct	30, 2018	DS DP	U-930	NCE	Nov 20, 2013
	7160870	Dec	08, 2021	DS DP	U-930	ODE	Nov 20, 2015
	7332481	May	24, 2021		U-930		
	7452874	May	24, 2021	DS DP			
	7473686	May	24, 2021	DS DP	U-930		
	7547719	Jul	13, 2025	DS DP	U-930		
	7790704	May	24, 2021		U-930		
	7795293	May	21, 2023		U-930		
	8062665	Aug	01, 2027	DP	U-930		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE			PATENT DELIST REQUESTED			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>										
N022291 004	6280959	Oct	30,	2018	DS	DP	U-930		NCE	Nov 20, 2013
	7160870	Dec	08,	2021	DS	DP	U-930		ODE	Nov 20, 2015
	7332481	May	24,	2021			U-930			
	7452874	May	24,	2021	DS	DP				
	7473686	May	24,	2021	DS	DP	U-930			
	7547719	Jul	13,	2025	DS	DP	U-930			
	7790704	May	24,	2021			U-930			
	7795293	May	21,	2023			U-930			
	8071129	Aug	01,	2027		DP	U-930			
<u>EMEDASTINE DIFUMARATE - EMADINE</u>										
N020706 001	5441958	Dec	08,	2013			U-404			
<u>EMTRICITABINE - EMTRIVA</u>										
N021500 001	5814639	Sep	29,	2015	DS	DP				
	5814639*PED	Mar	29,	2016						
	5914331	Jul	02,	2017	DS					
	5914331*PED	Jan	02,	2018						
	6642245	Nov	04,	2020			U-257			
	6642245	Nov	04,	2020			U-541			
	6642245*PED	May	04,	2021						
	6703396	Mar	09,	2021	DS	DP				
	6703396*PED	Sep	09,	2021						
<u>EMTRICITABINE - EMTRIVA</u>										
N021896 001	5814639	Sep	29,	2015	DS	DP				
	5814639*PED	Mar	29,	2016						
	5914331	Jul	02,	2017	DS					
	5914331*PED	Jan	02,	2018						
	6642245	Nov	04,	2020			U-257			
	6642245*PED	May	04,	2021						
	6703396	Mar	09,	2021	DS	DP				
	6703396*PED	Sep	09,	2021						
<u>EMTRICITABINE; RILPIVIRINE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u>										
N202123 001	5814639	Sep	29,	2015	DS	DP				
	5814639*PED	Mar	29,	2016						
	5914331	Jul	02,	2017	DS					
	5914331*PED	Jan	02,	2018						
	5922695	Jul	25,	2017	DS		U-257			
	5922695*PED	Jan	25,	2018						
	5935946	Jul	25,	2017	DS	DP	U-257			
	5935946*PED	Jan	25,	2018						
	5977089	Jul	25,	2017	DS	DP				
	5977089*PED	Jan	25,	2018			U-257			
	6043230	Jul	25,	2017			U-257			
	6043230*PED	Jan	25,	2018						
	6642245	Nov	04,	2020			U-257			
	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	14,	2023	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
EMTRICITABINE: TENOFOVIR DISOPROXIL FUMARATE - TRUVADA								
N021752 001	5814639	Sep	29, 2015	DS	DP			
	5814639*PED	Mar	29, 2016					
	5914331	Jul	02, 2017	DS	DP	U-248		
	5914331*PED	Jan	02, 2018	DS				
	5922695	Jul	25, 2017	DS		U-248		
	5922695	Jul	25, 2017	DS		U-541		
	5922695	Jul	25, 2017	DS		U-1170		
	5922695*PED	Jan	25, 2018					
	5935946	Jul	25, 2017	DS	DP	U-541		
	5935946	Jul	25, 2017	DS	DP	U-248		
	5935946	Jul	25, 2017	DS	DP	U-1170		
	5935946*PED	Jan	25, 2018					
	5977089	Jul	25, 2017	DS	DP	U-1170		
	5977089	Jul	25, 2017	DS	DP	U-541		
	5977089	Jul	25, 2017	DS	DP	U-248		
	5977089*PED	Jan	25, 2018					
	6043230	Jul	25, 2017		DP	U-248		
	6043230	Jul	25, 2017		DP	U-1170		
	6043230	Jul	25, 2017		DP	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					
ENFUVIRTIDE - FUZEON								
N021481 001	5464933	Jun	07, 2013					
	6133418	Nov	17, 2014	DS	DP			
	6475491	Jun	07, 2015			U-248		
ENOXAPARIN SODIUM - LOVENOX								
N020164 009	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 001	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 002	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 003	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 004	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 005	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 006	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 007	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		
ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)								
N020164 008	5389618	Feb	14, 2012	DS	DP	U-545		
	RE38743	Feb	14, 2012	DS	DP	U-545		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ENTACAPONE - COMTAN</u>						
N020796 001	5446194	Oct 19, 2013	DS			
	6599530	Sep 14, 2018	DP U-219			
<u>ENTECAVIR - BARACLUDÉ</u>						
N021797 001	5206244	Feb 21, 2015	DS		D-127	Oct 15, 2013
<u>ENTECAVIR - BARACLUDÉ</u>						
N021797 002	5206244	Feb 21, 2015	DS		D-127	Oct 15, 2013
<u>ENTECAVIR - BARACLUDÉ</u>						
N021798 001	5206244	Feb 21, 2015	DS		D-127	Oct 15, 2013
<u>EPINASTINE HYDROCHLORIDE - ELESTAT</u>						
N021565 001	7429602	Nov 29, 2020	U-765	Y		
<u>EPINASTINE HYDROCHLORIDE - EPINASTINE HYDROCHLORIDE</u>						
A090870 001					PC	Oct 29, 2011
<u>EPINEPHRINE - ADRENACCLICK</u>						
N020800 003	5665071	May 27, 2013	DP			
<u>EPINEPHRINE - ADRENACCLICK</u>						
N020800 004	5665071	May 27, 2013	DP			
<u>EPINEPHRINE - EPIPEN</u>						
N019430 001	7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
<u>EPINEPHRINE - EPIPEN JR.</u>						
N019430 002	7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
<u>EPINEPHRINE - TWINJECT 0.15</u>						
N020800 002	7297136	Jan 18, 2025	DP			
	7621891	Feb 04, 2025	DP			
<u>EPINEPHRINE - TWINJECT 0.3</u>						
N020800 001	7297136	Jan 18, 2025	DP			
	7621891	Feb 04, 2025	DP			
<u>EPINEPHRINE; LIDOCAINE HYDROCHLORIDE - LIDOSITE TOPICAL SYSTEM KIT</u>						
N021504 001	5246418	Sep 30, 2013	DS DP			
	5873850	Sep 30, 2013	DS DP			
	6377847	Sep 30, 2013	DS DP			
	6385488	Sep 30, 2013	DS DP			
	6629968	Jun 30, 2020	DS DP			
	6635045	Jun 29, 2021	DS DP			
	6862473	Sep 30, 2013	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
EPLERENONE - INSPRA						
N021437 001	6410054	Dec 08, 2019	U-3			
	6410054	Dec 08, 2019	U-537			
	6410054*PED	Jun 08, 2020				
	6410524	Nov 05, 2019	U-467			
	6410524*PED	May 05, 2020				
	6495165	Dec 08, 2019	U-3			
	6495165	Dec 08, 2019	U-537			
	6495165*PED	Jun 08, 2020				
	6534093	Dec 08, 2019	U-537			
	6534093	Dec 08, 2019	U-3			
	6534093*PED	Jun 08, 2020				
	6558707	Dec 08, 2019	DP U-537			
	6558707*PED	Jun 08, 2020				
	6747020	Nov 05, 2019	U-587			
	6747020*PED	May 05, 2020				
	6863902	Apr 10, 2020	DP U-664			
	6863902*PED	Oct 10, 2020				
	7157101	Dec 08, 2019	DP U-664			
	7157101*PED	Jun 08, 2020				
EPLERENONE - INSPRA						
N021437 002	6410054	Dec 08, 2019	U-3			
	6410054	Dec 08, 2019	U-537			
	6410054*PED	Jun 08, 2020				
	6410524	Nov 05, 2019	U-467			
	6410524*PED	May 05, 2020				
	6495165	Dec 08, 2019	U-3			
	6495165	Dec 08, 2019	U-537			
	6495165*PED	Jun 08, 2020				
	6534093	Dec 08, 2019	U-3			
	6534093	Dec 08, 2019	U-537			
	6534093*PED	Jun 08, 2020				
	6558707	Dec 08, 2019	DP U-537			
	6558707*PED	Jun 08, 2020				
	6747020	Nov 05, 2019	U-587			
	6747020*PED	May 05, 2020				
	6863902	Apr 10, 2020	DP U-664			
	6863902*PED	Oct 10, 2020				
	7157101	Dec 08, 2019	DP U-664			
	7157101*PED	Jun 08, 2020				
EPLERENONE - INSPRA						
N021437 003	6410054	Dec 08, 2019	U-3			
	6410054	Dec 08, 2019	U-537			
	6410054*PED	Jun 08, 2020				
	6410524	Nov 05, 2019	U-467			
	6410524*PED	May 05, 2020				
	6495165	Dec 08, 2019	U-3			
	6495165	Dec 08, 2019	U-537			
	6495165*PED	Jun 08, 2020				
	6534093	Dec 08, 2019	U-3			
	6534093	Dec 08, 2019	U-537			
	6534093*PED	Jun 08, 2020				
	6558707	Dec 08, 2019	DP U-537			
	6558707*PED	Jun 08, 2020				
	6747020	Nov 05, 2019	U-587			
	6747020*PED	May 05, 2020				
	6863902	Apr 10, 2020	DP U-664			
	6863902*PED	Oct 10, 2020				
	7157101	Dec 08, 2019	DP U-664			
	7157101*PED	Jun 08, 2020				
EPROSARTAN MESYLATE - TEVETEN						
N020738 004	5656650	Aug 12, 2014	U-3			
EPROSARTAN MESYLATE - TEVETEN						
N020738 005	5656650	Aug 12, 2014	U-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
<u>EPROSARTAN MESYLATE - TEVETEN</u>						
	N020738 006	5656650	Aug 12, 2014	U-3		
<u>EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE - TEVETEN HCT</u>						
	N021268 001	5656650	Aug 12, 2014	U-3		
<u>EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE - TEVETEN HCT</u>						
	N021268 002	5656650	Aug 12, 2014	U-3		
<u>EPTIFIBATIDE - INTEGRILIN</u>						
	N020718 001	5686570	Nov 11, 2014			
		5747447	May 05, 2015			
		5756451	Nov 11, 2014			
		5807825	Sep 15, 2015	U-244		
		5968902	Jun 02, 2015	U-453		
<u>EPTIFIBATIDE - INTEGRILIN</u>						
	N020718 002	5686570	Nov 11, 2014			
		5747447	May 05, 2015			
		5756451	Nov 11, 2014			
		5807825	Sep 15, 2015	U-244		
		5968902	Jun 02, 2015	U-453		
<u>ERIBULIN MESYLATE - HALAVEN</u>						
	N201532 001	6214865	Jun 16, 2019	DS		
		6469182	Jun 16, 2019		U-1096	
		7470720	Jun 16, 2019	DP		
<u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u>						
	N021743 001	5747498	Nov 08, 2018	DS DP	U-659	
		6900221	Nov 09, 2020	DS DP	U-875	
		6900221	Nov 09, 2020	DS DP	U-659	
		6900221	Nov 09, 2020	DS DP	U-1046	
		7087613	Nov 09, 2020		U-659	
		7087613	Nov 09, 2020		U-1045	
		RE41065	Nov 08, 2018	DS DP		
<u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u>						
	N021743 002	5747498	Nov 08, 2018	DS DP	U-659	
		6900221	Nov 09, 2020	DS DP	U-875	
		6900221	Nov 09, 2020	DS DP	U-659	
		6900221	Nov 09, 2020	DS DP	U-1046	
		7087613	Nov 09, 2020		U-659	
		7087613	Nov 09, 2020		U-1045	
		RE41065	Nov 08, 2018	DS DP		
<u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u>						
	N021743 003	5747498	Nov 08, 2018	DS DP	U-659	
		6900221	Nov 09, 2020	DS DP	U-875	
		6900221	Nov 09, 2020	DS DP	U-1046	
		6900221	Nov 09, 2020	DS DP	U-659	
		7087613	Nov 09, 2020		U-659	
		7087613	Nov 09, 2020		U-1045	
		RE41065	Nov 08, 2018	DS DP		
<u>ERTAPENEM SODIUM - INVANZ</u>						
	N021337 001	5478820	Nov 21, 2015	DS DP	U-160	
		5478820*PED	May 21, 2016			
		5652233	Feb 02, 2013	DS DP	U-160	
		5652233*PED	Aug 02, 2013			
		5952323	May 15, 2017	DP		
		5952323*PED	Nov 15, 2017			
		7342005	Feb 02, 2013	DP		
		7342005*PED	Aug 02, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ESCITALOPRAM OXALATE - LEXAPRO</u>							
N021323 001	6916941	Aug	12, 2022	DS DP		NPP	Mar 19, 2012
	6916941*PED	Feb	12, 2023	DS DP			
	7420069	Aug	12, 2022	DP			
	7420069*PED	Feb	12, 2023				
	RE34712	Sep	14, 2011	DS DP			
	RE34712*PED	Mar	14, 2012				
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>							
N021323 002	6916941	Aug	12, 2022	DS DP		NPP	Mar 19, 2012
	6916941*PED	Feb	12, 2023	DS DP			
	7420069	Aug	12, 2022	DP			
	7420069*PED	Feb	12, 2023				
	RE34712	Sep	14, 2011	DS DP			
	RE34712*PED	Mar	14, 2012				
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>							
N021323 003	6916941	Aug	12, 2022	DS DP		NPP	Mar 19, 2012
	6916941*PED	Feb	12, 2023	DS DP			
	7420069	Aug	12, 2022	DP			
	7420069*PED	Feb	12, 2023				
	RE34712	Sep	14, 2011	DS DP			
	RE34712*PED	Mar	14, 2012				
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>							
N021365 001	RE34712	Sep	14, 2011	DS DP		NPP	Mar 19, 2012
	RE34712*PED	Mar	14, 2012				
<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC</u>							
N019386 006	6310094	Jan	12, 2021				
	6310094*PED	Jul	12, 2021				
	6528540	Jan	12, 2021				
	6528540*PED	Jul	12, 2021				
<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC</u>							
N019386 007	6310094	Jan	12, 2021				
	6310094*PED	Jul	12, 2021				
	6528540	Jan	12, 2021				
	6528540*PED	Jul	12, 2021				
<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER</u>							
N019386 005	6310094	Jan	12, 2021				
	6310094*PED	Jul	12, 2021				
	6528540	Jan	12, 2021				
	6528540*PED	Jul	12, 2021				
<u>ESMOLOL HYDROCHLORIDE - BREVIBLOC IN PLASTIC CONTAINER</u>							
N019386 004	6310094	Jan	12, 2021				
	6310094*PED	Jul	12, 2021				
	6528540	Jan	12, 2021				
	6528540*PED	Jul	12, 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
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
N021153 001	5690960	Nov 25, 2014	DP U-373			
	5690960	Nov 25, 2014	DP U-729			
	5690960	Nov 25, 2014	DP U-770			
	5690960*PED	May 25, 2015	U-373			
	5714504	Feb 03, 2015	DP U-729			
	5714504	Feb 03, 2015	DP U-770			
	5714504	Feb 03, 2015	DP U-373			
	5714504*PED	Aug 03, 2015	U-373			
	5877192	May 27, 2014	DP U-729			
	5877192	May 27, 2014	DP U-770			
	5877192	May 27, 2014	DP U-373			
	5877192*PED	Nov 27, 2014	U-373			
	5900424	May 04, 2016	DS			
	5900424	May 04, 2016	DS			
	5900424	May 04, 2016	DS			
	5900424*PED	Nov 04, 2016	U-373			
	6147103	Oct 09, 2018				
	6147103*PED	Apr 09, 2019				
	6166213	Oct 09, 2018				
	6166213*PED	Apr 09, 2019				
	6191148	Oct 09, 2018				
	6191148*PED	Apr 09, 2019				
	6369085	May 25, 2018	DS DP U-770			
	6369085	May 25, 2018	DS DP U-729			
	6369085*PED	Nov 25, 2018				
	6428810	Nov 03, 2019	DP U-770			
	6428810	Nov 03, 2019	DP U-469			
	6428810	Nov 03, 2019	DP U-729			
	6428810*PED	May 03, 2020	U-469			
	6875872	May 27, 2014	DS			
	6875872*PED	Nov 27, 2014				
	7411070	May 25, 2018	DS			
	7411070*PED	Nov 25, 2018				
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
N021153 002	5690960	Nov 25, 2014	DP U-770			
	5690960	Nov 25, 2014	DP U-373			
	5690960	Nov 25, 2014	DP U-729			
	5690960*PED	May 25, 2015	U-373			
	5714504	Feb 03, 2015	DP U-770			
	5714504	Feb 03, 2015	DP U-729			
	5714504	Feb 03, 2015	DP U-373			
	5714504*PED	Aug 03, 2015	U-373			
	5877192	May 27, 2014	DP U-770			
	5877192	May 27, 2014	DP U-729			
	5877192	May 27, 2014	DP U-373			
	5877192*PED	Nov 27, 2014	U-373			
	5900424	May 04, 2016	DS			
	5900424	May 04, 2016	DS			
	5900424	May 04, 2016	DS			
	5900424*PED	Nov 04, 2016	U-373			
	6147103	Oct 09, 2018				
	6147103*PED	Apr 09, 2019				
	6166213	Oct 09, 2018				
	6166213*PED	Apr 09, 2019				
	6191148	Oct 09, 2018				
	6191148*PED	Apr 09, 2019				
	6369085	May 25, 2018	DS DP U-729			
	6369085	May 25, 2018	DS DP U-770			
	6369085*PED	Nov 25, 2018				
	6428810	Nov 03, 2019	DP U-770			
	6428810	Nov 03, 2019	DP U-729			
	6428810	Nov 03, 2019	DP U-469			
	6428810*PED	May 03, 2020	U-469			
	6875872	May 27, 2014	DS			
	6875872*PED	Nov 27, 2014				
	7411070	May 25, 2018	DS			
	7411070*PED	Nov 25, 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
ESOMEPRAZOLE MAGNESIUM - NEXIUM						
N021957 001	5690960	Nov 25, 2014	DP U-1207		M-86	Jun 18, 2012
	5690960	Nov 25, 2014	DP U-729		PED	Aug 27, 2011
	5690960	Nov 25, 2014	DP U-773		PED	Dec 18, 2012
	5690960*PED	May 25, 2015				
	5714504	Feb 03, 2015	DP U-729			
	5714504	Feb 03, 2015	DP U-773			
	5714504	Feb 03, 2015	DP U-1207			
	5714504*PED	Aug 03, 2015				
	5877192	May 27, 2014	U-773			
	5877192	May 27, 2014	U-729			
	5877192*PED	Nov 27, 2014				
	5900424	May 04, 2016	DS U-1207			
	5900424	May 04, 2016	DS U-729			
	5900424	May 04, 2016	DS U-773			
	5900424*PED	Nov 04, 2016				
	6369085	May 25, 2018	DS DP U-729			
	6369085	May 25, 2018	DS DP U-773			
	6369085	May 25, 2018	DS DP U-1207			
	6369085*PED	Nov 25, 2018				
	6428810	Nov 03, 2019	DP U-729			
	6428810	Nov 03, 2019	DP U-773			
	6428810	Nov 03, 2019	DP U-1207			
	6428810*PED	May 03, 2020				
	6875872	May 27, 2014	DS			
	6875872*PED	Nov 27, 2014				
	7411070	May 25, 2018	DS			
	7411070*PED	Nov 25, 2018				
ESOMEPRAZOLE MAGNESIUM - NEXIUM						
N021957 002	5690960	Nov 25, 2014	DP U-1207		M-86	Jun 18, 2012
	5690960	Nov 25, 2014	DP U-729		PED	Aug 27, 2011
	5690960	Nov 25, 2014	DP U-773		PED	Dec 18, 2012
	5690960*PED	May 25, 2015				
	5714504	Feb 03, 2015	DP U-1207			
	5714504	Feb 03, 2015	DP U-729			
	5714504	Feb 03, 2015	DP U-773			
	5714504*PED	Aug 03, 2015				
	5877192	May 27, 2014	U-729			
	5877192	May 27, 2014	U-773			
	5877192*PED	Nov 27, 2014				
	5900424	May 04, 2016	DS U-1207			
	5900424	May 04, 2016	DS U-729			
	5900424	May 04, 2016	DS U-773			
	5900424*PED	Nov 04, 2016				
	6369085	May 25, 2018	DS DP U-1207			
	6369085	May 25, 2018	DS DP U-729			
	6369085	May 25, 2018	DS DP U-773			
	6369085*PED	Nov 25, 2018				
	6428810	Nov 03, 2019	DP U-1207			
	6428810	Nov 03, 2019	DP U-729			
	6428810	Nov 03, 2019	DP U-773			
	6428810*PED	May 03, 2020				
	6875872	May 27, 2014	DS			
	6875872*PED	Nov 27, 2014				
	7411070	May 25, 2018	DS			
	7411070*PED	Nov 25, 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
ESOMEPRAZOLE MAGNESIUM - NEXIUM							
N022101 001	5690960	Nov	25, 2014	DP U-858		M-86	Jun 18, 2012
	5690960*PED	May	25, 2015			PED	Dec 18, 2012
	5714504	Feb	03, 2015	DP U-858		PED	Aug 27, 2011
	5714504*PED	Aug	03, 2015				
	5877192	May	27, 2014	U-858			
	5877192*PED	Nov	27, 2014				
	5900424	May	04, 2016	DS U-858			
	5900424*PED	Nov	04, 2016				
	6369085	May	25, 2018	DS DP U-858			
	6369085*PED	Nov	25, 2018				
	6428810	Nov	03, 2019	DP U-858			
	6428810*PED	May	03, 2020				
	6875872	May	27, 2014	DS			
	6875872*PED	Nov	27, 2014				
	7411070	May	25, 2018	DS			
	7411070*PED	Nov	25, 2018				
ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO							
N022511 001	5714504	Feb	03, 2015	DP U-1053		NC	Apr 30, 2013
	5714504*PED	Aug	03, 2015				
	5900424	May	04, 2016	DS U-1053			
	5900424*PED	Nov	04, 2016				
	6369085	May	25, 2018	DS DP U-1053			
	6369085*PED	Nov	25, 2018				
	6875872	May	27, 2014	DS			
	6875872*PED	Nov	27, 2014				
	6926907	Feb	28, 2023	DP U-1052			
	7411070	May	25, 2018	DS U-1053			
	7411070*PED	Nov	25, 2018				
	7745466	Oct	13, 2018	DP U-1053			
ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO							
N022511 002	5714504	Feb	03, 2015	DP U-1053		NC	Apr 30, 2013
	5714504*PED	Aug	03, 2015				
	5900424	May	04, 2016	DS U-1053			
	5900424*PED	Nov	04, 2016				
	6369085	May	25, 2018	DS DP U-1053			
	6369085*PED	Nov	25, 2018				
	6875872	May	27, 2014	DS			
	6875872*PED	Nov	27, 2014				
	6926907	Feb	28, 2023	DP U-1052			
	7411070	May	25, 2018	DS U-1053			
	7411070*PED	Nov	25, 2018				
	7745466	Oct	13, 2018	DP U-1053			
ESOMEPRAZOLE SODIUM - NEXIUM IV							
N021689 001	5877192	May	27, 2014	U-643			
	5877192*PED	Nov	27, 2014				
	6143771	May	27, 2014	DP U-643			
ESOMEPRAZOLE SODIUM - NEXIUM IV							
N021689 002	5877192	May	27, 2014	U-643			
	5877192*PED	Nov	27, 2014				
	6143771	May	27, 2014	DP U-643			
ESTRADIOL - ELESTRIN							
N021813 001	7198801	Jun	25, 2022	DP			
	7470433	Aug	03, 2021	DP			
ESTRADIOL - EVAMIST							
N022014 001	6299900	Feb	19, 2017	DP U-889			
	6299900	Feb	19, 2017	DP U-888			
	6818226	Feb	19, 2017	DP U-888			
	6818226	Feb	19, 2017	DP U-889			
	6923983	Feb	19, 2017	DP U-888			
	6923983	Feb	19, 2017	DP U-889			
	6978945	Nov	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
<u>ESTRADIOL - MENOSTAR</u>						
N021674 001	5891868	Nov 21, 2017	DP	U-594		
	6692763	Nov 21, 2017	DP	U-594		
<u>ESTRADIOL - VAGIFEM</u>						
N020908 002	7018992	Sep 17, 2022		U-1023		D-122 Nov 25, 2012
<u>ESTRADIOL - VIVELLE-DOT</u>						
N020538 005	5474783	Dec 12, 2012				
	5656286	Aug 12, 2014				
	5958446	Dec 12, 2012				
	6024976	Jan 07, 2014				
<u>ESTRADIOL - VIVELLE-DOT</u>						
N020538 006	5474783	Dec 12, 2012				
	5656286	Aug 12, 2014				
	5958446	Dec 12, 2012				
	6024976	Jan 07, 2014				
<u>ESTRADIOL - VIVELLE-DOT</u>						
N020538 007	5474783	Dec 12, 2012				
	5656286	Aug 12, 2014				
	5958446	Dec 12, 2012				
	6024976	Jan 07, 2014				
<u>ESTRADIOL - VIVELLE-DOT</u>						
N020538 008	5474783	Dec 12, 2012				
	5656286	Aug 12, 2014				
	5958446	Dec 12, 2012				
	6024976	Jan 07, 2014				
<u>ESTRADIOL - VIVELLE-DOT</u>						
N020538 009	5474783	Dec 12, 2012	DP			
	5656286	Aug 12, 2014	DP			
	5958446	Dec 12, 2012	DP			
	6024976	Jan 07, 2014	DP			
<u>ESTRADIOL ACETATE - FMRING</u>						
N021367 001	5855906	Dec 19, 2015		U-508		
<u>ESTRADIOL ACETATE - FMRING</u>						
N021367 002	5855906	Dec 19, 2015		U-508		
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N021633 001	6962908	Dec 21, 2021	DP			
	7572779	Oct 02, 2025		U-904		
	7799771	Dec 21, 2021	DP			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N021633 002	6962908	Dec 21, 2021	DP			
	7572779	Oct 02, 2025		U-904		
	7799771	Dec 21, 2021	DP			
<u>ESTRADIOL ACETATE - FEMTRACE</u>						
N021633 003	6962908	Dec 21, 2021	DP			
	7572779	Oct 02, 2025		U-904		
	7799771	Dec 21, 2021	DP			
<u>ESTRADIOL HEMIHYDRATE - ESTRASORB</u>						
N021371 001	5629021	Jan 31, 2015	DP			
<u>ESTRADIOL; LEVONORGESTREL - CLIMARA PRO</u>						
N021258 001	5676968	Oct 14, 2014	DP			
<u>ESTRADIOL; NORETHINDRONE ACETATE - COMBIPATCH</u>						
N020870 001	5474783	Dec 12, 2012				
	5656286	Aug 12, 2014				
	5958446	Dec 12, 2012				
	6024976	Jan 07, 2014				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ESTRADIOL; NORETHINDRONE ACETATE - COMBIPATCH</u>						
N020870 002	5474783	Dec 12, 2012				
	5656286	Aug 12, 2014				
	5958446	Dec 12, 2012				
	6024976	Jan 07, 2014				
<u>ESTRADIOL; NORGESTIMATE - PREFEST</u>						
N021040 001	5382573	Jan 17, 2012				
	6747019	Mar 20, 2020		U-311		
	7320970	Mar 30, 2020	DP	U-844		
<u>ESTROGENS, CONJUGATED - PREMARIN</u>						
N020216 001					I-579	Nov 07, 2011
<u>ESTROGENS, CONJUGATED SYNTHETIC A - CENESTIN</u>						
N020992 001	5908638	Jul 26, 2015		DP		
<u>ESTROGENS, CONJUGATED SYNTHETIC A - CENESTIN</u>						
N020992 002	5908638	Jul 26, 2015				
<u>ESTROGENS, CONJUGATED SYNTHETIC A - CENESTIN</u>						
N020992 003	5908638	Jul 26, 2015				
<u>ESTROGENS, CONJUGATED SYNTHETIC A - CENESTIN</u>						
N020992 004	5908638	Jul 26, 2015				
<u>ESTROGENS, CONJUGATED SYNTHETIC A - CENESTIN</u>						
N020992 005	5908638	Jul 26, 2015				
<u>ESTROGENS, CONJUGATED SYNTHETIC A - SYNTHETIC CONJUGATED ESTROGENS A</u>						
N021788 001					NP	Nov 28, 2011
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u>						
N021443 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	
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u>						
N021443 002	6660726	Mar 08, 2021	DS	DP	U-904	
	6660726	Mar 08, 2021	DS	DP	U-905	
	6855703	Feb 12, 2021	DS	DP	U-905	
	6855703	Feb 12, 2021	DS	DP	U-904	
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u>						
N021443 003	6660726	Mar 08, 2021	DS	DP	U-905	
	6660726	Mar 08, 2021	DS	DP	U-904	
	6855703	Feb 12, 2021	DS	DP	U-904	
	6855703	Feb 12, 2021	DS	DP	U-905	
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u>						
N021443 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	
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u>						
N021443 005	6660726	Mar 08, 2021	DS	DP	U-905	
	6660726	Mar 08, 2021	DS	DP	U-904	
	6855703	Feb 12, 2021	DS	DP	U-904	
	6855703	Feb 12, 2021	DS	DP	U-905	
<u>ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE - PREMPHASE 14/14</u>						
N020527 002	5547948	Jan 17, 2015				
<u>ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE - PREMPRO</u>						
N020527 001	5547948	Jan 17, 2015				
<u>ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE - PREMPRO</u>						
N020527 003	5547948	Jan 17, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE - PREMPRO</u>						
N020527 004	5547948	Jan 17, 2015				
<u>ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE - PREMPRO</u>						
N020527 005	5547948	Jan 17, 2015				
<u>ESZOPICLONE - LUNESTA</u>						
N021476 001	6319926	Jan 16, 2012		U-620		
	6444673	Feb 14, 2014	DS DP			
	6864257	Aug 30, 2012		U-629		
	7381724	Jan 16, 2012	DS DP	U-629		
<u>ESZOPICLONE - LUNESTA</u>						
N021476 002	6319926	Jan 16, 2012		U-620		
	6444673	Feb 14, 2014	DS DP			
	6864257	Aug 30, 2012		U-629		
	7381724	Jan 16, 2012	DS DP	U-629		
<u>ESZOPICLONE - LUNESTA</u>						
N021476 003	6319926	Jan 16, 2012		U-620		
	6444673	Feb 14, 2014	DS DP			
	6864257	Aug 30, 2012		U-629		
	7381724	Jan 16, 2012	DS DP	U-629		
<u>ETHINYL ESTRADIOL; ETONOGESTREL - NUVARING</u>						
N021187 001	5989581	Apr 08, 2018				
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - LOSEASONIQUE</u>						
N022262 001	7615545	Jun 15, 2023		U-1		
	7855190	Dec 05, 2028		U-1		
	7858605	Jun 23, 2023	DP			
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - LYBREL</u>						
N021864 001	6500814	Sep 03, 2018		U-1		
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - PREVEN EMERGENCY CONTRACEPTIVE KIT</u>						
N020946 001	6156742	Dec 05, 2020		U-374		
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONALE</u>						
N021544 001	5898032	Jun 23, 2017		U-1		
	RE39861	Jun 23, 2017		U-828		
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u>						
N021840 001	7320969	Jan 30, 2024		U-828		
	7615545	Jun 15, 2023		U-1		
	7855190	Dec 05, 2028		U-1		
	7858605	Jun 23, 2023	DP			
<u>ETHINYL ESTRADIOL; NORELGESTROMIN - ORTHO EVRA</u>						
N021180 001	5876746	Nov 20, 2015	DP	U-514		
	5972377	Jun 07, 2015		U-514		
<u>ETHINYL ESTRADIOL; NORETHINDRONE - FEMCON FE</u>						
N021490 001	6667050	Apr 06, 2019	DP	U-1		
<u>ETHINYL ESTRADIOL; NORETHINDRONE - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>						
A078965 001					PC	Sep 21, 2011
<u>ETHINYL ESTRADIOL; NORETHINDRONE - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>						
N022573 001	5552394	Jul 22, 2014		U-828		
	6667050	Apr 06, 2019	DP	U-828		
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u>						
N022501 001	5552394	Jul 22, 2014		U-1090		
	7704984	Feb 02, 2029		U-1090		
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LOESTRIN 24 FE</u>						
N021871 001	5552394	Jul 22, 2014		U-1		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ETHINYLN ESTRADIOL; NORGESTIMATE - ORTHO TRI-CYCLEN LO</u>						
N021241 001	6214815	Jun 09, 2019		U-112		
	6214815*PED	Dec 09, 2019				
<u>ETRAVIRINE - INTELENCE</u>						
N022187 001	6878717	Nov 05, 2019		U-256		
	6878717	Nov 05, 2019		U-1016		
	7037917	Dec 13, 2020	DS DP	U-256		
	7037917	Dec 13, 2020	DS DP	U-1016		
	7887845	Mar 25, 2019	DP			
	8003789	Nov 01, 2019				
<u>ETRAVIRINE - INTELENCE</u>						
N022187 002	6878717	Nov 05, 2019		U-1016		
	7037917	Dec 13, 2020	DS DP	U-1016		
	7887845	Mar 25, 2019	DP			
	8003789	Nov 01, 2019				
<u>EVEROLIMUS - AFINITOR</u>						
N022334 001	5665772	Sep 09, 2019	DS DP		I-638	May 05, 2014
	6004973	Jul 12, 2016	DP		I-630	Oct 29, 2013
	7297703	Dec 06, 2019	DP		NCE	Mar 30, 2014
					ODE	May 05, 2018
					ODE	Oct 29, 2017
<u>EVEROLIMUS - AFINITOR</u>						
N022334 002	5665772	Sep 09, 2019	DS DP		I-638	May 05, 2014
	6004973	Jul 12, 2016	DP		I-630	Oct 29, 2013
	7297703	Dec 06, 2019	DP		NCE	Mar 30, 2014
					ODE	May 05, 2018
					ODE	Oct 29, 2017
<u>EVEROLIMUS - AFINITOR</u>						
N022334 003	5665772	Sep 09, 2019			I-638	May 05, 2014
					I-630	Oct 29, 2013
					NCE	Mar 30, 2014
					ODE	May 05, 2018
					ODE	Oct 29, 2017
<u>EVEROLIMUS - ZORTRESS</u>						
N021560 001	5665772	Sep 09, 2019	DS DP	U-1049	NP	Apr 20, 2013
	6004973	Jul 12, 2016	DP	U-1049	NCE	Mar 30, 2014
	6239124	Aug 11, 2017		U-1049		
	6440990	Sep 24, 2013	DP	U-1049		
	6455518	Jul 29, 2017		U-1049		
<u>EVEROLIMUS - ZORTRESS</u>						
N021560 002	5665772	Sep 09, 2019	DS DP	U-1049	NP	Apr 20, 2013
	6004973	Jul 12, 2016	DP	U-1049	NCE	Mar 30, 2014
	6239124	Aug 11, 2017		U-1049		
	6440990	Sep 24, 2013	DP	U-1049		
	6455518	Jul 29, 2017		U-1049		
<u>EVEROLIMUS - ZORTRESS</u>						
N021560 003	5665772	Sep 09, 2019	DS DP	U-1049	NP	Apr 20, 2013
	6004973	Jul 12, 2016	DP	U-1049	NCE	Mar 30, 2014
	6239124	Aug 11, 2017		U-1049		
	6440990	Sep 24, 2013	DP	U-1049		
	6455518	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
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N021773 001	5424286	Dec 01, 2016	U-653			
	5424286	Dec 01, 2016	U-1108			
	6858576	Jan 06, 2017	U-656			
	6872700	Jan 14, 2020	U-654			
	6902744	Jan 14, 2020	DP			
	6956026	Jan 07, 2018	U-687			
	7297761	Oct 15, 2017	DP			
	7521423	Oct 15, 2017	DP			
	7741269	Jan 07, 2018	U-1074			
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
N021773 002	5424286	Dec 01, 2016	U-1108			
	5424286	Dec 01, 2016	U-653			
	6858576	Jan 06, 2017	U-656			
	6872700	Jan 14, 2020	U-654			
	6902744	Jan 14, 2020	DP			
	6956026	Jan 07, 2018	U-687			
	7297761	Oct 15, 2017	DP			
	7521423	Oct 15, 2017	DP			
	7741269	Jan 07, 2018	U-1074			
<u>EZETIMIBE - ZETIA</u>						
N021445 001	5846966	Sep 21, 2013	U-1172		PED	Dec 05, 2011
	5846966	Sep 21, 2013	U-474			
	5846966	Sep 21, 2013	U-473			
	5846966*PED	Mar 21, 2014				
	7030106	Jan 25, 2022	DP			
	7030106*PED	Jul 25, 2022				
	7612058	Jan 25, 2022	U-1027			
	7612058	Jan 25, 2022	U-1173			
	7612058*PED	Jul 25, 2022				
	RE37721	Oct 25, 2016	DS DP U-473			
	RE37721*PED	Apr 25, 2017				
	RE42461	Oct 25, 2016	DS DP U-1173			
	RE42461	Oct 25, 2016	DS DP U-473			
	RE42461*PED	Apr 25, 2017				
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 001	5846966	Sep 21, 2013	DP U-473		PED	Dec 05, 2011
	5846966	Sep 21, 2013	DP U-593			
	5846966*PED	Mar 21, 2014				
	RE37721	Oct 25, 2016	DS DP U-473	Y		
	RE37721*PED	Apr 25, 2017		Y		
	RE42461	Oct 25, 2016	DS DP U-473			
	RE42461*PED	Apr 25, 2017				
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 002	5846966	Sep 21, 2013	DP U-473		PED	Dec 05, 2011
	5846966	Sep 21, 2013	DP U-593			
	5846966*PED	Mar 21, 2014				
	RE37721	Oct 25, 2016	DS DP U-473	Y		
	RE37721*PED	Apr 25, 2017		Y		
	RE42461	Oct 25, 2016	DS DP U-473			
	RE42461*PED	Apr 25, 2017				
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>						
N021687 003	5846966	Sep 21, 2013	DP U-473			
	5846966	Sep 21, 2013	DP U-593			
	5846966*PED	Mar 21, 2014				
	RE37721	Oct 25, 2016	DS DP U-473	Y		
	RE37721*PED	Apr 25, 2017		Y		
	RE42461	Oct 25, 2016	DS DP U-473			
	RE42461*PED	Apr 25, 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
EZETIMIBE; SIMVASTATIN - VYTORIN							
N021687 004	5846966	Sep	21, 2013	DP U-473		PED	Dec 05, 2011
	5846966	Sep	21, 2013	DP U-593			
	5846966*PED	Mar	21, 2014				
	RE37721	Oct	25, 2016	DS DP U-473	Y		
	RE37721*PED	Apr	25, 2017		Y		
	RE42461	Oct	25, 2016	DS DP U-473			
	RE42461*PED	Apr	25, 2017				
EZOGABINE - POTIGA							
N022345 001						NCE	Jun 10, 2016
EZOGABINE - POTIGA							
N022345 002						NCE	Jun 10, 2016
EZOGABINE - POTIGA							
N022345 003						NCE	Jun 10, 2016
EZOGABINE - POTIGA							
N022345 004						NCE	Jun 10, 2016
FAMCICLOVIR - FAMVIR							
N020363 001	5840763	Sep	01, 2015	U-96		M-98	Jan 31, 2014
	5840763*PED	Mar	01, 2016			M-54	Dec 24, 2012
	5866581	Oct	04, 2014	U-96		PED	Jun 24, 2013
	5866581*PED	Apr	04, 2015				
	5916893	Sep	01, 2015	U-96			
	5916893*PED	Mar	01, 2016				
	6124304	Oct	04, 2014	U-96			
	6124304*PED	Apr	04, 2015				
FAMCICLOVIR - FAMVIR							
N020363 002	5840763	Sep	01, 2015	U-96		M-98	Jan 31, 2014
	5840763*PED	Mar	01, 2016			M-54	Dec 24, 2012
	5866581	Oct	04, 2014	U-96		PED	Jun 24, 2013
	5866581*PED	Apr	04, 2015				
	5916893	Sep	01, 2015	U-96			
	5916893*PED	Mar	01, 2016				
	6124304	Oct	04, 2014	U-96			
	6124304*PED	Apr	04, 2015				
FAMCICLOVIR - FAMVIR							
N020363 003	5840763	Sep	01, 2015	U-96		M-98	Jan 31, 2014
	5840763*PED	Mar	01, 2016			M-54	Dec 24, 2012
	5866581	Oct	04, 2014	U-96		PED	Jun 24, 2013
	5866581*PED	Apr	04, 2015				
	5916893	Sep	01, 2015	U-96			
	5916893*PED	Mar	01, 2016				
	6124304	Oct	04, 2014	U-96			
	6124304*PED	Apr	04, 2015				
FAMOTIDINE - FLUXID							
N021712 001	6024981	Apr	09, 2018	DP			
	6221392	Apr	09, 2018	DP			
FAMOTIDINE - FLUXID							
N021712 002	6024981	Apr	09, 2018	DP			
	6221392	Apr	09, 2018	DP			
FAMOTIDINE - PEPCID AC							
N020325 001	5854267	Dec	29, 2015	U-267			
	5854267*PED	Jun	29, 2016	U-267			
FAMOTIDINE - PEPCID AC							
N020801 001	5667794	May	02, 2015			Y	
	5667794*PED	Nov	02, 2015				
	5854267	Dec	29, 2015	U-267			
	5854267*PED	Jun	29, 2016				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FAMOTIDINE - PEPCID AC							
N020801 002	6814978	Aug	26, 2021	DP			
	6814978*PED	Feb	26, 2022				
FAMOTIDINE - PEPCID AC (GELTAB)							
N020902 001	5854267	Dec	29, 2015	U-368			
	5854267*PED	Jun	29, 2016	U-368			
FAMOTIDINE; IBUPROFEN - DUEXIS							
N022519 001	8067033	Jul	18, 2026	DP		NC	
	8067451	Jul	18, 2026	DP	U-1196		Apr 23, 2014
FEBUXOSTAT - ULORIC							
N021856 001	5614520	Mar	25, 2014	DS	DP U-954	NCE	
	6225474	Jun	18, 2019	DS			Feb 13, 2014
	7361676	Mar	08, 2024	DP			
FEBUXOSTAT - ULORIC							
N021856 002	5614520	Mar	25, 2014	DS	DP U-954	NCE	
	6225474	Jun	18, 2019	DS			Feb 13, 2014
	7361676	Mar	08, 2024	DP			
FENOFRIBRATE - ANTARA (MICRONIZED)							
N021695 001	7101574	Aug	20, 2020	DS	DP		
	7863331	Aug	08, 2020		U-1107		
	7863331	Aug	08, 2020		U-1106		
FENOFRIBRATE - ANTARA (MICRONIZED)							
N021695 003	7101574	Aug	20, 2020	DS	DP		
	7863331	Aug	08, 2020		U-1106		
	7863331	Aug	08, 2020		U-1107		
FENOFRIBRATE - FENOGLIDE							
N022118 001	7658944	Dec	09, 2024	DP			
FENOFRIBRATE - FENOGLIDE							
N022118 002	7658944	Dec	09, 2024	DP			
FENOFRIBRATE - LIPOFEN							
N021612 001	5545628	Jan	10, 2015	DP	U-701		
FENOFRIBRATE - LIPOFEN							
N021612 002	5545628	Jan	10, 2015	DP	U-701		
FENOFRIBRATE - LIPOFEN							
N021612 003	5545628	Jan	10, 2015	DP	U-701		
FENOFRIBRATE - TRICOR							
N021203 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			
FENOFRIBRATE - TRICOR							
N021203 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			
FENOFRIBRATE - TRICOR							
N021656 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			
	7276249	Feb	21, 2023	DP			
	7320802	Feb	21, 2023		U-847		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FENOFIBRATE - TRICOR							
N021656 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		
FENOFIBRATE - TRIGLIDE							
N021350 001	6696084	Sep	11, 2021	DS	DP	U-680	
FENOFIBRATE - TRIGLIDE							
N021350 002	6696084	Sep	11, 2021	DS	DP	U-680	
FENOFIBRIC ACID - FIBRICOR							
N022418 001	7569612	Aug	20, 2027		U-1000		
	7741373	Aug	20, 2027		U-1059		
	7741374	Aug	20, 2027		U-1061		
	7741374	Aug	20, 2027		U-1060		
	7915247	Aug	20, 2027		U-1061		
	7915247	Aug	20, 2027		U-1059		
	7915247	Aug	20, 2027		U-1000		
FENOFIBRIC ACID - FIBRICOR							
N022418 002	7569612	Aug	20, 2027		U-1000		
	7741373	Aug	20, 2027		U-1059		
	7741374	Aug	20, 2027		U-1061		
	7741374	Aug	20, 2027		U-1060		
	7915247	Aug	20, 2027		U-1061		
	7915247	Aug	20, 2027		U-1059		
	7915247	Aug	20, 2027		U-1000		
FENTANYL CITRATE - ABSTRAL							
N022510 001	6759059	Sep	24, 2019	DP	U-767		
	6761910	Sep	24, 2019	DP	U-767		
	7910132	Sep	24, 2019	DP	U-767		
FENTANYL CITRATE - ABSTRAL							
N022510 002	6759059	Sep	24, 2019	DP	U-767		
	6761910	Sep	24, 2019	DP	U-767		
	7910132	Sep	24, 2019	DP	U-767		
FENTANYL CITRATE - ABSTRAL							
N022510 003	6759059	Sep	24, 2019	DP	U-767		
	6761910	Sep	24, 2019	DP	U-767		
	7910132	Sep	24, 2019	DP	U-767		
FENTANYL CITRATE - ABSTRAL							
N022510 004	6759059	Sep	24, 2019	DP	U-767		
	6761910	Sep	24, 2019	DP	U-767		
	7910132	Sep	24, 2019	DP	U-767		
FENTANYL CITRATE - ABSTRAL							
N022510 005	6759059	Sep	24, 2019	DP	U-767		
	6761910	Sep	24, 2019	DP	U-767		
	7910132	Sep	24, 2019	DP	U-767		
FENTANYL CITRATE - ABSTRAL							
N022510 006	6759059	Sep	24, 2019	DP	U-767		
	6761910	Sep	24, 2019	DP	U-767		
	7910132	Sep	24, 2019	DP	U-767		
FENTANYL CITRATE - FENTORA							
N021947 001	6200604	Mar	26, 2019		U-767		
	6974590	Mar	26, 2019		U-767		
	7862832	Jun	15, 2028	DP			
	7862833	Jun	15, 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
FENTANYL CITRATE - FENTORA						
N021947 002	6200604	Mar 26, 2019		U-767		
	6974590	Mar 26, 2019		U-767		
	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
FENTANYL CITRATE - FENTORA						
N021947 003	6200604	Mar 26, 2019		U-767		
	6974590	Mar 26, 2019		U-767		
	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
FENTANYL CITRATE - FENTORA						
N021947 004	6200604	Mar 26, 2019		U-767		
	6974590	Mar 26, 2019		U-767		
	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
FENTANYL CITRATE - FENTORA						
N021947 005	6200604	Mar 26, 2019		U-767		
	6974590	Mar 26, 2019		U-767		
	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
FENTANYL CITRATE - FENTORA						
N021947 006	6200604	Mar 26, 2019		U-767		
	6974590	Mar 26, 2019		U-767		
FENTANYL CITRATE - LAZANDA						
N022569 001	6432440	Apr 20, 2018	DP	U-1169	NDF	Jun 30, 2014
FENTANYL CITRATE - LAZANDA						
N022569 002	6432440	Apr 20, 2018	DP	U-1169	NDF	Jun 30, 2014
FENTANYL CITRATE - ONSOLIS						
N022266 001	6159498	Oct 18, 2016	DP		NP	Jul 16, 2012
	7579019	Jan 31, 2019	U-767			
FENTANYL CITRATE - ONSOLIS						
N022266 002	6159498	Oct 18, 2016	DP		NP	Jul 16, 2012
	7579019	Jan 31, 2019	U-767			
FENTANYL CITRATE - ONSOLIS						
N022266 003	6159498	Oct 18, 2016	DP		NP	Jul 16, 2012
	7579019	Jan 31, 2019	U-767			
FENTANYL CITRATE - ONSOLIS						
N022266 004	6159498	Oct 18, 2016	DP		NP	Jul 16, 2012
	7579019	Jan 31, 2019	U-767			
FENTANYL CITRATE - ONSOLIS						
N022266 005	6159498	Oct 18, 2016	DP		NP	Jul 16, 2012
	7579019	Jan 31, 2019	U-767			
FENTANYL HYDROCHLORIDE - IONSYS						
N021338 001	5697896	Dec 16, 2014	DP			
	5843014	Dec 01, 2015	DP			
	6169920	Jan 02, 2018	DP			
	6171294	Jun 05, 2015		U-736		
	6181963	Nov 02, 2019	DP			
	6195582	Jan 28, 2019	DP	U-736		
	6216033	Jun 05, 2015	DP			
	6317629	Jun 02, 2012	DP			
	6425892	Jun 05, 2015		U-736		
	6842640	Jun 02, 2015	DP			
	6881208	Apr 19, 2022		U-736		
	6975902	Apr 01, 2024	DP			
	7018370	Jun 05, 2015		U-736		
	7027859	Sep 26, 2014	DP			
	7302293	Jun 05, 2015	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>FERUMOXYTOL - FERAHEME</u>							
N022180 001	6599498	Mar	08, 2020	DS DP		NP	Jun 30, 2012
	7553479	Mar	08, 2020	DS DP			
	7871597	Mar	08, 2020	DS DP			
<u>FESOTERODINE FUMARATE - TOVIAZ</u>							
N022030 001	6858650	May	11, 2019	DS U-913		NCE	Oct 31, 2013
	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			
<u>FESOTERODINE FUMARATE - TOVIAZ</u>							
N022030 002	6858650	May	11, 2019	DS U-913		NCE	Oct 31, 2013
	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			
<u>FEXOFENADINE HYDROCHLORIDE - ALLEGRA</u>							
N020625 001	5578610	Nov	26, 2013	U-192			
	5578610*PED	May	26, 2014	U-192			
	5738872	Feb	28, 2015				
	5738872*PED	Aug	28, 2015				
	5855912	Feb	28, 2015				
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015				
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017	U-138			
	6037353*PED	Sep	14, 2017	U-138			
	6113942	Feb	28, 2015				
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012	U-138			
	6187791*PED	Nov	11, 2012	U-138			
	6399632	May	11, 2012	U-468			
	6399632*PED	Nov	11, 2012	U-468			
	7135571	May	18, 2014	DS			
	7135571*PED	Nov	18, 2014	DS			
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014	DS			
<u>FEXOFENADINE HYDROCHLORIDE - ALLEGRA</u>							
N021963 001	5578610	Nov	26, 2013	DS DP U-772			
	6037353	Mar	14, 2017	U-772			
	6187791	May	11, 2012	U-772			
	6399632	May	11, 2012	U-772			
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014	DS			
<u>FEXOFENADINE HYDROCHLORIDE - ALLEGRA ALLERGY</u>							
N020872 007	5578610	Nov	26, 2013	DS DP U-1160			
	5578610*PED	May	26, 2014	DP			
	5855912	Feb	28, 2015				
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015				
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017	U-1160			
	6037353*PED	Sep	14, 2017				
	6113942	Feb	28, 2015				
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012	U-1160			
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012	U-1160			
	6399632*PED	Nov	11, 2012				
	7135571	May	18, 2014	DS	U-1160		
	7135571*PED	Nov	18, 2014	DS			
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014	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
FEXOFENADINE HYDROCHLORIDE - ALLEGRA ALLERGY							
N020872 010	5578610	Nov	26, 2013	DS	DP	U-1160	
	5578610*PED	May	26, 2014				
	5855912	Feb	28, 2015		DP		
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015		DP		
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017			U-1160	
	6037353*PED	Sep	14, 2017				
	6113942	Feb	28, 2015		DP		
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012			U-1160	
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012			U-1160	
	6399632*PED	Nov	11, 2012				
	7135571	May	18, 2014	DS		U-1160	
	7135571*PED	Nov	18, 2014				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
FEXOFENADINE HYDROCHLORIDE - ALLEGRA HIVES							
N020872 008	5578610	Nov	26, 2013	DS	DP	U-1160	
	5578610*PED	May	26, 2014				
	5855912	Feb	28, 2015		DP		
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015		DP		
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017			U-1160	
	6037353*PED	Sep	14, 2017				
	6113942	Feb	28, 2015		DP		
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012			U-1160	
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012			U-1160	
	6399632*PED	Nov	11, 2012				
	7135571	May	18, 2014	DS		U-1160	
	7135571*PED	Nov	18, 2014				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
FEXOFENADINE HYDROCHLORIDE - ALLEGRA HIVES							
N020872 009	5578610	Nov	26, 2013	DS	DP	U-1160	
	5578610*PED	May	26, 2014				
	5855912	Feb	28, 2015		DP		
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015		DP		
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017			U-1160	
	6037353*PED	Sep	14, 2017				
	6113942	Feb	28, 2015		DP		
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012			U-1160	
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012			U-1160	
	6399632*PED	Nov	11, 2012				
	7135571	May	18, 2014	DS		U-1160	
	7135571*PED	Nov	18, 2014				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY							
N020872 005	5578610	Nov	26, 2013	DS DP U-1160			
	5578610*PED	May	26, 2014				
	5855912	Feb	28, 2015	DP			
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015	DP			
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017		U-1160		
	6037353*PED	Sep	14, 2017				
	6113942	Feb	28, 2015	DP			
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012		U-1160		
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012		U-1160		
	6399632*PED	Nov	11, 2012				
	7135571	May	18, 2014	DS		U-1160	
	7135571*PED	Nov	18, 2014				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY							
N021909 002	5578610	Nov	26, 2013	DS DP U-1158			
	5738872	Feb	28, 2015	DP			
	6037353	Mar	14, 2017		U-1158		
	6187791	May	11, 2012		U-1158		
	6399632	May	11, 2012		U-1158		
	7138524	May	18, 2014	DS			
FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY							
N201373 001	5578610	Nov	26, 2013	DS DP U-1157			
	5578610*PED	May	26, 2014				
	6037353	Mar	14, 2017		U-1157		
	6037353*PED	Sep	14, 2017				
	6187791	May	11, 2012		U-1157		
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012		U-1157		
	6399632*PED	Nov	11, 2012				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES							
N020872 006	5578610	Nov	26, 2013	DS DP U-1160			
	5578610*PED	May	26, 2014				
	5855912	Feb	28, 2015	DP			
	5855912*PED	Aug	28, 2015				
	5932247	Feb	28, 2015	DP			
	5932247*PED	Aug	28, 2015				
	6037353	Mar	14, 2017		U-1160		
	6037353*PED	Sep	14, 2017				
	6113942	Feb	28, 2015	DP			
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012		U-1160		
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012		U-1160		
	6399632*PED	Nov	11, 2012				
	7135571	May	18, 2014	DS		U-1160	
	7135571*PED	Nov	18, 2014				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES							
N021909 003	5578610	Nov	26, 2013	DS DP U-1158			
	5738872	Feb	28, 2015	DP			
	6037353	Mar	14, 2017		U-1158		
	6187791	May	11, 2012		U-1158		
	6399632	May	11, 2012		U-1158		
	7138524	May	18, 2014	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
<u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u>							
N201373 002	5578610	Nov	26, 2013	DS DP	U-1157		
	5578610*PED	May	26, 2014				
	6037353	Mar	14, 2017		U-1157		
	6037353*PED	Sep	14, 2017				
	6187791	May	11, 2012		U-1157		
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012		U-1157		
	6399632*PED	Nov	11, 2012				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
<u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION</u>							
N020786 002	5578610	Nov	26, 2013	DS DP	U-1159		
	5578610*PED	May	26, 2014				
	5855912	Feb	28, 2015		DP		
	5855912*PED	Aug	28, 2015				
	6037353	Mar	14, 2017		U-1159		
	6037353*PED	Sep	14, 2017		U-138		
	6039974	Jul	31, 2018		DP		
	6113942	Feb	28, 2015		DP		
	6113942*PED	Aug	28, 2015				
	6187791	May	11, 2012		U-1159		
	6187791*PED	Nov	11, 2012		U-138		
	6399632	May	11, 2012		U-1159		
	6399632*PED	Nov	11, 2012		U-468		
	7135571	May	18, 2014	DS	U-1159		
	7135571*PED	Nov	18, 2014				
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
<u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION</u>							
N021704 002	5578610	Nov	26, 2013	DS DP	U-1159		
	5578610*PED	May	26, 2014				
	6037353	Mar	14, 2017		U-1159		
	6037353*PED	Sep	14, 2017				
	6187791	May	11, 2012		U-1159		
	6187791*PED	Nov	11, 2012				
	6399632	May	11, 2012		U-1159		
	6399632*PED	Nov	11, 2012				
	6613357	Dec	25, 2020		DP U-1159		
	7138524	May	18, 2014	DS			
	7138524*PED	Nov	18, 2014				
	RE39069	May	29, 2018		DP		
<u>FIDAXOMICIN - DIFICID</u>							
N201699 001	7378508	Jul	31, 2027	DS DP		NCE	
	7863249	Jul	31, 2027	DS DP			May 27, 2016
	7906489	Mar	04, 2027		U-319		
<u>FINASTERIDE - PROPECIA</u>							
N020788 001	5547957	Oct	15, 2013		U-236		
	5571817	Nov	05, 2013		U-259		
	5886184	Nov	19, 2012				
<u>FINASTERIDE - PROSCAR</u>							
N020180 001	5886184	Nov	19, 2012	DS			
	5942519	Oct	23, 2018		U-280		
<u>FINGOLIMOD - GILENYA</u>							
N022527 001	5604229	Feb	18, 2014	DS	U-1086	M-106	
						NCE	Sep 21, 2015
<u>FLUDARABINE PHOSPHATE - OFORTA</u>							
N022273 001	7148207	Dec	20, 2022		DP U-944	NDF	
	7547776	Dec	10, 2018	DS		ODE	Dec 18, 2015
<u>FLUOCINOLONE ACETONIDE - RETISERT</u>							
N021737 001	6217895	Mar	22, 2019		DP U-708	ODE	
	6548078	Mar	22, 2019	DP	U-708		Apr 08, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN - TRI-LUMA</u>							
N021112 001	7915243	Mar	22, 2026	DP			
	7939516	May	04, 2025	DP			
<u>FLUOCINONIDE - VANOS</u>							
N021758 001	6765001	Dec	21, 2021	DP			
	7220424	Jan	07, 2023		U-861		
	7794738	Sep	11, 2022		U-1084		
<u>FLUOROURACIL - CARAC</u>							
N020985 001	6670335	Jun	02, 2021	DP	U-68		
<u>FLUOXETINE HYDROCHLORIDE - PROZAC</u>							
N018936 001	6960577	Nov	01, 2017		U-963		I-589 Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE - PROZAC</u>							
N018936 003	6960577	Nov	01, 2017		U-963		I-589 Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE - PROZAC</u>							
N018936 006	6960577	Nov	01, 2017		U-963		I-589 Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE - PROZAC WEEKLY</u>							
N021235 001	5910319	May	29, 2017		U-396		
	5985322	May	29, 2017		U-397		
	RE39030	May	29, 2017	DP	U-396		
	RE39030	May	29, 2017	DP	U-397		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>							
N021520 001	5229382*PED	Oct	23, 2011				I-593 Mar 19, 2012
	6960577	Nov	01, 2017		U-962		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>							
N021520 002	5229382*PED	Oct	23, 2011				I-593 Mar 19, 2012
	5945416	Mar	24, 2017	DS	DP		
	6960577	Nov	01, 2017		U-962	Y	
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>							
N021520 003	5229382*PED	Oct	23, 2011				I-593 Mar 19, 2012
	5945416	Mar	24, 2017	DS	DP		
	6960577	Nov	01, 2017		U-962	Y	
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>							
N021520 004	5229382*PED	Oct	23, 2011				I-593 Mar 19, 2012
	5945416	Mar	24, 2017	DS	DP		
	6960577	Nov	01, 2017		U-962	Y	
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>							
N021520 005	5229382*PED	Oct	23, 2011				I-593 Mar 19, 2012
	5945416	Mar	24, 2017	DS	DP		
	6960577	Nov	01, 2017		U-962	Y	
<u>FLUTICASONE FUROATE - VERAMYST</u>							
N022051 001	6858596	Aug	03, 2021		DP	U-808	
	7101866	Aug	03, 2021	DS	DP	U-808	
	7541350	Aug	03, 2021		DP	U-988	
<u>FLUTICASONE PROPIONATE - CUTIVATE</u>							
N021152 001	7300669	Oct	20, 2019		DP	U-835	
<u>FLUTICASONE PROPIONATE - FLOVENT DISKUS 100</u>							
N020833 002	5590645*PED	Sep	01, 2011				
	5860419*PED	Sep	01, 2011				
	5873360	Feb	23, 2016		DP		
	5873360*PED	Aug	23, 2016				
	6032666*PED	Sep	01, 2011				
	6378519*PED	Sep	01, 2011				
	6536427*PED	Sep	01, 2011				
	6792945*PED	Sep	01, 2011				
	7225808*PED	Sep	01, 2011				
	7389775*PED	Sep	01, 2011				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE - FLOVENT DISKUS 250						
N020833 003	5590645*PED	Sep 01, 2011				
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016		DP		
	5873360*PED	Aug 23, 2016				
	6032666*PED	Sep 01, 2011				
	6378519*PED	Sep 01, 2011				
	6536427*PED	Sep 01, 2011				
	6792945*PED	Sep 01, 2011				
	7225808*PED	Sep 01, 2011				
	7389775*PED	Sep 01, 2011				
FLUTICASONE PROPIONATE - FLOVENT DISKUS 50						
N020833 001	5590645*PED	Sep 01, 2011				
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016		DP		
	5873360*PED	Aug 23, 2016				
	6032666*PED	Sep 01, 2011				
	6378519*PED	Sep 01, 2011				
	6536427*PED	Sep 01, 2011				
	6792945*PED	Sep 01, 2011				
	7225808*PED	Sep 01, 2011				
	7389775*PED	Sep 01, 2011				
FLUTICASONE PROPIONATE - FLOVENT HFA						
N021433 001	5658549	Aug 19, 2014	DP	U-710		
	5658549*PED	Feb 19, 2015				
	5674472	Oct 07, 2014	DP			
	5674472*PED	Apr 07, 2015				
	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013				
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018				
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6510969	Dec 23, 2017	DP			
	6510969*PED	Jun 23, 2018				
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	6743413	Jun 01, 2021		U-581		
	6743413*PED	Dec 01, 2021				
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018				
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018				
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018				
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2019				
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018				
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019				
	7500444	Jan 04, 2025				
	7500444*PED	Jul 04, 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
FLUTICASONE PROPIONATE - FLOVENT HFA						
N021433 002	5658549	Aug 19, 2014	DP U-710			
	5658549*PED	Feb 19, 2015				
	5674472	Oct 07, 2014	DP			
	5674472*PED	Apr 07, 2015				
	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013				
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018				
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6510969	Dec 23, 2017	DP			
	6510969*PED	Jun 23, 2018				
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	6743413	Jun 01, 2021	U-581			
	6743413*PED	Dec 01, 2021				
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018				
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018				
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018				
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2019				
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018				
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019				
	7500444	Jan 04, 2025	DP			
	7500444*PED	Jul 04, 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
FLUTICASONE PROPIONATE - FLOVENT HFA						
N021433 003	5658549	Aug 19, 2014	DP U-710			
	5658549*PED	Feb 19, 2015		U-710		
	5674472	Oct 07, 2014	DP			
	5674472*PED	Apr 07, 2015				
	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013				
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018	DP			
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6510969	Dec 23, 2017	DP			
	6510969*PED	Jun 23, 2018	DP			
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	6743413	Jun 01, 2021	U-581			
	6743413*PED	Dec 01, 2021	U-581			
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018				
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018				
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018				
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2019				
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018				
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019				
	7500444	Jan 04, 2025	DP			
	7500444*PED	Jul 04, 2025				
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50						
N021077 001	5590645*PED	Sep 01, 2011			M-84	Mar 31, 2012
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666*PED	Sep 01, 2011				
	6378519*PED	Sep 01, 2011				
	6536427*PED	Sep 01, 2011				
	6792945*PED	Sep 01, 2011				
	7225808*PED	Sep 01, 2011				
	7389775*PED	Sep 01, 2011				
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50						
N021077 002	5590645*PED	Sep 01, 2011			M-84	Mar 31, 2012
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666*PED	Sep 01, 2011				
	6378519*PED	Sep 01, 2011				
	6536427*PED	Sep 01, 2011				
	6792945*PED	Sep 01, 2011				
	7225808*PED	Sep 01, 2011				
	7389775*PED	Sep 01, 2011				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50						
N021077 003	5590645*PED	Sep 01, 2011			M-84	Mar 31, 2012
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666*PED	Sep 01, 2011				
	6378519*PED	Sep 01, 2011				
	6536427*PED	Sep 01, 2011				
	6792945*PED	Sep 01, 2011				
	7225808*PED	Sep 01, 2011				
	7389775*PED	Sep 01, 2011				
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA						
N021254 001	5658549	Aug 19, 2014	DP	U-738		
	5658549*PED	Feb 19, 2015		U-738		
	5674472	Oct 07, 2014	DP			
	5674472*PED	Apr 07, 2015				
	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013				
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018				
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6510969	Dec 23, 2017				
	6510969*PED	Jun 23, 2018				
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	6743413	Jun 01, 2021		U-841		
	6743413*PED	Dec 01, 2021		U-841		
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018				
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018				
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018				
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2018				
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018				
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019				
	7500444	Jan 04, 2025	DP			
	7500444*PED	Jul 04, 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
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA						
N021254 002	5658549	Aug 19, 2014	DP U-738			
	5658549*PED	Feb 19, 2015	DP U-738			
	5674472	Oct 07, 2014	DP			
	5674472*PED	Apr 07, 2015	DP			
	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013	DP			
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018	DP			
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6510969	Dec 23, 2017	DP			
	6510969*PED	Jun 23, 2018	DP			
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	6743413	Jun 01, 2021	U-841			
	6743413*PED	Dec 01, 2021	U-841			
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018				
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018				
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018				
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2018				
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018				
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019				
	7500444	Jan 04, 2025	DP			
	7500444*PED	Jul 04, 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
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N021254 003	5658549	Aug 19, 2014	DP U-738			
	5658549*PED	Feb 19, 2015	DP U-738			
	5674472	Oct 07, 2014	DP			
	5674472*PED	Apr 07, 2015	DP			
	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018	DP			
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018	DP			
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013	DP			
	6315173	Dec 23, 2017	DP			
	6315173*PED	Jun 23, 2018	DP			
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018	DP			
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018	DP			
	6510969	Dec 23, 2017	DP			
	6510969*PED	Jun 23, 2018	DP			
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015	DP			
	6743413	Jun 01, 2021	U-841			
	6743413*PED	Dec 01, 2021	U-841			
	6938796	Jan 16, 2018	DP			
	6938796*PED	Jul 16, 2018	DP			
	6966467	Dec 23, 2017	DP			
	6966467*PED	Jun 23, 2018	DP			
	6997349	Jan 16, 2018	DP			
	6997349*PED	Jul 16, 2018	DP			
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2018	DP			
	7143908	Jan 16, 2018	DP			
	7143908*PED	Jul 16, 2018	DP			
	7350676	Aug 24, 2018	DP			
	7350676*PED	Feb 24, 2019	DP			
	7500444	Jan 04, 2025	DP			
	7500444*PED	Jul 04, 2025	DP			
<u>FLUVASTATIN SODIUM - LESCOL</u>						
N020261 001	5354772	Oct 11, 2011	U-109			
	5354772	Oct 11, 2011	U-413			
	5354772*PED	Apr 11, 2012				
	5356896	Dec 12, 2011				
	5356896*PED	Jun 12, 2012				
<u>FLUVASTATIN SODIUM - LESCOL XL</u>						
N020261 002	5354772	Oct 11, 2011	U-109			
	5354772	Oct 11, 2011	U-413			
	5354772*PED	Apr 11, 2012				
	5356896	Dec 12, 2011				
	5356896*PED	Jun 12, 2012				
<u>FLUVOXAMINE MALEATE - LUVOX CR</u>						
N022033 001	7465462	May 10, 2020	DP U-929			
<u>FLUVOXAMINE MALEATE - LUVOX CR</u>						
N022033 002	7465462	May 10, 2020	DP U-929			
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM</u>						
N020582 001	5767251	Jun 16, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>FOLLITROPIN ALFA/BETA - FOLLISTIM</u>						
	N020582 002	5767251	Jun 16, 2015			
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u>						
	N021211 001	5767251	Jun 16, 2015	DS	D-133	Aug 22, 2014
	5929028	Jan 14, 2018		DP U-567	I-306	Jun 28, 2013
	7446090	Aug 23, 2019		DP		
	7563763	Aug 23, 2019		U-1183		
	7563763	Aug 23, 2019		U-993		
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u>						
	N021211 002	5767251	Jun 16, 2015	DS	D-133	Aug 22, 2014
	5929028	Jan 14, 2018		DP U-567	I-306	Jun 28, 2013
	7446090	Aug 23, 2019		DP		
	7563763	Aug 23, 2019		U-1183		
	7563763	Aug 23, 2019		U-993		
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u>						
	N021211 003	5767251	Jun 16, 2015	DS	D-133	Aug 22, 2014
	5929028	Jan 14, 2018		DP U-567		
	7446090	Aug 23, 2019		DP		
	7563763	Aug 23, 2019		U-1183		
	7563763	Aug 23, 2019		U-993		
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u>						
	N021211 004	5767251	Jun 16, 2015	DS	D-133	Aug 22, 2014
	5929028	Jan 14, 2018		DP U-567	I-306	Jun 28, 2013
	7446090	Aug 23, 2019		DP		
	7563763	Aug 23, 2019		U-993		
	7563763	Aug 23, 2019		U-1183		
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u>						
	N021273 001				I-306	Jun 28, 2013
<u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u>						
	N021273 002				I-306	Jun 28, 2013
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N020378 001	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N020378 002	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N020378 003	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N020378 004	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015		DS		
	7563763	Aug 23, 2019		DP		
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N020378 005	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015		DS		
	7563763	Aug 23, 2019		DP		
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N021765 001	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
	N021765 003	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF</u>						
	N021765 002	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015		DS		
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF PEN</u>						
	N021684 001	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015		DS		
	7446090	Aug 23, 2019		DP		
	7741268	Apr 02, 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
FOLLITROPIN ALFA/BETA - GONAL-F RFF PEN							
N021684 002	5767067	Jun	16, 2015	DS			
	5767251	Jun	16, 2015	DS			
	7446090	Aug	23, 2019	DP			
	7741268	Apr	02, 2024	DP			
FOLLITROPIN ALFA/BETA - GONAL-F RFF PEN							
N021684 003	5767067	Jun	16, 2015	DS			
	5767251	Jun	16, 2015	DS			
	7446090	Aug	23, 2019	DP			
	7741268	Apr	02, 2024	DP			
FOMEPIZOLE - ANTIZOL							
N020696 001	7553863	Jun	30, 2027	DS	DP		
FOMIVIRSEN SODIUM - VITRAVENE PRESERVATIVE FREE							
N020961 001	5442049	Aug	15, 2012				
	5595978	Aug	15, 2012			U-522	
FORMOTEROL FUMARATE - FORADIL							
N020831 001	6488027	Mar	08, 2019				
	6887459	Nov	28, 2020			U-762	
FORMOTEROL FUMARATE - PERFOROMIST							
N022007 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		
FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA							
N022518 001	5889015	Jan	27, 2014		U-1068		
	5889015*PED	Jul	27, 2014				
	6057307	Jan	27, 2014	DP	U-1068		
	6057307*PED	Jul	27, 2014				
	6068832	Aug	27, 2017	DP	U-1068		
	6677323	Jan	27, 2014		U-1068		
	7067502	May	21, 2020	DP	U-1068		
	7566705	May	08, 2020	DP	U-1068		
FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA							
N022518 002	5889015	Jan	27, 2014		U-1068		
	5889015*PED	Jul	27, 2014				
	6057307	Jan	27, 2014	DP	U-1068		
	6057307*PED	Jul	27, 2014				
	6068832	Aug	27, 2017	DP	U-1068		
	6677323	Jan	27, 2014		U-1068		
	7067502	May	21, 2020	DP	U-1068		
	7566705	May	08, 2020	DP	U-1068		
FOSAMPRENAVIR CALCIUM - LEXIVA							
N021548 001	6436989	Dec	24, 2017	DS	DP	U-257	
	6436989*PED	Jun	24, 2018				
	6514953	Jul	15, 2019	DS	DP	U-257	
	6514953*PED	Jan	15, 2020				
FOSAMPRENAVIR CALCIUM - LEXIVA							
N022116 001	6436989	Dec	24, 2017	DS	DP	U-257	
	6436989*PED	Jun	24, 2018				
FOSAPREPITANT DIMEGLUMINE - EMEND							
N022023 001	5512570	Mar	04, 2014		U-850		
	5538982	Jul	23, 2013		U-850		
	5691336	Mar	04, 2019	DS	DP		
	5716942	Feb	10, 2015		U-850		
	7214692	Sep	18, 2012		U-850		
FOSAPREPITANT DIMEGLUMINE - EMEND							
N022023 002	5691336	Mar	04, 2019	DS	DP		
						D-128	Nov 12, 2013
						NCE	Jan 25, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>FOSPROPOFOL DISODIUM - LUSEDRA</u>							
N022244 001	6204257	Aug	07, 2018	DS	DP	U-945	
	6872838	Aug	07, 2018	DS			NCE Dec 12, 2013
<u>FROVATRIPTAN SUCCINATE - FROVA</u>							
N021006 001	5464864	Nov	07, 2015			U-436	
	5616603	Apr	01, 2014			U-436	
	5637611	Jun	10, 2014			U-436	
	5827871	Oct	27, 2015			U-436	
	5962501	Dec	16, 2013			U-436	
<u>FULVESTRANT - FASLODEX</u>							
N021344 001	6774122	Jan	09, 2021			U-596	
	6774122*PED	Jul	09, 2021				D-126 Sep 09, 2013
	7456160	Jan	09, 2021			U-596	M-103 May 17, 2014
	7456160*PED	Jul	09, 2021				PED Nov 17, 2014
							PED Mar 09, 2014
<u>GABAPENTIN - GABAPENTIN</u>							
A078974 001							PC Aug 22, 2011
<u>GABAPENTIN - GRALISE</u>							
N022544 001	6340475	Sep	19, 2016	DP			
	6488962	Jun	20, 2020	DP			NP Jan 28, 2014
	6635280	Sep	19, 2016	DP			
	6723340	Oct	25, 2021	DP			
	7438927	Feb	26, 2024			U-1114	
	7731989	Oct	25, 2022	DP			
<u>GABAPENTIN - GRALISE</u>							
N022544 002	6340475	Sep	19, 2016	DP			
	6488962	Jun	20, 2020	DP			NP Jan 28, 2014
	6635280	Sep	19, 2016	DP			
	6723340	Oct	25, 2021	DP			
	7438927	Feb	26, 2024			U-1114	
	7731989	Oct	25, 2022	DP			
<u>GABAPENTIN - NEURONTIN</u>							
N020235 001	6054482	Apr	25, 2017				
	6054482*PED	Oct	25, 2017				
<u>GABAPENTIN - NEURONTIN</u>							
N020235 002	6054482	Apr	25, 2017				
	6054482*PED	Oct	25, 2017				
<u>GABAPENTIN - NEURONTIN</u>							
N020235 003	6054482	Apr	25, 2017				
	6054482*PED	Oct	25, 2017				
<u>GABAPENTIN - NEURONTIN</u>							
N020882 001	6054482	Apr	25, 2017				
	6054482*PED	Oct	25, 2017				
<u>GABAPENTIN - NEURONTIN</u>							
N020882 002	6054482	Apr	25, 2017				
	6054482*PED	Oct	25, 2017				
<u>GABAPENTIN - NEURONTIN</u>							
N021129 001	6054482	Apr	25, 2017				
	6054482*PED	Oct	25, 2017				
	7256216	May	28, 2022				
	7256216*PED	Nov	28, 2022	DP			
<u>GABAPENTIN ENACARBIL - HORIZANT</u>							
N022399 001	6818787	Nov	06, 2022	DS	DP		
	8026279	Nov	10, 2026	DS	DP		NCE Apr 06, 2016
	8048917	Nov	06, 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
GABAPENTIN ENACARBIL - HORIZANT							
N022399 002	6818787	Nov	06, 2022	DS DP		NCE	Apr 06, 2016
	8026279	Nov	10, 2026	DS DP			
	8048917	Nov	06, 2022	DS DP			
GADOBENATE DIMEGLUMINE - MULTIHANCE							
N021357 001	4916246	Apr	10, 2012	DS		NPP	Mar 17, 2013
GADOBENATE DIMEGLUMINE - MULTIHANCE							
N021357 002	4916246	Apr	10, 2012	DS		NPP	Mar 17, 2013
GADOBENATE DIMEGLUMINE - MULTIHANCE							
N021357 003	4916246	Apr	10, 2012	DS		NPP	Mar 17, 2013
GADOBENATE DIMEGLUMINE - MULTIHANCE							
N021357 004	4916246	Apr	10, 2012	DS		NPP	Mar 17, 2013
GADOBENATE DIMEGLUMINE - MULTIHANCE MULTIPACK							
N021358 001	4916246	Apr	10, 2012	DS			
GADOBENATE DIMEGLUMINE - MULTIHANCE MULTIPACK							
N021358 002	4916246	Apr	10, 2012	DS			
GADOBUTROL - GADAVIST							
N201277 001	5980864	Nov	09, 2016	DS DP U-1119		NCE	Mar 14, 2016
GADOBUTROL - GADAVIST							
N201277 002	5980864	Nov	09, 2016	DS DP U-1119		NCE	Mar 14, 2016
GADOBUTROL - GADAVIST							
N201277 003	5980864	Nov	09, 2016	DS DP U-1119		NCE	Mar 14, 2016
GADOBUTROL - GADAVIST							
N201277 004	5980864	Nov	09, 2016	DS DP U-1119		NCE	Mar 14, 2016
GADOBUTROL - GADAVIST							
N201277 005	5980864	Nov	09, 2016	DS DP U-1119		NCE	Mar 14, 2016
GADODIAMIDE - OMNISCAN							
N020123 001	5362475	Nov	08, 2011	DS		NCE	Dec 22, 2013
	5560903	Oct	01, 2013	DP			
GADODIAMIDE - OMNISCAN							
N022066 001	5560903	Oct	01, 2013	DP			
GADODIAMIDE - OMNISCAN							
N022066 002	5362475	Nov	08, 2011	DS		NCE	Dec 22, 2013
	5560903	Oct	01, 2013	DP			
GADOFOSVESET TRISODIUM - ABLAVAR							
N021711 001	5362475	Nov	08, 2011	DS		NCE	Dec 22, 2013
	6676929	May	26, 2015	DP			
	7011815	Feb	01, 2015		U-1112		
	7060250	May	26, 2015	DS			
	7229606	May	26, 2015		U-1112		
	8017105	May	26, 2015	DS			
GADOFOSVESET TRISODIUM - ABLAVAR							
N021711 002	5362475	Nov	08, 2011	DS		NCE	Dec 22, 2013
	6676929	May	26, 2015	DP			
	7011815	Feb	01, 2015		U-1112		
	7060250	May	26, 2015	DS			
	7229606	May	26, 2015		U-1112		
	8017105	May	26, 2015	DS			
GADOPENTETATE DIMEGLUMINE - MAGNEVIST							
N019596 001	5362475	Nov	08, 2011				
GADOPENTETATE DIMEGLUMINE - MAGNEVIST							
N021037 001	5362475	Nov	08, 2011				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
GADOTERIDOL - PROHANCE						
N020131 001	5474756	Dec 12, 2012		U-480		
	5846519	Dec 08, 2015				
	6143274	Dec 12, 2012		U-480		
GADOTERIDOL - PROHANCE MULTIPACK						
N021489 001	5474756	Dec 12, 2012		U-536		
	5846519	Dec 08, 2015				
	6143274	Dec 12, 2012		U-536		
GADOXETATE DISODIUM - EOViST						
N022090 001					NCE	Jul 03, 2013
GALANTAMINE HYDROBROMIDE - RAZADYNE						
N021169 001	6099863	Jun 06, 2017				
	6358527	Jun 06, 2017		DP U-322		
GALANTAMINE HYDROBROMIDE - RAZADYNE						
N021169 002	6099863	Jun 06, 2017				
	6358527	Jun 06, 2017		DP U-322		
GALANTAMINE HYDROBROMIDE - RAZADYNE						
N021169 003	6099863	Jun 06, 2017				
	6358527	Jun 06, 2017		DP U-322		
GALANTAMINE HYDROBROMIDE - RAZADYNE ER						
N021615 001	7160559	Dec 20, 2019		DP		
GALANTAMINE HYDROBROMIDE - RAZADYNE ER						
N021615 002	7160559	Dec 20, 2019		DP		
GALANTAMINE HYDROBROMIDE - RAZADYNE ER						
N021615 003	7160559	Dec 20, 2019		DP		
GANCICLOVIR - VITRASERT						
N020569 001	5378475	Jan 03, 2012				
GANCICLOVIR - ZIRGAN						
N022211 001					NDF	Sep 15, 2012
					ODE	Sep 15, 2016
GANIRELIX ACETATE - GANIRELIX ACETATE INJECTION						
N021057 001	4801577	Feb 05, 2012	DS DP			
	5767082	Jun 16, 2015				
GATIFLOXACIN - ZYMAR						
N021493 001	5880283	Dec 05, 2015				
	5880283*PED	Jun 05, 2016				
	6333045	Aug 20, 2019				
	6333045*PED	Feb 20, 2020	DP			
GATIFLOXACIN - ZYMAXID						
N022548 001	5880283	Dec 05, 2015	DS		NP	May 18, 2013
	5880283*PED	Jun 05, 2016				
	6333045	Aug 20, 2019	DS DP			
	6333045*PED	Feb 20, 2020				
GEFITINIB - IRESSA						
N021399 001	5457105	Jan 19, 2013				
	5616582	Jan 19, 2013				
	5770599	May 05, 2017	DS DP U-881			
GEMCITABINE HYDROCHLORIDE - GEMCITABINE HYDROCHLORIDE						
A077983 001					PC	Jul 24, 2011
GEMCITABINE HYDROCHLORIDE - GEMCITABINE HYDROCHLORIDE						
A077983 002					PC	Jul 24, 2011
GEMCITABINE HYDROCHLORIDE - GEMZAR						
N020509 001	5464826	Nov 07, 2012	U-146			
	5464826*PED	May 07, 2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
GEMCITABINE HYDROCHLORIDE - GEMZAR						
N020509 002	5464826	Nov 07, 2012		U-146		
	5464826*PED	May 07, 2013				
GEMIFLOXACIN MESYLATE - FACTIVE						
N021158 001	5633262	Jun 15, 2015				
	5776944	Apr 04, 2017	DS DP			
	5962468	Jun 15, 2015		U-282		
	6262071	Sep 21, 2019		U-513		
	6331550	Sep 21, 2019		U-511		
	6340689	Sep 14, 2019		U-512		
	6455540	Sep 21, 2019		U-511		
	6723734	Mar 20, 2018	DS DP			
	6803376	Sep 21, 2019	DS DP	U-608		
	6803376	Sep 21, 2019	DS DP	U-609		
GEMTUZUMAB OZOGAMICIN - MYLOTARG						
N021174 001	5585089	Dec 17, 2013				
	5606040	Feb 25, 2014				
	5693762	Dec 02, 2014				
	5739116	Apr 14, 2015				
	5767285	Jun 16, 2015				
	5773001	Jun 30, 2015		U-320		
GLATIRAMER ACETATE - COPAXONE						
N020622 001	5981589	May 24, 2014			I-594	Feb 27, 2012
	6054430	May 24, 2014				
	6342476	May 24, 2014		U-441		
	6362161	May 24, 2014		U-441		
	6620847	May 24, 2014	DS			
	6939539	May 24, 2014	DS			
	7199098	May 24, 2014	DS			
GLATIRAMER ACETATE - COPAXONE						
N020622 002	5981589	May 24, 2014			I-594	Feb 27, 2012
	6054430	May 24, 2014				
	6342476	May 24, 2014		U-441		
	6362161	May 24, 2014		U-441		
	6620847	May 24, 2014	DS			
	6939539	May 24, 2014	DS			
	7199098	May 24, 2014	DS			
GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT						
N021925 001	6150383	Jun 19, 2016		U-753		
	6211205	Jun 19, 2016		U-753		
	6303640	Aug 09, 2016		U-753		
	6329404	Jun 19, 2016	DP	U-753		
	7538125	Jun 19, 2016		DP		
	7700128	Jan 30, 2027		DP		
	8071130	Jun 08, 2028		DP		
GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT						
N021925 002	6150383	Jun 19, 2016		U-753		
	6211205	Jun 19, 2016		U-753		
	6303640	Aug 09, 2016		U-753		
	6329404	Jun 19, 2016	DP	U-753		
	7538125	Jun 19, 2016		DP		
	7700128	Jan 30, 2027		DP		
	8071130	Jun 08, 2028		DP		
GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL						
N021700 001	5002953	Sep 17, 2011	DS DP	U-690		
	5002953	Sep 17, 2011	DS DP	U-781		
	5002953*PED	Mar 17, 2012		U-781		
	5741803	Apr 21, 2015	DS DP	U-781		
	5741803	Apr 21, 2015	DS DP	U-690		
	5741803*PED	Oct 21, 2015				
	7358366	Apr 19, 2020	DS			
	7358366*PED	Oct 19, 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>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>							
N021700 002	5002953	Sep	17, 2011	DS	DP U-690		
	5002953	Sep	17, 2011	DS	DP U-781		
	5002953*PED	Mar	17, 2012				
	5741803	Apr	21, 2015	DS	DP U-690		
	5741803	Apr	21, 2015	DS	DP U-781		
	5741803*PED	Oct	21, 2015				
	7358366	Apr	19, 2020	DS			
	7358366*PED	Oct	19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>							
N021700 003	5002953	Sep	17, 2011	DS	DP U-690		
	5002953	Sep	17, 2011	DS	DP U-781		
	5002953*PED	Mar	17, 2012				
	5741803	Apr	21, 2015	DS	DP U-690		
	5741803	Apr	21, 2015	DS	DP U-781		
	5741803*PED	Oct	21, 2015				
	7358366	Apr	19, 2020	DS			
	7358366*PED	Oct	19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>							
N021700 004	5002953	Sep	17, 2011	DS	DP U-840		
	5002953*PED	Mar	17, 2012				
	7358366	Apr	19, 2020	DS			
	7358366*PED	Oct	19, 2020				
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>							
N021700 005	5002953	Sep	17, 2011	DS	DP U-840		
	5002953*PED	Mar	17, 2012				
	7358366	Apr	19, 2020	DS			
	7358366*PED	Oct	19, 2020				
<u>GLIPIZIDE - GLUCOTROL XL</u>							
N020329 001	5591454	Jan	07, 2014			U-150	
<u>GLIPIZIDE - GLUCOTROL XL</u>							
N020329 002	5591454	Jan	07, 2014			U-150	
<u>GLIPIZIDE - GLUCOTROL XL</u>							
N020329 003	5591454	Jan	07, 2014			U-111	
<u>GLUTAMINE - NUTRESTORE</u>							
N021667 001	5288703	Oct	07, 2011			U-898	
<u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u>							
N021178 001	6303146	Jul	14, 2019			U-412	
	6303146*PED	Jan	14, 2020			U-412	
<u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u>							
N021178 002	6303146	Jul	14, 2019			U-412	
	6303146*PED	Jan	14, 2020			U-412	
<u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u>							
N021178 003	6303146	Jul	14, 2019			U-412	
	6303146*PED	Jan	14, 2020			U-412	
<u>GLYCOPYRROLATE - CUVPOSA</u>							
N022571 001	7638552	Aug	20, 2023			U-1076	
	7816396	Aug	20, 2023			U-1076	
<u>GLYCOPYRROLATE - ROBINUL</u>							
N012827 001	7091236	Apr	24, 2024			U-877	
<u>GLYCOPYRROLATE - ROBINUL FORTE</u>							
N012827 002	7091236	Apr	24, 2024			U-877	
<u>GOSERELIN ACETATE - ZOLADEX</u>							
N019726 001	7118552	Apr	13, 2022			DP	
	7220247	Apr	09, 2022			DP	
	7500964	Feb	26, 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
<u>GOSERELIN ACETATE - ZOLADEX</u>							
N020578 001	7118552	Apr	13, 2022	DP			
	7220247	Apr	09, 2022	DP			
	7500964	Feb	26, 2021	DP			
<u>GRANISETRON - SANCUSO</u>							
N022198 001	7608282	Oct	22, 2024	DP	U-1011	NE	Sep 12, 2011
						NDF	Sep 12, 2011
<u>GRANISETRON HYDROCHLORIDE - KYTRIL</u>							
N020239 001	5952340	Sep	14, 2016	U-519		M-102	Apr 29, 2014
	6294548	May	04, 2019				
<u>GRANISETRON HYDROCHLORIDE - KYTRIL</u>							
N020239 002	5952340	Sep	14, 2016	U-519		M-102	Apr 29, 2014
	6294548	May	04, 2019				
<u>GRANISETRON HYDROCHLORIDE - KYTRIL</u>							
N020239 003						M-102	Apr 29, 2014
<u>GRANISETRON HYDROCHLORIDE - KYTRIL</u>							
N020239 004	5952340	Sep	14, 2016	U-519		M-102	Apr 29, 2014
	6294548	May	04, 2019				
<u>GREPAFLOXACIN HYDROCHLORIDE - RAXAR</u>							
N020695 001	5563138	Oct	08, 2013				
<u>GUAIFENESIN - MUCINEX</u>							
N021282 001	6372252	Apr	28, 2020	U-489			
	6955821	Apr	28, 2020	DP	U-489		
	7838032	Apr	28, 2020	DP			
<u>GUAIFENESIN - MUCINEX</u>							
N021282 002	6372252	Apr	28, 2020	U-489			
	6955821	Apr	28, 2020	DP	U-489		
	7838032	Apr	28, 2020	DP			
<u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u>							
N021585 001	6372252	Apr	28, 2020	DP			
	6955821	Apr	28, 2020	DP	U-686		
	7838032	Apr	28, 2020	DP			
<u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u>							
N021585 002	6372252	Apr	28, 2020	DP			
	6955821	Apr	28, 2020	DP	U-686		
	7838032	Apr	28, 2020	DP			
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>							
N022037 001	5854290	Sep	21, 2015	U-494		I-635	Feb 25, 2014
	6287599	Dec	20, 2020	DP		NP	Sep 02, 2012
	6811794	Jul	04, 2022	DP	U-494		
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>							
N022037 002	5854290	Sep	21, 2015	U-494		I-635	Feb 25, 2014
	6287599	Dec	20, 2020	DP		NP	Sep 02, 2012
	6811794	Jul	04, 2022	DP	U-494		
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>							
N022037 003	5854290	Sep	21, 2015	U-494		I-635	Feb 25, 2014
	6287599	Dec	20, 2020	DP		NP	Sep 02, 2012
	6811794	Jul	04, 2022	DP	U-494		
<u>GUANFACINE HYDROCHLORIDE - INTUNIV</u>							
N022037 004	5854290	Sep	21, 2015	U-494		I-635	Feb 25, 2014
	6287599	Dec	20, 2020	DP		NP	Sep 02, 2012
	6811794	Jul	04, 2022	DP	U-494		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u>						
N022555 001	6034267	Mar 08, 2016		U-1087		NP May 28, 2013
	7247655	Mar 08, 2016		DP U-1087		
	7348361	Apr 22, 2019		DP U-1087		
	7530461	Jan 11, 2017		DP U-1087		
<u>HISTRELIN ACETATE - SUPPRELIN LA</u>						
N022058 001	8062652	Jun 16, 2026		U-1197		ODE May 03, 2014
<u>HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT</u>						
N021859 001	7767429	Sep 23, 2027	DS DP			
<u>HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE - BIDIL</u>						
N020727 001	6465463	Sep 08, 2020		U-71		
	6784177	Sep 08, 2020		U-71		
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN - AVALIDE</u>						
N020758 001	5270317	Sep 30, 2011				
	5270317*PED	Mar 30, 2012				
	5994348	Jun 07, 2015				
	5994348*PED	Dec 07, 2015				
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN - AVALIDE</u>						
N020758 002	5270317	Sep 30, 2011				
	5270317*PED	Mar 30, 2012				
	5994348	Jun 07, 2015				
	5994348*PED	Dec 07, 2015				
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN - AVALIDE</u>						
N020758 003	5270317	Sep 30, 2011	DS DP			
	5270317*PED	Mar 30, 2012				
	5994348	Jun 07, 2015	DP			
	5994348*PED	Dec 07, 2015				
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - BENICAR HCT</u>						
N021532 002	5616599	Apr 25, 2016	DS DP	U-500		
	5616599*PED	Oct 25, 2016				
	6878703	Nov 19, 2021		U-3		Y
	6878703*PED	May 19, 2022				
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - BENICAR HCT</u>						
N021532 003	5616599	Apr 25, 2016	DS DP	U-500		
	5616599*PED	Oct 25, 2016				
	6878703	Nov 19, 2021		U-3		Y
	6878703*PED	May 19, 2022				
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - BENICAR HCT</u>						
N021532 005	5616599	Apr 25, 2016	DS DP	U-500		
	5616599*PED	Oct 25, 2016				
	6878703	Nov 19, 2021		U-3		Y
	6878703*PED	May 19, 2022				
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u>						
N021162 001	5591762	Jan 07, 2014	DS DP	U-3		
	6358986	Jan 10, 2020				
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u>						
N021162 002	5591762	Jan 07, 2014	DS DP	U-3		
	6358986	Jan 10, 2020				
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u>						
N021162 003	5591762	Jan 07, 2014	DS DP	U-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
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>						
N020818 001	5399578	Mar 21, 2012	U-3		I-567	Jul 31, 2011
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	U-3			
	6294197*PED	Dec 18, 2017				
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>						
N020818 002	5399578	Mar 21, 2012	U-3		I-567	Jul 31, 2011
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	U-3			
	6294197*PED	Dec 18, 2017				
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>						
N020818 003	5399578	Mar 21, 2012	U-3		I-567	Jul 31, 2011
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	U-3			
	6294197*PED	Dec 18, 2017				
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>						
N020818 004	5399578	Mar 21, 2012	U-3		I-567	Jul 31, 2011
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	U-3			
	6294197*PED	Dec 18, 2017				
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>						
N020818 005	5399578	Mar 21, 2012	U-3		I-567	Jul 31, 2011
	5399578*PED	Sep 21, 2012				
	6294197	Jun 18, 2017	U-3			
	6294197*PED	Dec 18, 2017				
<u>HYDROCODONE BITARTRATE; IBUPROFEN - VICOPROFEN</u>						
N020716 001	6348216	Jun 10, 2017				
	6599531	Jun 10, 2017				
<u>HYDROCORTISONE BUTYRATE - LOCOID</u>						
N022076 001	7378405	Dec 19, 2026	DP			
	7981877	Jan 23, 2025	DP			
<u>HYDROCORTISONE BUTYRATE - LOCOID LIPOCREAM</u>						
N020769 001	5635497	Jun 03, 2014			I-613	Oct 19, 2012
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019034 003	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019034 004	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019034 005	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019891 001	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019892 001	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019892 002	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u>						
N019892 003	6589960	Nov 09, 2020	DS DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N019034 001	6589960	Nov 09, 2020	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u>						
N019034 002	6589960	Nov 09, 2020	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - EXALGO</u>						
N021217 001	5702725	Jul 07, 2014	DP U-1043		NDF	Mar 01, 2013
	5914131	Jul 07, 2014	DP U-1043			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>HYDROMORPHONE HYDROCHLORIDE - EXALGO</u>						
N021217 002	5702725	Jul 07, 2014	DP U-1043		NDF	Mar 01, 2013
	5914131	Jul 07, 2014	DP U-1043			
<u>HYDROMORPHONE HYDROCHLORIDE - EXALGO</u>						
N021217 003	5702725	Jul 07, 2014	DP U-1043		NDF	Mar 01, 2013
	5914131	Jul 07, 2014	DP U-1043			
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N021044 001	5958452	Nov 04, 2014	DP			
	5965161	Nov 04, 2014	DP U-616			
	5968551	Dec 24, 2011	DP			
	6294195	Dec 24, 2011	DP			
	6335033	Nov 04, 2014	DP U-616			
	6589960	Nov 09, 2020	DP			
	6706281	Nov 04, 2014	DP U-616			
	6743442	Nov 04, 2014	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N021044 002	5958452	Nov 04, 2014	DP			
	5965161	Nov 04, 2014	DP U-616			
	5968551	Dec 24, 2011	DP			
	6294195	Dec 24, 2011	DP			
	6335033	Nov 04, 2014	DP U-616			
	6589960	Nov 09, 2020	DP			
	6706281	Nov 04, 2014	DP U-616			
	6743442	Nov 04, 2014	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N021044 003	5958452	Nov 04, 2014	DP			
	5965161	Nov 04, 2014	DP U-616			
	5968551	Dec 24, 2011	DP			
	6294195	Dec 24, 2011	DP			
	6335033	Nov 04, 2014	DP U-616			
	6589960	Nov 09, 2020	DP			
	6706281	Nov 04, 2014	DP U-616			
	6743442	Nov 04, 2014	DP			
<u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u>						
N021044 004	5958452	Nov 04, 2014	DP			
	5965161	Nov 04, 2014	DP U-616			
	5968551	Dec 24, 2011	DP			
	6294195	Dec 24, 2011	DP			
	6335033	Nov 04, 2014	DP U-616			
	6589960	Nov 09, 2020	DP			
	6706281	Nov 04, 2014	DP U-616			
	6743442	Nov 04, 2014	DP			
<u>HYDROXOCOBALAMIN - CYANOKIT</u>						
N022041 001	5834448	Nov 14, 2016	DP		ODE	Dec 15, 2013
<u>HYDROXOCOBALAMIN - CYANOKIT</u>						
N022041 002	5834448	Nov 14, 2016	DP U-789		ODE	Dec 15, 2013
<u>HYDROXYPROGESTERONE CAPROATE - MAKENA</u>						
N021945 001					ODE	Feb 03, 2018
<u>IBANDRONATE SODIUM - BONIVA</u>						
N021455 001	4927814	Mar 17, 2012	DS DP U-700			
	4927814	Mar 17, 2012	DS DP U-642			
	6143326	Apr 21, 2017	U-642			
	6294196	Oct 07, 2019	DP			
<u>IBANDRONATE SODIUM - BONIVA</u>						
N021455 002	4927814	Mar 17, 2012	DS DP U-700			
	4927814	Mar 17, 2012	DS DP U-642			
	6294196	Oct 07, 2019	DP			
	7192938	May 06, 2023	U-798			
	7410957	May 06, 2023	U-887			
	7718634	May 06, 2023	U-642			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>IBANDRONATE SODIUM - BONIVA</u>								
N021858 001	4927814 5662918	Mar 17, 2012 Sep 02, 2014		DS DP	U-700 DP			
<u>IBUPROFEN - CALDOLOR</u>								
N022348 001	6727286	Nov 27, 2021		DP	U-981		NP	Jun 11, 2012
<u>IBUPROFEN - CALDOLOR</u>								
N022348 002	6727286	Nov 27, 2021		DP	U-981		NP	Jun 11, 2012
<u>IBUPROFEN - CHILDREN'S MOTRIN</u>								
N020516 001	5374659 5374659*PED	Dec 20, 2011 Jun 20, 2012						
<u>IBUPROFEN - CHILDREN'S MOTRIN</u>								
N020603 001	5374659 5374659*PED	Dec 20, 2011 Jun 20, 2012						
<u>IBUPROFEN - MIDOL LIQUID GELS</u>								
N021472 001	6251426	Jun 25, 2018						
<u>IBUPROFEN - MOTRIN</u>								
N019842 001	5374659 5374659*PED	Dec 20, 2011 Jun 20, 2012						
<u>IBUPROFEN - MOTRIN</u>								
N020135 001	5320855*PED	Dec 14, 2011						
<u>IBUPROFEN - MOTRIN</u>								
N020135 002	5320855*PED	Dec 14, 2011						
<u>IBUPROFEN LYSINE - NEOPROFEN</u>								
N021903 001	6342530 6342530 6344479	Nov 14, 2020 Nov 14, 2020 Mar 20, 2021	DS DP DS	U-794 U-1127 U-794		Y	ODE	Apr 13, 2013
<u>IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - CHILDREN'S MOTRIN COLD</u>								
N021128 001	6211246	Jun 10, 2019						
<u>ICATIBANT ACETATE - FIRAZYR</u>								
N022150 001	5648333	Jul 15, 2014	DS	DP	U-1187		NCE	Aug 25, 2016
							ODE	Aug 25, 2018
<u>ICODEXTRIN - EXTRANEAL</u>								
N021321 001	6077836 6248726	Jun 20, 2017 Jun 19, 2018		U-495 U-495				
<u>ILOPERIDONE - FANAPT</u>								
N022192 001	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>								
N022192 002	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>								
N022192 003	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>								
N022192 004	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>								
N022192 005	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>								
N022192 006	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>								
N022192 007	RE39198	Nov 15, 2016	DS	DP	U-971		NCE	May 06, 2014
<u>ILOPROST - VENTAVIS</u>								
N021779 001							ODE	Dec 29, 2011

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ILOPROST - VENTAVIS</u>						
	N021779 002				ODE	Dec 29, 2011
<u>ILOPROST - VENTAVIS</u>						
	N021779 003				ODE	Dec 29, 2011
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N021335 001	5521184	Jan 04, 2015	DS DP			
	5521184*PED	Jul 04, 2015				
	6894051	May 23, 2019	DS DP	U-649		
	6894051*PED	Nov 23, 2019				
	6958335	Dec 19, 2021		U-791		
	6958335*PED	Jun 19, 2022				
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N021335 002	5521184	Jan 04, 2015	DS DP	U-649		
	5521184*PED	Jul 04, 2015				
	6894051	May 23, 2019	DS DP	U-649		
	6894051*PED	Nov 23, 2019				
	6958335	Dec 19, 2021		U-791		
	6958335*PED	Jun 19, 2022				
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N021588 001	5521184	Jan 04, 2015	DS DP		I-583	Dec 19, 2011
	5521184*PED	Jul 04, 2015			ODE	Oct 19, 2013
	6894051	May 23, 2019	DS DP	U-649	ODE	Oct 19, 2013
	6894051*PED	Nov 23, 2019			ODE	Oct 19, 2013
	6958335	Dec 19, 2021		U-791	ODE	Oct 19, 2013
	6958335*PED	Jun 19, 2022			ODE	Oct 19, 2013
	7544799	Jan 16, 2019	DS DP		ODE	Oct 19, 2013
	7544799*PED	Jul 16, 2019				
<u>IMATINIB MESYLATE - GLEEVEC</u>						
N021588 002	5521184	Jan 04, 2015			I-583	Dec 19, 2011
	5521184*PED	Jul 04, 2015			ODE	Oct 19, 2013
	6894051	May 23, 2019	DS DP	U-649	ODE	Oct 19, 2013
	6894051*PED	Nov 23, 2019			ODE	Oct 19, 2013
	6958335	Dec 19, 2021		U-791	ODE	Oct 19, 2013
	6958335*PED	Jun 19, 2022			ODE	Oct 19, 2013
	7544799	Jan 16, 2019	DS DP		ODE	Oct 19, 2013
	7544799*PED	Jul 16, 2019				
<u>IMIGLUCERASE - CEREZYME</u>						
N020367 001	5549892	Aug 27, 2013		U-252		
<u>IMIGLUCERASE - CEREZYME</u>						
N020367 002	5549892	Aug 27, 2013		U-252		
<u>IMIQUIMOD - ALDARA</u>						
N020723 001	7696159	Apr 01, 2024	DS	U-1048		
	7696159	Apr 01, 2024	DS	U-1047		
	7696159*PED	Oct 01, 2024				
<u>IMIQUIMOD - ZYCLARA</u>						
N022483 001					I-636	Mar 24, 2014
					NP	Mar 25, 2013
<u>IMIQUIMOD - ZYCLARA</u>						
N022483 002					NS	Jul 15, 2014
<u>INDACATEROL MALEATE - ARCAPTA NEOHALER</u>						
N022383 001	6878721	Oct 10, 2020	DS DP	U-1168	NCE	Jul 01, 2016
<u>INDINAVIR SULFATE - CRIXIVAN</u>						
N020685 001	5413999	May 09, 2012		U-132		
	6645961	Mar 04, 2018	DP			
	6689761	Feb 10, 2021		U-554		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
INDINAVIR SULFATE - CRIXIVAN							
N020685 003	5413999	May	09, 2012		U-132		
	6645961	Mar	04, 2018	DP			
	6689761	Feb	10, 2021		U-554		
INDINAVIR SULFATE - CRIXIVAN							
N020685 005	5413999	May	09, 2012		U-132		
	6645961	Mar	04, 2018	DP			
	6689761	Feb	10, 2021		U-554		
INDINAVIR SULFATE - CRIXIVAN							
N020685 006	5413999	May	09, 2012		U-132		
	6645961	Mar	04, 2018	DP			
	6689761	Feb	10, 2021		U-554		
INDIUM IN-111 PENTETREOTIDE KIT - OCTREOSCAN							
N020314 001	5384113	Jan	24, 2012	DP			
	5753627	May	19, 2015		U-1125		
	5776894	Jul	07, 2015	DS DP			
INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT - NOVOLOG MIX 50/50							
N021810 001	5547930	Sep	28, 2013	DS DP			
	5618913	Jun	07, 2014	DS DP			
	5618913*PED	Dec	07, 2014				
	5834422	Sep	28, 2013	DP U-471			
	5840680	Sep	28, 2013	DS DP	U-471		
	5866538	Jun	20, 2017		DP		
	5866538*PED	Dec	20, 2017				
INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT - NOVOLOG MIX 70/30							
N021172 001	5547930	Sep	28, 2013	DS DP			
	5618913	Jun	07, 2014	DS DP			
	5618913*PED	Dec	07, 2014				
	5834422	Sep	28, 2013	DP U-471			
	5840680	Sep	28, 2013	DS DP	U-471		
	5866538	Jun	19, 2017		DP		
INSULIN ASPART RECOMBINANT - NOVOLOG							
N020986 001	5618913	Jun	07, 2014	DS DP			
	5618913*PED	Dec	07, 2014				
	5866538	Jun	20, 2017				
	5866538*PED	Dec	20, 2017				
INSULIN DETEMIR RECOMBINANT - LEVEMIR							
N021536 001	5750497	May	16, 2019	DS DP	U-668		
	5866538	Jun	20, 2017		DP		
	6011007	Feb	02, 2014	DS DP	U-668		
	6869930	Feb	02, 2014	DS DP	U-668		
INSULIN GLARGINE RECOMBINANT - LANTUS							
N021081 001	5656722	Aug	12, 2014	DS DP	U-948		
	5656722*PED	Feb	12, 2015				
	7476652	Jul	23, 2023		DP		
	7476652*PED	Jan	23, 2024				
	7713930	Jun	13, 2023		DP		
	7713930*PED	Dec	13, 2023				
	7918833	Sep	23, 2027		DP		
	7918833*PED	Mar	23, 2028				
INSULIN GLULISINE RECOMBINANT - APIDRA							
N021629 001	6221633	Jun	18, 2018	DS DP	U-471	NPP	Oct 24, 2011
	6960561	Jan	25, 2023	DP	U-471		
	7452860	Mar	22, 2022		DP		
	7696162	Mar	22, 2022	DP	U-471		
INSULIN GLULISINE RECOMBINANT - APIDRA							
N021629 002	6221633	Jun	18, 2018	DS DP	U-471	NPP	Oct 24, 2011
	6960561	Jan	25, 2023	DP	U-471		
	7452860	Mar	22, 2022		DP		
	7696162	Mar	22, 2022	DP	U-471		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u>						
N021629 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		
<u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 50/50</u>						
N021018 001	5461031	Jun 26, 2014				
	5474978	Jun 16, 2014				
	5514646	May 07, 2013				
	5747642	Jun 16, 2014				
<u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 75/25</u>						
N021017 001	5461031	Jun 16, 2014				
	5474978	Jun 16, 2014				
	5514646	May 07, 2013				
	5747642	Jun 16, 2014				
<u>INSULIN LISPRO RECOMBINANT - HUMALOG</u>						
N020563 001	5474978	Jun 16, 2014		U-534		
	5514646	May 07, 2013		U-534		
<u>INSULIN LISPRO RECOMBINANT - HUMALOG PEN</u>						
N020563 002	5474978	Jun 16, 2014		U-534		
	5514646	May 07, 2013		U-534		
<u>INSULIN RECOMBINANT HUMAN - EXUBERA</u>						
N021868 001	5740794	Apr 21, 2015		DP		
	5997848	Mar 07, 2014		U-704		
	6051256	Mar 07, 2014		DP		
	6257233	May 14, 2019		U-704		
	6423344	Mar 07, 2014		DP		
	6543448	Sep 21, 2014		DP		
	6546929	May 14, 2019		U-704		
	6582728	Jun 24, 2020		DP		
	6592904	Mar 07, 2014		DP		
	6685967	Sep 11, 2018		DP		
	6737045	Mar 07, 2014		U-704		
<u>INSULIN RECOMBINANT HUMAN - EXUBERA</u>						
N021868 002	5740794	Apr 21, 2015		DP		
	5997848	Mar 07, 2014		U-704		
	6051256	Mar 07, 2014		DP		
	6257233	May 14, 2019		U-704		
	6423344	Mar 07, 2014		DP		
	6543448	Sep 21, 2014		DP		
	6546929	May 14, 2019		U-704		
	6582728	Jun 24, 2020		DP		
	6592904	Mar 07, 2014		DP		
	6685967	Sep 11, 2018		DP		
	6737045	Mar 07, 2014		U-704		
<u>INSULIN RECOMBINANT HUMAN - HUMULIN R</u>						
N018780 001					NR	Mar 25, 2014
<u>INSULIN RECOMBINANT HUMAN - HUMULIN R PEN</u>						
N018780 005					NR	Mar 25, 2014
<u>IOBENGUANE SULFATE I-123 - ADREVIEW</u>						
N022290 001					NCE	Sep 19, 2013
					ODE	Sep 19, 2015
<u>IODIXANOL - VISIPAQUE 270</u>						
N020351 001	5349085	Sep 20, 2011				
	5366722	Nov 22, 2011		DP		
	RE36418	Jul 12, 2011		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>IODIXANOL - VISIPAQUE 270</u>						
N020808 001	5349085	Sep 20, 2011				
	5366722	Nov 22, 2011	DP			
	RE36418	Jul 12, 2011	DP			
<u>IODIXANOL - VISIPAQUE 320</u>						
N020351 002	5349085	Sep 20, 2011				
	RE36418	Jul 12, 2011	DP			
<u>IODIXANOL - VISIPAQUE 320</u>						
N020808 002	5349085	Sep 20, 2011				
	RE36418	Jul 12, 2011	DP			
<u>IOFLUPANE I-123 - DATSCAN</u>						
N022454 001					NCE	Jan 14, 2016
<u>IOPROMIDE - ULTRAVIST (PHARMACY BULK)</u>						
N021425 002					I-619	Dec 30, 2012
<u>IOPROMIDE - ULTRAVIST 370</u>						
N020220 001					I-619	Dec 30, 2012
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>						
N021527 001	5676930	Oct 14, 2014	DP			
	5683677	Nov 04, 2014	DP			
	6739333	May 26, 2020	DP			
	6983743	May 26, 2020	DP			
<u>IRBESARTAN - AVAPRO</u>						
N020757 001	5270317	Sep 30, 2011				
	5270317*PED	Mar 30, 2012				
	6342247	Jun 07, 2015				
	6342247*PED	Dec 07, 2015				
<u>IRBESARTAN - AVAPRO</u>						
N020757 002	5270317	Sep 30, 2011				
	5270317*PED	Mar 30, 2012				
	6342247	Jun 07, 2015				
	6342247*PED	Dec 07, 2015				
<u>IRBESARTAN - AVAPRO</u>						
N020757 003	5270317	Sep 30, 2011				
	5270317*PED	Mar 30, 2012				
	6342247	Jun 07, 2015				
	6342247*PED	Dec 07, 2015				
<u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u>						
N020571 001	6403569	Apr 28, 2020		U-449		
	6403569*PED	Oct 28, 2020				
	6794370	May 01, 2020		U-606		
	6794370*PED	Nov 01, 2020				
<u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u>						
N020571 002	6403569	Apr 28, 2020		U-449		
	6403569*PED	Oct 28, 2020				
	6794370	May 01, 2020		U-606		
	6794370*PED	Nov 01, 2020				
<u>IRON DEXTRAN - DEXFERRUM</u>						
A040024 001	5624668	Sep 29, 2015				
<u>ITRACONAZOLE - ONMEL</u>						
N022484 001	6509038	May 12, 2017	DP	U-1054		
	7081255	May 12, 2017	DP	U-1054		
<u>ITRACONAZOLE - SPORANOX</u>						
N020083 001	5633015	May 27, 2014				
<u>ITRACONAZOLE - SPORANOX</u>						
N020657 001	5707975	Jan 13, 2015				
	6407079	Jun 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
<u>ITRACONAZOLE - SPORANOX</u>						
N020966 001	6407079	Jun 18, 2019				
<u>IXABEPILONE - IXEMPRAL KIT</u>						
N022065 001	6670384	Jan 23, 2022	DP U-959		M-61	Oct 18, 2014
	6670384	Jan 23, 2022	DP U-960		NCE	Oct 16, 2012
	6670384*PED	Jul 23, 2022			PED	Apr 18, 2015
	7022330	Jan 23, 2022	DP U-958		PED	Apr 16, 2013
	7022330*PED	Jul 23, 2022				
	7125899	May 26, 2018	DS DP U-957			
	7125899*PED	Nov 26, 2018				
	7312237	Aug 21, 2024	U-965			
	7312237*PED	Feb 21, 2025				
	RE41393	Feb 08, 2022	U-961			
	RE41393*PED	Aug 08, 2022				
	RE41911	May 26, 2018	DS DP U-961			
	RE41911*PED	Nov 26, 2018				
<u>IXABEPILONE - IXEMPRAL KIT</u>						
N022065 002	6670384	Jan 23, 2022	DP U-960		M-61	Oct 18, 2014
	6670384	Jan 23, 2022	DP U-959		NCE	Oct 16, 2012
	6670384*PED	Jul 23, 2022			PED	Apr 18, 2015
	7022330	Jan 23, 2022	DP U-958		PED	Apr 16, 2013
	7022330*PED	Jul 23, 2022				
	7125899	May 26, 2018	DS DP U-957			
	7125899*PED	Nov 26, 2018				
	7312237	Aug 21, 2024	U-965			
	7312237*PED	Feb 21, 2025				
	RE41393	Feb 08, 2022	U-961			
	RE41393*PED	Aug 08, 2022				
	RE41911	May 26, 2018	DS DP U-961			
	RE41911*PED	Nov 26, 2018				
<u>KETOCONAZOLE - EXTINA</u>						
N021738 001	7553835	Oct 19, 2018	DP U-245			
<u>KETOCONAZOLE - KETOCONAZOLE</u>						
A091550 001					PC	Feb 21, 2012
<u>KETOCONAZOLE - NIZORAL A-D</u>						
N020310 001	5456851	Apr 07, 2014				
<u>KETOCONAZOLE - XOLEGEL</u>						
N021946 001	7179475	Dec 04, 2018	DP U-792			
<u>KETOROLAC TROMETHAMINE - ACULAR LS</u>						
N021528 001	8008338	May 24, 2027	U-1181			
<u>KETOROLAC TROMETHAMINE - ACUVAIL</u>						
N022427 001	7842714	Aug 15, 2029	DS DP		NP	Jul 22, 2012
<u>KETOROLAC TROMETHAMINE - SPRIX</u>						
N022382 001	6333044	Dec 25, 2018	DP U-1057		NDF	May 14, 2013
	7476689	Oct 11, 2012	DP U-1056			
<u>LACOSAMIDE - VIMPAT</u>						
N022253 001	5654301	Aug 05, 2014	DS DP U-914		NCE	Oct 28, 2013
	RE38551	Mar 17, 2017	DS DP U-914			
<u>LACOSAMIDE - VIMPAT</u>						
N022253 002	5654301	Aug 05, 2014	DS DP U-914		NCE	Oct 28, 2013
	RE38551	Mar 17, 2017	DS DP U-914			
<u>LACOSAMIDE - VIMPAT</u>						
N022253 003	5654301	Aug 05, 2014	DS DP U-914		NCE	Oct 28, 2013
	RE38551	Mar 17, 2017	DS DP U-914			
<u>LACOSAMIDE - VIMPAT</u>						
N022253 004	5654301	Aug 05, 2014	DS DP U-914		NCE	Oct 28, 2013
	RE38551	Mar 17, 2017	DS DP U-914			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE			PATENT DELIST REQUESTED			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>LACOSAMIDE - VIMPAT</u>										
N022254 001	5654301	Aug 05, 2014	DS	DP	U-911			NCE	Oct 28, 2013	
	RE38551	Mar 17, 2017	DS	DP	U-911					
<u>LACOSAMIDE - VIMPAT</u>										
N022255 001	5654301	Aug 05, 2014	DS	DP	U-914			NCE	Oct 28, 2013	
	RE38551	Mar 17, 2017	DS	DP	U-914					
<u>LAMIVUDINE - EPIVIR</u>										
N020564 001	5905082	May 18, 2016	DS	DP	U-248					
	5905082*PED	Nov 18, 2016								
<u>LAMIVUDINE - EPIVIR</u>										
N020564 003	5905082	May 18, 2016								
	5905082*PED	Nov 18, 2016								
<u>LAMIVUDINE - EPIVIR-HBV</u>										
N021003 001	5905082	May 18, 2016								
	5905082*PED	Nov 18, 2016								
	RE39155	Jul 02, 2013					U-250			
	RE39155*PED	Jan 02, 2014								
<u>LAMIVUDINE - EPIVIR-HBV</u>										
N021004 001	6004968	Mar 20, 2018								
	6004968*PED	Sep 20, 2018								
	RE39155	Jul 02, 2013					U-250			
	RE39155*PED	Jan 02, 2014								
<u>LAMIVUDINE; ZIDOVUDINE - COMBIVIR</u>										
N020857 001	5859021	May 15, 2012	DS	DP	U-248					
	5905082	May 18, 2016	DS	DP	U-248					
	5905082*PED	Nov 18, 2016					U-248			
<u>LAMOTRIGINE - LAMICTAL CD</u>										
N020764 001	5698226	Jan 29, 2012								
	5698226*PED	Jul 29, 2012								
<u>LAMOTRIGINE - LAMICTAL CD</u>										
N020764 002	5698226	Jan 29, 2012								
	5698226*PED	Jul 29, 2012								
<u>LAMOTRIGINE - LAMICTAL CD</u>										
N020764 003	5698226	Jan 29, 2012								
	5698226*PED	Jul 29, 2012								
<u>LAMOTRIGINE - LAMICTAL CD</u>										
N020764 004	5698226	Jan 29, 2012								
	5698226*PED	Jul 29, 2012								
<u>LAMOTRIGINE - LAMICTAL ODT</u>										
N022251 001	7919115	Jan 04, 2029	DS	DP						
<u>LAMOTRIGINE - LAMICTAL ODT</u>										
N022251 002	7919115	Jan 04, 2029	DS	DP						
<u>LAMOTRIGINE - LAMICTAL ODT</u>										
N022251 003	7919115	Jan 04, 2029	DS	DP						
<u>LAMOTRIGINE - LAMICTAL ODT</u>										
N022251 004	7919115	Jan 04, 2029	DS	DP						
<u>LAMOTRIGINE - LAMICTAL XR</u>										
N022115 001								I-644	Apr 25, 2014	
								I-622	Jan 29, 2013	
								NDF	May 29, 2012	
								PED	Nov 29, 2012	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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 XR</u>						
	N022115 002				I-644 I-622 NDF PED	Apr 15, 2014 Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
	N022115 003				I-644 I-622 NDF PED	Apr 25, 2014 Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
	N022115 004				I-644 I-622 NDF PED	Apr 25, 2014 Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
	N022115 005				I-644 I-622 NDF PED	Apr 25, 2014 Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
	N022115 006				I-622 I-644 NDF PED	Jan 29, 2013 Apr 25, 2014 May 29, 2012 Nov 29, 2012
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
	N022074 001 5595760	Mar 08, 2020	DP U-831		D-131 NCE ODE	Mar 04, 2014 Aug 30, 2012 Aug 30, 2014
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
	N022074 002 5595760	Mar 08, 2020	DP U-831		D-131 NCE ODE	Mar 04, 2014 Aug 30, 2012 Aug 30, 2014
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>						
	N022074 003 5595760	Mar 08, 2020	DP U-831		D-131 NCE ODE	Mar 04, 2014 Aug 30, 2012 Aug 30, 2014
<u>LANSOPRAZOLE - PREVACID</u>						
	N020406 001				M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
	N020406 002				M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
	N021281 001				M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
	N021281 002				M-85 PED	Oct 28, 2011 Apr 28, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
LANSOPRAZOLE - PREVACID							
N021428 001	5464632	Nov	07, 2012			M-85	Oct 28, 2011
	5464632*PED	May	07, 2013			PED	Apr 28, 2012
	6328994	May	17, 2019				
	6328994*PED	Nov	17, 2019				
	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				
LANSOPRAZOLE - PREVACID							
N021428 002	5464632	Nov	07, 2012			M-85	Oct 28, 2011
	5464632*PED	May	07, 2013			PED	Apr 28, 2012
	6328994	May	17, 2019				
	6328994*PED	Nov	17, 2019				
	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				
LANSOPRAZOLE - PREVACID 24 HR							
N022327 001						NP	May 18, 2012
LANTHANUM CARBONATE - FOSRENOL							
N021468 001	7396841	Aug	17, 2021	DP	U-947		
	7396841*PED	Feb	17, 2022				
LANTHANUM CARBONATE - FOSRENOL							
N021468 002	5968976	Oct	26, 2018	DP	U-613		
	7381428	Aug	26, 2024		U-890		
	7465465	Aug	26, 2024	DP			
LANTHANUM CARBONATE - FOSRENOL							
N021468 003	5968976	Oct	26, 2018	DP	U-613		
	7381428	Aug	26, 2024		U-890		
	7465465	Aug	26, 2024	DP			
LANTHANUM CARBONATE - FOSRENOL							
N021468 004	5968976	Oct	26, 2018	DP	U-613		
	7381428	Aug	26, 2024		U-890		
	7465465	Aug	26, 2024	DP			
LAPATINIB DITOSYLATE - TYKERB							
N022059 001	6391874	Jul	11, 2017	DS	DP U-800	I-620	Jan 29, 2013
	6713485	Sep	29, 2020	DS	DP U-800	NCE	Mar 13, 2012
	6727256	Jan	08, 2019	DS	DP U-800		
	6828320	Jul	11, 2017		U-800		
	7157466	Nov	19, 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
<u>LENALIDOMIDE - REVLIMIT</u>						
N021880 001	5635517	Oct 04, 2019	DS	U-866	ODE	Jun 29, 2013
	6045501	Aug 28, 2018		U-694	ODE	Dec 27, 2012
	6281230	Jul 24, 2016		U-769		
	6315720	Oct 23, 2020		U-694		
	6555554	Jul 24, 2016	DP			
	6561976	Aug 28, 2018		U-694		
	6561977	Oct 23, 2020		U-694		
	6755784	Oct 23, 2020		U-694		
	6908432	Aug 28, 2018		U-694		
	7119106	Jul 24, 2016	DP			
	7189740	Apr 11, 2023		U-1079		
	7465800	Apr 22, 2026	DS DP			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1165		
<u>LENALIDOMIDE - REVLIMIT</u>						
N021880 002	5635517	Oct 04, 2019	DS	U-866	ODE	Jun 29, 2013
	6045501	Aug 28, 2018		U-694	ODE	Dec 27, 2012
	6281230	Jul 24, 2016		U-769		
	6315720	Oct 23, 2020		U-694		
	6555554	Jul 24, 2016	DP			
	6561976	Aug 28, 2018		U-694		
	6561977	Oct 23, 2020		U-694		
	6755784	Oct 23, 2020		U-694		
	6908432	Aug 28, 2018		U-694		
	7119106	Jul 24, 2016	DP			
	7189740	Apr 11, 2023		U-1079		
	7465800	Apr 22, 2026	DS DP			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1165		
<u>LENALIDOMIDE - REVLIMIT</u>						
N021880 003	5635517	Oct 04, 2019	DS	U-866	ODE	Jun 29, 2013
	6045501	Aug 28, 2018		U-694		
	6281230	Jul 24, 2016		U-769		
	6315720	Oct 23, 2020		U-694		
	6555554	Jul 24, 2016	DP			
	6561976	Aug 28, 2018		U-694		
	6561977	Oct 23, 2020		U-694		
	6755784	Oct 23, 2020		U-694		
	6908432	Aug 28, 2018		U-694		
	7119106	Jul 24, 2016	DP			
	7189740	Apr 11, 2023		U-1079		
	7465800	Apr 22, 2026	DS DP			
	7968569	Oct 07, 2023		U-1165		
<u>LENALIDOMIDE - REVLIMIT</u>						
N021880 004	5635517	Oct 04, 2019	DS	U-866	ODE	Jun 29, 2013
	6045501	Aug 28, 2018		U-694		
	6281230	Jul 24, 2016		U-769		
	6315720	Oct 23, 2020		U-694		
	6555554	Jul 24, 2016	DP			
	6561976	Aug 28, 2018		U-694		
	6561977	Oct 23, 2020		U-694		
	6755784	Oct 23, 2020		U-694		
	6908432	Aug 28, 2018		U-694		
	7119106	Jul 24, 2016	DP			
	7189740	Apr 11, 2023		U-1079		
	7465800	Apr 22, 2026	DS DP			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1165		
<u>LENALIDOMIDE - REVLIMIT</u>						
N021880 005					ODE	Dec 27, 2012
					ODE	Jun 29, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LEUPROLIDE ACETATE - ELIGARD</u>						
N021343 001	5324519	Oct 20, 2011				
	5599552	Feb 04, 2014				
	6395293	Sep 28, 2013	DP	U-801		
	6565874	Oct 28, 2018	DP	U-801		
	6626870	Mar 27, 2020	DP			
	6773714	Oct 28, 2018	DP	U-801		
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N021379 001	5324519	Oct 20, 2011				
	5599552	Feb 04, 2014				
	6395293	Sep 28, 2013	DP			
	6565874	Oct 28, 2018	DP	U-801		
	6626870	Mar 27, 2020	DP			
	6773714	Oct 28, 2018	DP	U-801		
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N021488 001	5324519	Oct 20, 2011				
	5599552	Feb 04, 2014				
	6395293	Sep 28, 2013	DP			
	6565874	Oct 28, 2018	DP	U-801		
	6626870	Mar 27, 2020	DP			
	6773714	Oct 28, 2018	DP	U-801		
<u>LEUPROLIDE ACETATE - ELIGARD</u>						
N021731 001	5599552	Feb 04, 2014	DP	U-621		
	6395293	Sep 28, 2013	DP			
	6565874	Oct 28, 2018	DP	U-621		
	6626870	Mar 27, 2020	DP			
	6773714	Oct 28, 2018	DP	U-621		
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N019732 001	5575987	Sep 02, 2013				
	5631020	May 20, 2014				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N020011 001	5575987	Sep 02, 2013				
	5631021	May 20, 2014				
	5716640	Sep 02, 2013				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N020517 001	5480656	Jan 02, 2013				
	5575987	Sep 02, 2013				
	5631020	May 20, 2014				
	5643607	Jan 02, 2013				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N020517 002	5480656	Jan 02, 2013				
	5575987	Sep 02, 2013				
	5631020	May 20, 2014				
	5643607	Jan 02, 2013				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N020517 003	6036976	Dec 13, 2016	DP	D-132	Jun 17, 2014	
	7429559	Dec 13, 2016	DP	NS	Jun 17, 2014	
<u>LEUPROLIDE ACETATE - LUPRON DEPOT</u>						
N020708 001	5480656	Jan 02, 2013				
	5575987	Sep 02, 2013				
	5631020	May 20, 2014				
	5643607	Jan 02, 2013				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 002	5575987	Sep 02, 2013			M-107	Oct 08, 2014
	5631020	May 20, 2014				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 003	5575987	Sep 02, 2013				
	5631020	May 20, 2014				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2013				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 004	5575987	Sep 02, 2013				
	5631020	May 20, 2014				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2013				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 005	5575987	Sep 02, 2013			M-107	Oct 08, 2014
	5631020	May 20, 2014				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 006	5575987	Sep 02, 2013			M-107	Oct 08, 2014
	5631020	May 20, 2014				
	5716640	Sep 02, 2013				
	6036976	Dec 13, 2016				
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 007	5575987	Sep 02, 2013	DP		NP	Aug 15, 2014
	5631020	May 20, 2014	DP			
	5716640	Sep 02, 2013	DP			
	6036976	Dec 13, 2016	DP			
<u>LEUPROLIDE ACETATE - LUPRON DEPOT-PED</u>						
N020263 008	5480656	Jan 02, 2013	DP		NP	Aug 15, 2014
	5575987	Sep 02, 2013	DP			
	5631020	May 20, 2014	DP			
	5643607	Jan 02, 2013	DP			
	5716640	Sep 02, 2013	DP			
	6036976	Dec 13, 2016	DP			
<u>LEUPROLIDE ACETATE - VIADUR</u>						
N021088 001	5728396	Jan 30, 2017		U-316		
	5932547	Jun 13, 2017				
	5985305	Jan 30, 2017				
	6113938	Jul 24, 2018				
	6124261	Jun 13, 2017				
	6132420	Jan 30, 2017				
	6156331	Jan 30, 2017				
	6235712	Jun 13, 2017				
	6375978	Dec 17, 2018				
	6395292	Jan 30, 2017				
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N020837 001	5362755	Mar 25, 2013		U-332		
	5547994	Aug 20, 2013		U-332		
	6451289	Mar 21, 2021				
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N020837 002	5362755	Mar 25, 2013		U-332		
	5547994	Aug 20, 2013		U-332		
	6451289	Mar 21, 2021				
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N020837 003	5362755	Mar 25, 2013		U-332		
	5547994	Aug 20, 2013		U-332		
	6451289	Mar 21, 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
<u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u>						
N020837 004	5362755	Mar 25, 2013		U-332		
	5547994	Aug 20, 2013		U-332		
	6451289	Mar 21, 2021	DP			
<u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u>						
N021730 001	5362755	Mar 25, 2013		U-636		
	5547994	Aug 20, 2013		U-636		
	5605674	Feb 25, 2014	DP			
	5836299	Nov 17, 2015	DP			
	7256310	Oct 08, 2024	DS DP	U-636		
<u>LEVETIRACETAM - KEPPRA XR</u>						
N022285 001	7858122	Sep 17, 2028	DP		NDF	Sep 12, 2011
<u>LEVETIRACETAM - KEPPRA XR</u>						
N022285 002	7858122	Sep 17, 2028	DP		NDF	Sep 12, 2011
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091093 001					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091093 002					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091261 002					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091285 001					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091285 002					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091291 002					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091399 001					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091399 002					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091430 001					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091430 002					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091557 001					PC	Mar 10, 2012
<u>LEVETIRACETAM - LEVETIRACETAM</u>						
A091557 002					PC	Mar 10, 2012
<u>LEVOBETAXOLOL HYDROCHLORIDE - BETAXON</u>						
N021114 001	5540918	Jul 30, 2013	DP			
	5540918*PED	Jan 30, 2014				
<u>LEVOBUPIVACAINE HYDROCHLORIDE - CHIROCAINE</u>						
N020997 001	5708011	Oct 13, 2014		U-276		
<u>LEVOBUPIVACAINE HYDROCHLORIDE - CHIROCAINE</u>						
N020997 002	5708011	Oct 13, 2014		U-276		
<u>LEVOBUPIVACAINE HYDROCHLORIDE - CHIROCAINE</u>						
N020997 003	5708011	Oct 13, 2014		U-276		
<u>LEVOCARNITINE - CARNITOR</u>						
N020182 001	6335369	Jan 18, 2021		U-433		
	6429230	Jan 18, 2021		U-433		
	6696493	Jan 18, 2021		U-433		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LEVO CETIRIZINE DIHYDROCHLORIDE - XYZAL</u>						
N022064 001	5698558	Sep 24, 2012	U-812		NPP	Aug 21, 2012
	5698558*PED	Mar 24, 2013			PED	Feb 21, 2013
<u>LEVO CETIRIZINE DIHYDROCHLORIDE - XYZAL</u>						
N022157 001	5698558	Sep 24, 2012	U-852		NPP	Aug 21, 2012
	5698558*PED	Mar 24, 2013			PED	Feb 21, 2013
<u>LEVOFLOXACIN - LEVAQUIN</u>						
N021721 001	6806256	Feb 26, 2022	DP			
	6806256*PED	Aug 26, 2022				
<u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u>						
N020140 001	6500829	Dec 31, 2019	DS DP		I-637	Apr 29, 2014
					ODE	Apr 29, 2018
					ODE	Mar 07, 2015
<u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u>						
N020140 002	6500829	Dec 31, 2019	DS DP		I-637	Apr 29, 2014
					ODE	Apr 29, 2018
					ODE	Mar 07, 2015
<u>LEVONORGESTREL - MIRENA</u>						
N021225 001	5785053	Dec 05, 2015	DP		I-610	Oct 01, 2012
<u>LEVONORGESTREL - PLAN B ONE-STEP</u>						
N021998 001					NP	Jul 10, 2012
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 001	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 002	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 003	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 004	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 005	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 006	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 007	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 008	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 009	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 010	6399101	Mar 30, 2020				
<u>LEVOTHYROXINE SODIUM - LEVO-T</u>						
N021342 011	6399101	Mar 30, 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>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 001	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 002	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 003	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 004	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 005	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 006	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 007	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 008	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 009	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 010	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 011	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>						
N021301 012	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Aug 14, 2022	DP		U-759	
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 002	7723390	Mar 14, 2024				
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 003	7723390	Mar 14, 2024				
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 004	7723390	Mar 14, 2024				
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 005	7723390	Mar 14, 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
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 006	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 007	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 008	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 009	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N021924 010	7723390	Mar 14, 2024	DP			
<u>LEVOTHYROXINE SODIUM - TIROSINT</u>						
N022121 001	7723390	Mar 14, 2024	DP			
<u>LIDOCAINE - DENTIPATCH</u>						
N020575 001	5332576	Jul 26, 2011				
<u>LIDOCAINE - DENTIPATCH</u>						
N020575 002	5332576	Jul 26, 2011				
<u>LIDOCAINE - LIDODERM</u>						
N020612 001	5411738	May 02, 2012				
	5601838	May 02, 2012			U-488	
	5741510	Mar 30, 2014	DP			
	5827529	Oct 27, 2015			U-486	
<u>LIDOCAINE HYDROCHLORIDE - AKTEN</u>						
N022221 001					NDF	Oct 07, 2011
<u>LIDOCAINE HYDROCHLORIDE - ZINGO</u>						
N022114 001	5630796	May 20, 2014			NPP	Jan 08, 2012
	5899880	May 04, 2016	DP			
	6004286	Mar 17, 2017	DP			
	6881200	Jun 11, 2016	DP			
<u>LIDOCAINE; PRILOCAINE - ORAQIX</u>						
N021451 001	6031007	Apr 01, 2017	DP	U-553		
<u>LIDOCAINE; TETRACAINE - LIDOCAINE AND TETRACAINE</u>						
N021717 001	5919479	Jul 28, 2015	DP			
	6528086	Sep 28, 2019	DP			
<u>LIDOCAINE; TETRACAINE - SYNERA</u>						
N021623 001	5658583	Jul 28, 2015	DP			
	5919479	Jul 28, 2015	DP			
	6306431	Jul 28, 2015	DP			
	6465006	Jul 28, 2015	DP			
	6546281	Jul 28, 2015	DP			
	6780426	Jul 28, 2015	DP			
<u>LINAGLIPTIN - TRADJENTA</u>						
N201280 001	6303661	Apr 24, 2017		U-774		
	6890898	Feb 02, 2019		U-493		
	7078381	Feb 02, 2019		U-493		
	7407955	Aug 12, 2023	DS	DP		
	7459428	Feb 02, 2019			U-493	
<u>LINEZOLID - ZYVOX</u>						
N021130 001	5688792	Nov 18, 2014	DS	U-319		
	5688792*PED	May 18, 2015				
	6514529	Mar 15, 2021		DP		
	6514529*PED	Sep 15, 2021				
	6559305	Jan 29, 2021	DS			
	6559305*PED	Jul 29, 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
<u>LINEZOLID - ZYVOX</u>							
N021130 002	5688792	Nov	18, 2014	DS	U-319		
	5688792*PED	May	18, 2015				
	6514529	Mar	15, 2021	DP			
	6514529*PED	Sep	15, 2021				
	6559305	Jan	29, 2021	DS			
	6559305*PED	Jul	29, 2021				
<u>LINEZOLID - ZYVOX</u>							
N021131 001	5688792	Nov	18, 2014		U-319		
	5688792*PED	May	18, 2015				
	6559305	Jan	29, 2021	DS			
	6559305*PED	Jul	29, 2021				
<u>LINEZOLID - ZYVOX</u>							
N021132 001	5688792	Nov	18, 2014	DS	U-319		
	5688792*PED	May	18, 2015				
	6559305	Jan	29, 2021	DS			
	6559305*PED	Jul	29, 2021				
<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>							
N022341 001	6268343	Aug	22, 2017	DS	DP U-968	NCE	
	6458924	Aug	22, 2017	DS	DP U-968		
	7235627	Aug	22, 2017	DS	DP		
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>							
N021977 001	7105486	Jun	29, 2023		U-727	M-82	
	7223735	Jun	29, 2023	DP		NPP	
	7655630	Feb	24, 2023	DS		NCE	
	7659253	Feb	24, 2023	DS	DP U-727		
	7659254	Feb	24, 2023		U-1034		
	7662787	Feb	24, 2023	DS			
	7671030	Feb	24, 2023	DP	U-727		
	7671031	Feb	28, 2023		U-727		
	7674774	Mar	18, 2023	DP	U-842		
	7678770	Mar	25, 2023		U-842		
	7678771	Mar	25, 2023	DP	U-842		
	7687466	Feb	24, 2023		DP		
	7687467	Apr	08, 2023	DP	U-842		
	7700561	Jun	29, 2023		DP		
	7718619	Feb	24, 2023	DP	U-842		
	7723305	Feb	24, 2023	DP	U-842		
<u>LISDEXAMFETAMINE Dimesylate - VYVANSE</u>							
N021977 002	7105486	Jun	29, 2023		U-727	M-82	
	7223735	Jun	29, 2023	DP		NPP	
	7655630	Feb	24, 2023	DS		NCE	
	7659253	Feb	24, 2023	DS	DP U-727		
	7659254	Feb	24, 2023		U-1034		
	7662787	Feb	24, 2023	DS			
	7671030	Feb	24, 2023	DP	U-727		
	7671031	Feb	28, 2023		U-727		
	7674774	Mar	18, 2023	DP	U-842		
	7678770	Mar	25, 2023		U-842		
	7678771	Mar	25, 2023	DP	U-842		
	7687466	Feb	24, 2023		DP		
	7687467	Apr	08, 2023	DP	U-842		
	7700561	Jun	29, 2023		DP		
	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
LISDEXAMFETAMINE Dimesylate - VYVANSE						
N021977 003	7105486	Jun 29, 2023		U-727	M-82	Apr 05, 2013
	7223735	Jun 29, 2023	DP		NPP	Nov 10, 2013
	7655630	Feb 24, 2023	DS		NCE	Feb 23, 2012
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7671030	Feb 24, 2023		DP U-727		
	7671031	Feb 28, 2023		U-727		
	7674774	Mar 18, 2023		DP U-842		
	7678770	Mar 25, 2023		U-842		
	7678771	Mar 25, 2023		DP U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Apr 08, 2023		DP U-842		
	7700561	Jun 29, 2023		DP		
	7718619	Feb 24, 2023		DP U-842		
	7723305	Feb 24, 2023		DP U-842		
LISDEXAMFETAMINE Dimesylate - VYVANSE						
N021977 004	7105486	Jun 29, 2023		U-842	M-82	Apr 05, 2013
	7223735	Jun 29, 2023	DP		NPP	Nov 10, 2013
	7655630	Feb 24, 2023	DS		NCE	Feb 23, 2012
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7671031	Feb 28, 2023		U-727		
	7678770	Mar 25, 2023		U-842		
	7687466	Feb 24, 2023	DP			
	7700561	Jun 29, 2023	DP			
LISDEXAMFETAMINE Dimesylate - VYVANSE						
N021977 005	7105486	Jun 29, 2023		U-842	M-82	Apr 05, 2013
	7223735	Jun 29, 2023	DP		NPP	Nov 10, 2013
	7655630	Feb 24, 2023	DS		NCE	Feb 23, 2012
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 28, 2023		U-727		
	7674774	Mar 18, 2023	DP	U-842		
	7678770	Mar 25, 2023		U-842		
	7678771	Mar 25, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Apr 08, 2023	DP	U-842		
	7700561	Jun 29, 2023		DP		
	7718619	Feb 24, 2023		DP U-842		
	7723305	Feb 24, 2023		DP U-842		
LISDEXAMFETAMINE Dimesylate - VYVANSE						
N021977 006	7105486	Jun 29, 2023		U-842	M-82	Apr 05, 2013
	7223735	Jun 29, 2023	DP		NPP	Nov 10, 2013
	7655630	Feb 24, 2023	DS		NCE	Feb 23, 2012
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 28, 2023		U-727		
	7674774	Mar 18, 2023	DP	U-842		
	7678770	Mar 25, 2023		U-842		
	7678771	Mar 25, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Apr 08, 2023	DP	U-842		
	7700561	Jun 29, 2023		DP		
	7718619	Feb 24, 2023		DP U-842		
	7723305	Feb 24, 2023		DP U-842		
LODOXAMIDE TROMETHAMINE - ALOMIDE						
N020191 001	5457126	Oct 10, 2012		U-117		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LOPERAMIDE HYDROCHLORIDE - IMODIUM A-D EZ CHEWS</u>						
N020448 001	5489436	Feb 06, 2013	DP			
	6814978	Aug 26, 2021	DP			
<u>LOPERAMIDE HYDROCHLORIDE; SIMETHICONE - IMODIUM MULTI-SYMP托M RELIEF</u>						
N020606 001	5489436	Feb 06, 2013				
	5679376	Oct 21, 2014		Y		
	5716641	May 21, 2012	U-226	Y		
<u>LOPERAMIDE HYDROCHLORIDE; SIMETHICONE - IMODIUM MULTI-SYMP托M RELIEF</u>						
N021140 001	6103260	Jul 17, 2017	DP			
<u>LOPINAVIR; RITONAVIR - KALETRA</u>						
N021226 001	5541206	Jul 30, 2013	DS	U-348		
	5541206*PED	Jan 30, 2014				
	5648497	Jul 15, 2014				
	5648497*PED	Jan 15, 2015				
	5886036	Nov 19, 2013	DS DP			
	5886036*PED	May 19, 2014				
	5914332	Dec 13, 2015		U-351		
	5914332*PED	Jun 13, 2016				
	5948436	Sep 13, 2013	DP			
	5948436*PED	Mar 13, 2014				
	6037157	Jun 26, 2016		U-346		
	6037157*PED	Dec 26, 2016				
	6232333	Nov 07, 2017				
	6232333*PED	May 07, 2018				
	6284767	Feb 15, 2016	DP	U-688		
	6284767	Feb 15, 2016	DP	U-401		
	6284767*PED	Aug 15, 2016				
	6458818	Nov 07, 2017				
	6458818*PED	May 07, 2018				
	6521651	Nov 07, 2017	DP			
	6521651*PED	May 07, 2018				
	6703403	Jun 26, 2016		U-257		
	6703403*PED	Dec 26, 2016				
	7141593	May 22, 2020	DP			
	7141593*PED	Nov 22, 2020				
	7432294	May 22, 2020	DP			
	7432294*PED	Nov 22, 2020				
<u>LOPINAVIR; RITONAVIR - KALETRA</u>						
N021251 001	5484801	Jan 28, 2014	DP		D-124	Apr 27, 2013
	5484801*PED	Jul 28, 2014				
	5541206	Jul 30, 2013	DS DP	U-348		
	5541206	Jul 30, 2013	DS DP	U-895		
	5541206*PED	Jan 30, 2014				
	5648497	Jul 15, 2014	DS			
	5648497*PED	Jan 15, 2015				
	5886036	Nov 19, 2013	DS DP	U-895		
	5886036*PED	May 19, 2014				
	5914332	Dec 13, 2015	DS DP	U-895		
	5914332	Dec 13, 2015	DS DP	U-351		
	5914332*PED	Jun 13, 2016				
	5948436	Sep 13, 2013	DP			
	5948436*PED	Mar 13, 2014				
	6037157	Jun 26, 2016		U-895		
	6037157	Jun 26, 2016		U-346		
	6037157*PED	Dec 26, 2016				
	6284767	Feb 15, 2016	DP	U-895		
	6284767	Feb 15, 2016	DP	U-401		
	6284767*PED	Aug 15, 2016				
	6703403	Jun 26, 2016		U-257		
	6703403	Jun 26, 2016		U-895		
	6703403*PED	Dec 26, 2016				
	6911214	Nov 28, 2021	DP	U-895		
	6911214*PED	May 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
<u>LOPINAVIR; RITONAVIR - KALETRA</u>								
N021906 001	5541206	Jul	30,	2013	DS DP U-688		D-124	Apr 27, 2013
	5541206*PED	Jan	30,	2014				
	5648497	Jul	15,	2014	DS DP			
	5648497*PED	Jan	15,	2015				
	5886036	Nov	19,	2013	DS DP U-895			
	5886036*PED	May	19,	2014				
	5914332	Dec	13,	2015	DS DP U-688			
	5914332*PED	Jun	13,	2016				
	6037157	Jun	26,	2016	U-688			
	6037157*PED	Dec	26,	2016				
	6284767	Feb	15,	2016	DP U-688			
	6284767*PED	Aug	15,	2016				
	6703403	Jun	26,	2016	U-688			
	6703403*PED	Dec	26,	2016				
	7148359	Jul	19,	2019	DP			
	7148359*PED	Jan	19,	2020				
	7364752	Nov	10,	2020	DP U-688			
	7364752*PED	May	10,	2021				
	8025899	Sep	26,	2027	DP			
	8025899*PED	Mar	26,	2028				
<u>LOPINAVIR; RITONAVIR - KALETRA</u>								
N021906 002	5541206	Jul	30,	2013	DS DP U-688		D-124	Apr 27, 2013
	5541206*PED	Jan	30,	2014				
	5648497	Jul	15,	2014	DS DP			
	5648497*PED	Jan	15,	2015				
	5886036	Nov	19,	2013	DS DP U-895			
	5886036*PED	May	19,	2014				
	5914332	Dec	13,	2015	DS DP U-688			
	5914332*PED	Jun	13,	2016				
	6037157	Jun	26,	2016	U-688			
	6037157*PED	Dec	26,	2016				
	6284767	Feb	15,	2016	DP U-688			
	6284767*PED	Aug	15,	2016				
	6703403	Jun	26,	2016	U-688			
	6703403*PED	Dec	26,	2016				
	7148359	Jul	19,	2019	DP			
	7148359*PED	Jan	19,	2020				
	7364752	Nov	10,	2020	DP U-688			
	7364752*PED	May	10,	2021				
	8025899	Sep	26,	2027	DP			
	8025899*PED	Mar	26,	2028				
<u>LORATADINE - CLARITIN</u>								
N020641 002	6132758	Jun	01,	2018				
<u>LORATADINE; PSEUDOEPHEDRINE SULFATE - CLARITIN-D 24 HOUR</u>								
N020470 002	5314697	Oct	23,	2012				
<u>LOTEPREDNOL ETABONATE - ALREX</u>								
N020803 001	4996335	Mar	09,	2012				
	4996335*PED	Sep	09,	2012				
	5540930	Oct	25,	2013				
	5540930*PED	Apr	25,	2014				
	5747061	Oct	25,	2013	DP U-576			
	5747061*PED	Apr	25,	2014				
<u>LOTEPREDNOL ETABONATE - LOTEMAX</u>								
N020583 001	4996335	Mar	09,	2012				
	4996335*PED	Sep	09,	2012				
	5540930	Oct	25,	2013				
	5540930*PED	Apr	25,	2014				
	5747061	Oct	25,	2013	DP U-575			
	5747061*PED	Apr	25,	2014				
<u>LOTEPREDNOL ETABONATE - LOTEMAX</u>								
N020841 001	5540930	Oct	25,	2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LOTEPREDNOL ETABONATE - LOTEMAX</u>						
	N200738 001				NDF	Apr 15, 2014
				PED		Oct 15, 2014
<u>LOTEPREDNOL ETABONATE; TOBRAMYCIN - ZYLET</u>						
N050804 001	4996335	Mar 09, 2012	DS DP U-920			
	4996335*PED	Sep 09, 2012				
	5540930	Oct 25, 2013	DP			
	5540930*PED	Apr 25, 2014				
	5747061	Oct 25, 2013	DP U-920			
	5747061*PED	Apr 25, 2014				
<u>LOVASTATIN - ALTOPREV</u>						
N021316 001	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018	U-456			
	6485748	Dec 12, 2017	DP			
<u>LOVASTATIN - ALTOPREV</u>						
N021316 002	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018	U-456			
	6485748	Dec 12, 2017	DP			
<u>LOVASTATIN - ALTOPREV</u>						
N021316 003	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018	U-456			
	6485748	Dec 12, 2017	DP			
<u>LOVASTATIN - ALTOPREV</u>						
N021316 004	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018	U-456			
	6485748	Dec 12, 2017	DP			
<u>LOVASTATIN; NIACIN - ADVICOR</u>						
N021249 001	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-1133			
	6080428	May 27, 2017	U-447			
	6080428	May 27, 2017	U-1132			
	6129930	Sep 20, 2013	DP U-1134			
	6129930	Sep 20, 2013	DP U-1133			
	6129930	Sep 20, 2013	DP U-448			
	6129930	Sep 20, 2013	DP U-1132			
	6406715	Sep 20, 2013	DP U-450			
	6469035	Mar 15, 2018	U-1131			
	6469035	Mar 15, 2018	U-768			
	6469035	Mar 15, 2018	U-1130			
	6469035	Mar 15, 2018	U-1129			
	6676967	Sep 20, 2013	U-548			
	6676967	Sep 20, 2013	U-1133			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1136			
	7011848	Sep 20, 2013	U-1137			
	7011848	Sep 20, 2013	U-712			
	7011848	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1136			
	7998506	Sep 20, 2013	U-1137			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LOVASTATIN; NIACIN - ADVICOR</u>						
N021249 002	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-1132			
	6080428	May 27, 2017	U-1133			
	6080428	May 27, 2017	U-447			
	6129930	Sep 20, 2013	DP U-448			
	6129930	Sep 20, 2013	DP U-1132			
	6129930	Sep 20, 2013	DP U-1134			
	6129930	Sep 20, 2013	DP U-1133			
	6406715	Sep 20, 2013	DP U-450			
	6469035	Mar 15, 2018	U-1129			
	6469035	Mar 15, 2018	U-768			
	6469035	Mar 15, 2018	U-1131			
	6469035	Mar 15, 2018	U-1130			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-548			
	6676967	Sep 20, 2013	U-1133			
	6676967	Sep 20, 2013	U-1132			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-712			
	7011848	Sep 20, 2013	U-1135			
	7011848	Sep 20, 2013	U-1137			
	7011848	Sep 20, 2013	U-1136			
	7998506	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1136			
	7998506	Sep 20, 2013	U-1137			
<u>LOVASTATIN; NIACIN - ADVICOR</u>						
N021249 003	6080428	May 27, 2017	U-1133			
	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-447			
	6080428	May 27, 2017	U-1132			
	6129930	Sep 20, 2013	DP U-1133			
	6129930	Sep 20, 2013	DP U-1134			
	6129930	Sep 20, 2013	DP U-1132			
	6129930	Sep 20, 2013	DP U-448			
	6406715	Sep 20, 2013	DP U-450			
	6469035	Mar 15, 2018	U-1131			
	6469035	Mar 15, 2018	U-1129			
	6469035	Mar 15, 2018	U-768			
	6469035	Mar 15, 2018	U-1130			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-1133			
	6676967	Sep 20, 2013	U-548			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1136			
	7011848	Sep 20, 2013	U-712			
	7011848	Sep 20, 2013	U-1137			
	7011848	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1137			
	7998506	Sep 20, 2013	U-1136			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>LOVASTATIN; NIACIN - ADVICOR</u>						
N021249 004	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-1132			
	6080428	May 27, 2017	U-1133			
	6080428	May 27, 2017	U-447			
	6129930	Sep 20, 2013	DP U-1133			
	6129930	Sep 20, 2013	DP U-1132			
	6129930	Sep 20, 2013	DP U-1134			
	6129930	Sep 20, 2013	DP U-448			
	6406715	Sep 20, 2013	DP			
	6469035	Mar 15, 2018	U-1130			
	6469035	Mar 15, 2018	U-1129			
	6469035	Mar 15, 2018	U-1131			
	6469035	Mar 15, 2018	U-768			
	6676967	Sep 20, 2013	U-548			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-1133			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1137			
	7011848	Sep 20, 2013	U-712			
	7011848	Sep 20, 2013	U-1136			
	7011848	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1135			
	7998506	Sep 20, 2013	U-1136			
	7998506	Sep 20, 2013	U-1137			
<u>LUBIPROSTONE - AMITIZA</u>						
N021908 001	5284858	Jul 14, 2014	DS			
	6414016	Sep 05, 2020		U-717		
	6583174	Oct 16, 2020	DP			
	7064148	Aug 30, 2022		U-739		
	7417067	Oct 16, 2020	DP			
	8026393	Oct 25, 2027	DP			
	8071613	Sep 05, 2020		U-1203		
	8088934	May 18, 2021	DS			
<u>LUBIPROSTONE - AMITIZA</u>						
N021908 002	5284858	Jul 14, 2014	DS			
	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			
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N200603 001	5532372	Jul 02, 2013	DS		NCE	Oct 28, 2015
<u>LURASIDONE HYDROCHLORIDE - LATUDA</u>						
N200603 002	5532372	Jul 02, 2013	DS		NCE	Oct 28, 2015
<u>LUTROPIN ALFA - LUVERIS</u>						
N021322 001	5767251	Jun 16, 2015	DS		ODE	Oct 08, 2011
<u>MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 25</u>						
N021910 001	5945449	Oct 31, 2017	DP U-785		ODE	Jul 26, 2013
	7300674	Mar 04, 2023	DP U-785			
<u>MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 35</u>						
N021910 002	5945449	Oct 31, 2017	DP U-785		ODE	Jul 26, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE</u>							
N022456 001	6489346	Jul	16, 2016	DP	U-588		
	6489346	Jul	16, 2016	DP	U-1021		
	6645988	Jul	16, 2016	DP			
	6699885	Jul	16, 2016	DP	U-588		
	6699885	Jul	16, 2016	DP	U-1021		
	7399772	Jul	16, 2016	DP	U-1021		
	7399772	Jul	16, 2016	DP	U-588		
<u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE</u>							
N022456 002	6489346	Jul	16, 2016	DP	U-1021		
	6489346	Jul	16, 2016	DP	U-588		
	6645988	Jul	16, 2016	DP			
	6699885	Jul	16, 2016	DP	U-588		
	6699885	Jul	16, 2016	DP	U-1021		
	7399772	Jul	16, 2016	DP	U-588		
	7399772	Jul	16, 2016	DP	U-1021		
<u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>							
N021850 001	6489346	Jul	16, 2016	DS	DP	U-588	
	6645988	Jul	16, 2016	DS	DP	U-588	
	6699885	Jul	16, 2016			U-588	
	7399772	Jul	16, 2016			U-588	
<u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>							
N021850 002	6489346	Jul	16, 2016	DS	DP	U-623	
	6645988	Jul	16, 2016	DS	DP	U-623	
	6699885	Jul	16, 2016			U-623	
	7399772	Jul	16, 2016			U-623	
<u>MAGNESIUM SULFATE ANHYDROUS; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u>							
N022372 001	6946149	Apr	30, 2022	DP	U-837	NC	Aug 05, 2013
<u>MALATHION - OVIDE</u>							
N018613 001	7560445	Feb	01, 2027	DS	DP	U-986	
	7977324	Aug	14, 2026		DP		
<u>MANGAFODIPIR TRISODIUM - TESLASCAN</u>							
N020652 001	4933456	Nov	27, 2011				
<u>MANNITOL - ARIDOL KIT</u>							
N022368 001	5817028	Feb	23, 2015		U-1091	NP	Oct 05, 2013
<u>MARAVIROC - SELZENTRY</u>							
N022128 001	6586430	Dec	01, 2019	DS	DP	U-824	
	6667314	Aug	06, 2021	DS	DP	U-824	
	7368460	Nov	25, 2022			U-824	
	7576097	May	25, 2021	DS			
<u>MARAVIROC - SELZENTRY</u>							
N022128 002	6586430	Dec	01, 2019	DS	DP	U-824	
	6667314	Aug	06, 2021	DS	DP	U-824	
	7368460	Nov	25, 2022			U-824	
	7576097	May	25, 2021	DS			
<u>MECASERMIN RECOMBINANT - INCRELEX</u>							
N021839 001	5681814	Sep	18, 2017	DP		ODE	Aug 30, 2012
	5824642	Jul	08, 2014		U-681		
	6207640	Apr	07, 2014		U-681		
<u>MECASERMIN RINFABATE RECOMBINANT - IPLEX</u>							
N021884 001	5681818	Oct	28, 2014	U-697		ODE	Dec 12, 2012
<u>MEDROXYPROGESTERONE ACETATE - DEPO-SUBQ PROVERA 104</u>							
N021583 001	6495534	May	15, 2020	DP			
<u>MEGESTROL ACETATE - MEGACE</u>							
N020264 001	5338732	Aug	16, 2011				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>MEGESTROL ACETATE - MEGACE ES</u>						
N021778 001	6592903 7101576	Sep 21, 2020 Apr 22, 2024	DP	U-755		
<u>MELOXICAM - MOBIC</u>						
N020938 001					ODE PED	Aug 11, 2012 Feb 11, 2013
<u>MELOXICAM - MOBIC</u>						
N020938 002					ODE PED	Aug 11, 2012 Feb 11, 2013
<u>MELOXICAM - MOBIC</u>						
N021530 001	6184220 6184220*PED	Mar 25, 2019 Sep 25, 2019	DP		ODE PED	Aug 11, 2012 Feb 11, 2013
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>						
N021487 001	5061703	Apr 11, 2015	U-539			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>						
N021487 002	5061703	Apr 11, 2015	U-539			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>						
N021627 001	5061703	Apr 11, 2015	U-539			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N022525 001	5061703 8039009	Apr 11, 2015 Mar 24, 2029	U-539 U-539		NDF	Jun 21, 2013
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N022525 002	5061703 8039009	Apr 11, 2015 Mar 24, 2029	U-539 U-539		NDF	Jun 21, 2013
<u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u>						
N022525 003	5061703 8039009	Apr 11, 2015 Mar 24, 2029	U-539 U-539		NDF	Jun 21, 2013
<u>MEQUINOL; TRETINOIN - SOLAGE</u>						
N020922 001	5194247 6353029	Dec 10, 2013 Aug 24, 2020	DP	U-294		
<u>MESALAMINE - APRISO</u>						
N022301 001	6551620	Apr 20, 2018	DS	DP	U-907	NP
<u>MESALAMINE - ASACOL</u>						
N019651 001	5541170 5541171	Jul 30, 2013 Jul 30, 2013		U-141 U-141		
<u>MESALAMINE - ASACOL HD</u>						
N021830 001	5541170 5541171 6893662	Jul 30, 2013 Jul 30, 2013 Nov 15, 2021	DP	U-141 U-141 U-141		
<u>MESALAMINE - LIALDA</u>						
N022000 001	6773720	Jun 08, 2020	DP		I-640	Jul 14, 2014
<u>MESALAMINE - SFROWASA</u>						
N019618 002	7645801	Jul 24, 2027	DS	DP		
<u>MESNA - MESNEX</u>						
N020855 001	5252341 5262169	Jul 16, 2011 Jul 16, 2011				
<u>METAXALONE - SKELAXIN</u>						
N013217 001	6407128 6683102	Dec 03, 2021 Dec 03, 2021	U-189 U-189			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
METAXALONE - SKELAXIN						
N013217 003	6407128	Dec 03, 2021		U-189		
	6683102	Dec 03, 2021		U-189		
	7122566	Feb 06, 2026		U-915		
	7714006	Dec 03, 2021		U-1050		
METFORMIN HYDROCHLORIDE - FORTAMET						
N021574 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			
METFORMIN HYDROCHLORIDE - FORTAMET						
N021574 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			
METFORMIN HYDROCHLORIDE - GLUCOPHAGE XR						
N021202 001	6475521	Mar 19, 2018				
	6660300	Mar 19, 2018		U-542		
METFORMIN HYDROCHLORIDE - GLUCOPHAGE XR						
N021202 004	6475521	Mar 19, 2018				
	6660300	Mar 19, 2018		U-542		
METFORMIN HYDROCHLORIDE - GLUMETZA						
N021748 001	6340475	Sep 19, 2016	DS DP	U-669		
	6488962	Jun 20, 2020	DS DP			
	6635280	Sep 19, 2016	DS DP			
	6723340	Oct 25, 2021	DS DP			
METFORMIN HYDROCHLORIDE - GLUMETZA						
N021748 002	6488962	Jun 20, 2020	DS DP			
	7780987	Mar 23, 2025	DS DP			
METFORMIN HYDROCHLORIDE - METFORMIN HYDROCHLORIDE						
A090692 001					PC	Mar 28, 2012
METFORMIN HYDROCHLORIDE - METFORMIN HYDROCHLORIDE						
A090692 002					PC	Mar 28, 2012
METFORMIN HYDROCHLORIDE - RIOMET						
N021591 001	6890957	Sep 14, 2023	DP			
METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET						
N021842 001	5965584	Jun 19, 2016	DP U-1055			
	6166042	Jun 19, 2016		U-679		
	6166043	Jun 19, 2016		U-679		
	6172090	Jun 19, 2016		U-679		
METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET						
N021842 002	5965584	Jun 19, 2016	DP U-1055			
	6166042	Jun 19, 2016		U-679		
	6166043	Jun 19, 2016		U-679		
	6172090	Jun 19, 2016		U-679		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N022024 001	5965584	Jun 19, 2016	DP U-973			
	6099859	Mar 20, 2018	DP			
	6166042	Jun 19, 2016		U-973		
	6166043	Jun 19, 2016		U-973		
	6172090	Jun 19, 2016		U-973		
	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-973		
	7919116	Mar 20, 2018		U-1120		
	7959946	Jul 31, 2026	DP			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
N022024 002	5965584	Jun 19, 2016	DP U-973			
	6099859	Mar 20, 2018	DP			
	6166042	Jun 19, 2016		U-973		
	6166043	Jun 19, 2016		U-973		
	6172090	Jun 19, 2016		U-973		
	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-973		
	7919116	Mar 20, 2018		U-1120		
	7959946	Jul 31, 2026	DP			
<u>METFORMIN HYDROCHLORIDE; REPAGLINIDE - PRANDIMET</u>						
N022386 001	6677358	Jun 12, 2018	DP U-546			
<u>METFORMIN HYDROCHLORIDE; REPAGLINIDE - PRANDIMET</u>						
N022386 002	6677358	Jun 12, 2018	DP U-546			
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N021410 001	5002953	Sep 17, 2011	DS DP U-493			
	5002953	Sep 17, 2011	DS DP U-690			
	5002953	Sep 17, 2011	DS DP U-691			
	5002953	Sep 17, 2011	DS DP U-734			
	5002953*PED	Mar 17, 2012				
	5741803	Apr 21, 2015	DS DP U-734	Y		
	5741803	Apr 21, 2015	DS DP U-493	Y		
	5741803*PED	Oct 21, 2015				
	5965584	Jun 19, 2016		U-493	Y	
	6166042	Jun 19, 2016		U-493	Y	
	6288095	Feb 11, 2017		U-493	Y	
	6288095*PED	Aug 11, 2017				
	7358366	Apr 19, 2020	DS			
	7358366*PED	Oct 19, 2020				
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>						
N021410 002	5002953	Sep 17, 2011	DS DP U-690			
	5002953	Sep 17, 2011	DS DP U-691			
	5002953	Sep 17, 2011	DS DP U-734			
	5002953	Sep 17, 2011	DS DP U-493			
	5002953*PED	Mar 17, 2012				
	5741803	Apr 21, 2015	DS DP U-493	Y		
	5741803	Apr 21, 2015	DS DP U-734	Y		
	5741803*PED	Oct 21, 2015				
	5965584	Jun 19, 2016		U-493	Y	
	6166042	Jun 19, 2016		U-493	Y	
	6288095	Feb 11, 2017		U-493	Y	
	6288095*PED	Aug 11, 2017				
	7358366	Apr 19, 2020	DS			
	7358366*PED	Oct 19, 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
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET								
N021410 003	5002953	Sep	17, 2011	DS	DP U-493			
	5002953	Sep	17, 2011	DS	DP U-691			
	5002953	Sep	17, 2011	DS	DP U-690			
	5002953	Sep	17, 2011	DS	DP U-734			
	5002953*PED	Mar	17, 2012					
	5741803	Apr	21, 2015	DS	DP U-493	Y		
	5741803	Apr	21, 2015	DS	DP U-734	Y		
	5741803*PED	Oct	21, 2015					
	5965584	Jun	19, 2016		U-493	Y		
	6166042	Jun	19, 2016		U-493	Y		
	6288095	Feb	11, 2017		U-493	Y		
	6288095*PED	Aug	11, 2017					
	7358366	Apr	19, 2020	DS				
	7358366*PED	Oct	19, 2020					
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET								
N021410 004	5002953	Sep	17, 2011	DS	DP U-690			
	5002953	Sep	17, 2011	DS	DP U-691			
	5002953	Sep	17, 2011	DS	DP U-734			
	5002953	Sep	17, 2011	DS	DP U-493			
	5002953*PED	Mar	17, 2012					
	5741803	Apr	21, 2015	DS	DP U-493	Y		
	5741803	Apr	21, 2015	DS	DP U-734	Y		
	5741803*PED	Oct	21, 2015					
	7358366	Apr	19, 2020	DS				
	7358366*PED	Oct	19, 2020					
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET								
N021410 005	5002953	Sep	17, 2011	DS	DP U-493			
	5002953	Sep	17, 2011	DS	DP U-691			
	5002953	Sep	17, 2011	DS	DP U-690			
	5002953	Sep	17, 2011	DS	DP U-734			
	5002953*PED	Mar	17, 2012					
	5741803	Apr	21, 2015	DS	DP U-493	Y		
	5741803	Apr	21, 2015	DS	DP U-734	Y		
	5741803*PED	Oct	21, 2015					
	7358366	Apr	19, 2020	DS				
	7358366*PED	Oct	19, 2020					
METFORMIN HYDROCHLORIDE; SAXAGLIPTIN - KOMBIGLYZE XR								
N200678 001	6395767	Feb	16, 2021	DS	DP U-1097		NCE	Jul 31, 2014
METFORMIN HYDROCHLORIDE; SAXAGLIPTIN - KOMBIGLYZE XR								
N200678 002	6395767	Feb	16, 2021	DS	DP U-1097		NCE	Jul 31, 2014
METFORMIN HYDROCHLORIDE; SAXAGLIPTIN - KOMBIGLYZE XR								
N200678 003	6395767	Feb	16, 2021	DS	DP U-1097		NCE	Jul 31, 2014
METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET								
N022044 001	6303661	Apr	24, 2017		U-802		NCE	Oct 16, 2011
	6699871	Jul	26, 2022	DS	DP U-802			
	6890898	Feb	02, 2019		U-803			
	6890898	Feb	02, 2019		U-1036			
	6890898	Feb	02, 2019		U-1038			
	7078381	Feb	02, 2019		U-803			
	7078381	Feb	02, 2019		U-1038			
	7078381	Feb	02, 2019		U-1036			
	7125873	Jul	26, 2022	DP	U-1038			
	7125873	Jul	26, 2022	DP	U-1036			
	7125873	Jul	26, 2022	DP	U-803			
	7326708	Apr	11, 2026	DS	DP U-802			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>							
N022044 002	6303661	Apr	24, 2017		U-802		
	6699871	Jul	26, 2022	DS DP	U-802		
	6890898	Feb	02, 2019		U-803		
	6890898	Feb	02, 2019		U-1038		
	6890898	Feb	02, 2019		U-1036		
	7078381	Feb	02, 2019		U-1038		
	7078381	Feb	02, 2019		U-1036		
	7078381	Feb	02, 2019		U-803		
	7125873	Jul	26, 2022		DP U-1036		
	7125873	Jul	26, 2022		DP U-1038		
	7125873	Jul	26, 2022		DP U-803		
	7326708	Apr	11, 2026	DS DP	U-802		
<u>METHYL AMINOLEVULINATE HYDROCHLORIDE - METVIXIA</u>							
N021415 001	6034267	Mar	08, 2016		U-804		
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>							
N021964 001	6559158	Nov	03, 2017		U-1185		
<u>METHYLNALTREXONE BROMIDE - RELISTOR</u>							
N021964 002	6559158	Nov	03, 2017		U-1185		
<u>METHYLPHENIDATE - DAYTRANA</u>							
N021514 001	5958446	Dec	12, 2012		DP		
	6210705	Sep	30, 2018		DP U-727		
	6348211	Sep	30, 2018		DP U-727		
<u>METHYLPHENIDATE - DAYTRANA</u>							
N021514 002	5958446	Dec	12, 2012		DP		
	6210705	Sep	30, 2018		DP U-727		
	6348211	Sep	30, 2018		DP U-727		
<u>METHYLPHENIDATE - DAYTRANA</u>							
N021514 003	5958446	Dec	12, 2012		DP		
	6210705	Sep	30, 2018		DP U-727		
	6348211	Sep	30, 2018		DP U-727		
<u>METHYLPHENIDATE - DAYTRANA</u>							
N021514 004	5958446	Dec	12, 2012		DP		
	6210705	Sep	30, 2018		DP U-727		
	6348211	Sep	30, 2018		DP U-727		
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>							
N021121 001	6919373	Jul	31, 2017		U-666		
	6919373*PED	Jan	31, 2018			M-88	
	6930129	Jul	31, 2017		U-666		
	6930129*PED	Jan	31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>							
N021121 002	6919373	Jul	31, 2017		U-666		
	6919373*PED	Jan	31, 2018			M-88	
	6930129	Jul	31, 2017		U-666		
	6930129*PED	Jan	31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>							
N021121 003	6919373	Jul	31, 2017		U-666		
	6919373*PED	Jan	31, 2018			M-88	
	6930129	Jul	31, 2017		U-666		
	6930129*PED	Jan	31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</u>							
N021121 004	6919373	Jul	31, 2017		U-666		
	6919373*PED	Jan	31, 2018			M-88	
	6930129	Jul	31, 2017		U-666		
	6930129*PED	Jan	31, 2018				
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>							
N021259 001	6344215	Oct	27, 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
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N021259 002	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N021259 003	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>						
N021259 004	6344215	Oct 27, 2020	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u>						
N021419 001	7691880	Oct 07, 2024	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</u>						
N021419 002	7691880	Oct 07, 2024	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u>						
N021284 001	5837284	Dec 04, 2015	DP			
	6228398	Nov 01, 2019	DP	U-472		
	6635284	Dec 04, 2015	DP	U-591		
	7431944	Dec 04, 2015	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u>						
N021284 002	5837284	Dec 04, 2015	DP			
	6228398	Nov 01, 2019	DP	U-472		
	6635284	Dec 04, 2015	DP	U-591		
	7431944	Dec 04, 2015	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u>						
N021284 003	5837284	Dec 04, 2015	DP			
	6228398	Nov 01, 2019	DP	U-472		
	6635284	Dec 04, 2015	DP	U-591		
	7431944	Dec 04, 2015	DP			
<u>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</u>						
N021284 004	5837284	Dec 04, 2015	DP			
	6228398	Nov 01, 2019	DP	U-472		
	6635284	Dec 04, 2015	DP	U-591		
	7431944	Dec 04, 2015	DP			
<u>METOCLOPRAMIDE HYDROCHLORIDE - METOZOLV ODT</u>						
N022246 001	6413549	Jul 11, 2017	DP			
<u>METOCLOPRAMIDE HYDROCHLORIDE - METOZOLV ODT</u>						
N022246 002	6413549	Jul 11, 2017	DP			
<u>METOCLOPRAMIDE HYDROCHLORIDE - REGLAN ODT</u>						
N021793 001	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<u>METOCLOPRAMIDE HYDROCHLORIDE - REGLAN ODT</u>						
N021793 002	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<u>METRONIDAZOLE - FLAGYL ER</u>						
N020868 001	6103262	Aug 15, 2017	DP	U-137		
<u>METRONIDAZOLE - METROGEL</u>						
N021789 001	6881726	Feb 21, 2022	DP	U-743		
	7348317	Feb 21, 2022	DP	U-743		
<u>METRONIDAZOLE - METROGEL-VAGINAL</u>						
N020208 001	5536743	Jul 16, 2013		U-137		
<u>METRONIDAZOLE - VANDAZOLE</u>						
N021806 001	7456207	Sep 22, 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
MICAFUNGIN SODIUM - MYCAMEINE							
N021506 002	5376634	Dec	27, 2011	DS DP			
	6107458	Mar	16, 2019	DS DP	U-650		
	6107458	Mar	16, 2019	DS DP	U-845		
	6265536	Sep	29, 2015	DS DP	U-650		
	6265536	Sep	29, 2015	DS DP	U-845		
	6774104	Jan	08, 2021	DP	U-845		
	6774104	Jan	08, 2021	DP	U-650		
MICAFUNGIN SODIUM - MYCAMEINE							
N021506 003	5376634	Dec	27, 2011	DS DP			
	6107458	Mar	16, 2019	DS DP	U-845		
	6107458	Mar	16, 2019	DS DP	U-650		
	6265536	Sep	29, 2015	DS DP	U-845		
	6265536	Sep	29, 2015	DS DP	U-650		
	6774104	Jan	08, 2021	DP	U-845		
	6774104	Jan	08, 2021	DP	U-650		
MICONAZOLE - ORAVIG							
N022404 001	6916485	Sep	11, 2022	DP	U-1051		
	7651698	Jan	08, 2026		U-1051		
MICONAZOLE NITRATE - MONISTAT 1 COMBINATION PACK							
N021308 001	5514698	Mar	21, 2014			Y	
	6153635	Nov	28, 2020			Y	
MIGLUSTAT - ZAVESCA							
N021348 001	5472969	May	13, 2013				
	5525616	Jun	11, 2013				
MILNACIPRAN HYDROCHLORIDE - SAVELLA							
N022256 001	6602911	Nov	05, 2021	U-882			
	6992110	Nov	05, 2021	U-882			
	7888342	Nov	05, 2021	U-882			
	7994220	Sep	19, 2029	U-819			
MILNACIPRAN HYDROCHLORIDE - SAVELLA							
N022256 002	6602911	Nov	05, 2021	U-882			
	6992110	Nov	05, 2021	U-882			
	7888342	Nov	05, 2021	U-882			
	7994220	Sep	19, 2029	U-819			
MILNACIPRAN HYDROCHLORIDE - SAVELLA							
N022256 003	6602911	Nov	05, 2021	U-882			
	6992110	Nov	05, 2021	U-882			
	7888342	Nov	05, 2021	U-882			
	7994220	Sep	19, 2029	U-819			
MILNACIPRAN HYDROCHLORIDE - SAVELLA							
N022256 004	6602911	Nov	05, 2021	U-882			
	6992110	Nov	05, 2021	U-882			
	7888342	Nov	05, 2021	U-882			
	7994220	Sep	19, 2029	U-819			
MINOCYCLINE HYDROCHLORIDE - SOLODYN							
N050808 001	5908838	Feb	19, 2018	U-917			
	7790705	Jun	24, 2025	U-1078			
	7919483	Mar	07, 2027	U-1078			
MINOCYCLINE HYDROCHLORIDE - SOLODYN							
N050808 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			
MINOCYCLINE HYDROCHLORIDE - SOLODYN							
N050808 003	5908838	Feb	19, 2018	U-917			
	7790705	Jun	24, 2025	U-1078			
	7919483	Mar	07, 2027	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
MINOCYCLINE HYDROCHLORIDE - SOLODYN						
N050808 004	5908838	Feb 19, 2018		U-917		
	7790705	Jun 24, 2025		U-1078		
	7919483	Mar 07, 2027		U-1078		
MINOCYCLINE HYDROCHLORIDE - SOLODYN						
N050808 005	5908838	Feb 19, 2018		U-917		
	7790705	Jun 24, 2025		U-1078		
	7919483	Mar 07, 2027		U-1078		
MINOCYCLINE HYDROCHLORIDE - SOLODYN						
N050808 006	5908838	Feb 19, 2018		U-917		
	7790705	Jun 24, 2025		U-1078		
	7919483	Mar 07, 2027		U-1078		
MINOCYCLINE HYDROCHLORIDE - SOLODYN						
N050808 007	5908838	Feb 19, 2018		U-917		
	7790705	Jun 24, 2025		U-1078		
	7919483	Mar 07, 2027		U-1078		
MINOCYCLINE HYDROCHLORIDE - SOLODYN						
N050808 008	5908838	Feb 19, 2018		U-917		
	7790705	Jun 24, 2025		U-1078		
	7919483	Mar 07, 2027		U-1078		
MINOXIDIL - MEN'S ROGAINE						
N021812 001	6946120	Apr 20, 2019		DP	U-702	
MODAFINIL - PROVIGIL						
N020717 001	7297346	Nov 29, 2023		DP		
	7297346*PED	May 29, 2024				
	RE37516	Oct 06, 2014			U-255	
	RE37516*PED	Apr 06, 2015				
MODAFINIL - PROVIGIL						
N020717 002	7297346	Nov 29, 2023		DP		
	7297346*PED	May 29, 2024			U-255	
	RE37516	Oct 06, 2014				
	RE37516*PED	Apr 06, 2015				
MOMETASONE FUROATE - ASMANEX TWISTHALER						
N021067 001	5394868	Jun 25, 2012		DP		
	5394868*PED	Dec 25, 2012				
	5687710	Nov 18, 2014		DP		
	5687710*PED	May 18, 2015				
	5829434	Nov 03, 2015		DP		
	5829434*PED	May 03, 2016				
	5889015	Jan 27, 2014			U-645	
	5889015*PED	Jul 27, 2014				
	6057307	Jan 27, 2014		DP	U-645	
	6057307*PED	Jul 27, 2014				
	6240918	Feb 20, 2017		DP		
	6240918*PED	Aug 20, 2017				
	6365581	Jan 27, 2014			U-645	
	6365581*PED	Jul 27, 2014				
	6503537	Mar 17, 2018		DP		
	6503537*PED	Sep 17, 2018				
	6677322	Jan 27, 2014			U-645	
	6677322*PED	Jul 27, 2014				
	6949532	Jan 27, 2014			U-645	
	6949532*PED	Jul 27, 2014				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
MOMETASONE FUROATE - ASMANEX TWISTHALER						
N021067 002	5394868	Jun 25, 2012	DP			
	5394868*PED	Dec 25, 2012				
	5687710	Nov 18, 2014	DP			
	5687710*PED	May 18, 2015				
	5829434	Nov 03, 2015	DP			
	5829434*PED	May 03, 2016				
	5889015	Jan 27, 2014		U-645		
	5889015*PED	Jul 27, 2014				
	6057307	Jan 27, 2014	DP	U-645		
	6057307*PED	Jul 27, 2014				
	6240918	Feb 20, 2017	DP			
	6240918*PED	Aug 20, 2017				
	6365581	Jan 27, 2014		U-645		
	6365581*PED	Jul 27, 2014				
	6503537	Mar 17, 2018	DP			
	6503537*PED	Sep 17, 2018				
	6677322	Jan 27, 2014		U-645		
	6677322*PED	Jul 27, 2014				
	6949532	Jan 27, 2014		U-645		
	6949532*PED	Jul 27, 2014				
MOMETASONE FUROATE MONOHYDRATE - NASONEX						
N020762 001	5837699	Jan 27, 2014	DP	U-625	I-626	May 26, 2013
	5837699*PED	Jul 27, 2014			M-99	Jan 19, 2014
	6127353	Oct 03, 2017	DS	DP		
	6127353*PED	Apr 03, 2018				
	6723713	Jan 27, 2014		U-625		
	6723713*PED	Jul 27, 2014				
MONTELUKAST SODIUM - SINGULAIR						
N020829 002	5565473	Feb 03, 2012	DS	DP	U-807	
	5565473	Feb 03, 2012	DS	DP	U-228	
	5565473	Feb 03, 2012	DS	DP	U-675	
	5565473*PED	Aug 03, 2012			U-228	
MONTELUKAST SODIUM - SINGULAIR						
N020830 001	5565473	Feb 03, 2012	DS	DP	U-675	
	5565473	Feb 03, 2012	DS	DP	U-807	
	5565473	Feb 03, 2012	DS	DP	U-228	
	5565473*PED	Aug 03, 2012			U-228	
MONTELUKAST SODIUM - SINGULAIR						
N020830 002	5565473	Feb 03, 2012	DS	DP	U-228	
	5565473	Feb 03, 2012	DS	DP	U-675	
	5565473	Feb 03, 2012	DS	DP	U-807	
	5565473*PED	Aug 03, 2012			U-228	
MONTELUKAST SODIUM - SINGULAIR						
N021409 001	5565473	Feb 03, 2012	DS	DP	U-675	
	5565473	Feb 03, 2012	DS	DP	U-807	
	5565473*PED	Aug 03, 2012				
	8007830	Oct 24, 2022		DP		
MORPHINE SULFATE - AVINZA						
N021260 001	6066339	Nov 25, 2017				
MORPHINE SULFATE - AVINZA						
N021260 002	6066339	Nov 25, 2017				
MORPHINE SULFATE - AVINZA						
N021260 003	6066339	Nov 25, 2017				
MORPHINE SULFATE - AVINZA						
N021260 004	6066339	Nov 25, 2017				
MORPHINE SULFATE - AVINZA						
N021260 005	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
<u>MORPHINE SULFATE - AVINZA</u>						
	N021260 006	6066339	Nov 25, 2017	DP		
<u>MORPHINE SULFATE - DEPODUR</u>						
N021671 001	5723147	Mar 03, 2015	DP	U-584		
	5807572	Sep 15, 2015	DP			
	5891467	Jan 31, 2017	DP			
	5931809	Jul 14, 2015		U-584		
	5962016	Jan 31, 2017	DP	U-584		
	5997899	Sep 01, 2016	DP			
	6171613	Oct 01, 2016	DP			
	6193998	Sep 01, 2016	DP			
	6241999	Sep 01, 2016	DP			
<u>MORPHINE SULFATE - DEPODUR</u>						
N021671 002	5723147	Mar 03, 2015	DP	U-584		
	5807572	Sep 15, 2015	DP			
	5891467	Jan 31, 2017	DP			
	5931809	Jul 14, 2015		U-584		
	5962016	Jan 31, 2017	DP	U-584		
	5997899	Sep 01, 2016	DP			
	6171613	Oct 01, 2016	DP			
	6193998	Sep 01, 2016	DP			
	6241999	Sep 01, 2016	DP			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 001	7682633	Jun 19, 2027		U-443	NC	Aug 13, 2012
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 002	7682633	Jun 19, 2027		U-443	NC	Aug 13, 2012
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 003	7682633	Jun 19, 2027		U-443	NC	Aug 13, 2012
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 004	7682633	Jun 19, 2027		U-443	NC	Aug 13, 2012
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 005	7682633	Jun 19, 2027		U-443	NC	Aug 13, 2012
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
<u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u>						
N022321 006	7682633	Jun 19, 2027		U-443	NC	Aug 13, 2012
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 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
<u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX</u>								
N021085 001	4990517	Dec	08,	2011	DS	DP	U-298	
	5607942	Mar	04,	2014			U-298	
	5849752	Dec	05,	2016			U-298	
	6610327	Oct	29,	2019	DP		U-298	
<u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u>								
N021277 001	4990517	Dec	08,	2011	DS	DP	U-298	
	5607942	Mar	04,	2014			U-298	
	5849752	Dec	05,	2016			U-298	
	6548079	Jul	25,	2020	DP		U-298	
<u>MOXIFLOXACIN HYDROCHLORIDE - MOXEZA</u>								
N022428 001	4990517	Dec	08,	2011	DS	DP	U-709	
	4990517*PED	Jun	08,	2012				NP
	5607942	Mar	04,	2014	DS	DP	U-709	PED
	5607942*PED	Sep	04,	2014				May 19, 2014
	6716830	Sep	29,	2019	DP			
	6716830*PED	Mar	29,	2020				
	7671070	Sep	29,	2019	DP		U-709	
	7671070*PED	Mar	29,	2020				
<u>MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX</u>								
N021598 001	4990517	Dec	08,	2011	DS	DP	U-709	
	4990517*PED	Jun	08,	2012				
	5607942	Mar	04,	2014	DS	DP	U-709	
	5607942*PED	Sep	04,	2014				
	6716830	Sep	29,	2019	DP			
	6716830*PED	Mar	29,	2020				
	7671070	Sep	29,	2019			U-709	
	7671070*PED	Mar	29,	2020				
<u>MUPIROCIN - CENTANY</u>								
N050788 001	6013657	Jul	08,	2018	DP			
<u>MUPIROCIN CALCIUM - BACTROBAN</u>								
N050746 001	6025389	Oct	20,	2014	DP		U-1122	
<u>MYCOPHENOLATE MOFETIL - CELLCEPT</u>								
N050759 001	5688529	Nov	18,	2014	DP			
<u>MYCOPHENOLATE MOFETIL HYDROCHLORIDE - CELLCEPT</u>								
N050758 001	5543408	Sep	15,	2013	DP			
<u>MYCOPHENOLIC ACID - MYFORTIC</u>								
N050791 001	6025391	Apr	10,	2017	DP		U-908	
	6172107	Apr	10,	2017	DP		U-908	
	6306900	Apr	10,	2017	DP			
<u>MYCOPHENOLIC ACID - MYFORTIC</u>								
N050791 002	6025391	Apr	10,	2017	DP		U-908	
	6172107	Apr	10,	2017	DP		U-908	
	6306900	Apr	10,	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
NALTREXONE - VIVITROL						
N021897 001	5792477	May 02, 2017	DP		I-631	Oct 12, 2013
	5916598	May 02, 2017	DP			
	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			
	6403114	May 02, 2017	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-1124		
	7919499	Oct 15, 2029		U-1123		
NAPROXEN SODIUM - NAPRELAN						
N020353 001	5637320	Jun 10, 2014				
NAPROXEN SODIUM - NAPRELAN						
N020353 002	5637320	Jun 10, 2014				
NAPROXEN SODIUM - NAPRELAN						
N020353 003	5637320	Jun 10, 2014				
NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET						
N021926 001	6060499	Aug 14, 2017	DP	U-867		
	6586458	Aug 14, 2017	DP	U-867		
	7332183	Oct 02, 2025	DP	U-867		
	8022095	Aug 14, 2017	DP	U-867		
NATEGLINIDE - STARLIX						
N021204 001	5463116	Oct 21, 2012				
	5488150	Jan 30, 2013				
	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			
NATEGLINIDE - STARLIX						
N021204 002	5463116	Oct 21, 2012				
	5488150	Jan 30, 2013				
	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			
NEBIVOLOL HYDROCHLORIDE - BYSTOLIC						
N021742 002	5759580	Jun 02, 2015	DP		NCE	
	6545040	Dec 17, 2021	DP	U-3		Dec 17, 2012
NEBIVOLOL HYDROCHLORIDE - BYSTOLIC						
N021742 003	5759580	Jun 02, 2015	DP		NCE	
	6545040	Dec 17, 2021	DP	U-3		Dec 17, 2012
NEBIVOLOL HYDROCHLORIDE - BYSTOLIC						
N021742 004	5759580	Jun 02, 2015	DP		NCE	
	6545040	Dec 17, 2021	DP	U-3		Dec 17, 2012
NEBIVOLOL HYDROCHLORIDE - BYSTOLIC						
N021742 005	5759580	Jun 02, 2015	DP		NCE	
	6545040	Dec 17, 2021	DP	U-3		Dec 17, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>NEDOCROMIL SODIUM - ALOCRIL</u>						
	N021009 001 RE38628	Aug 22, 2012		U-304		
<u>NELARABINE - ARRANON</u>						
	5424295	Jun 13, 2017	DS DP		ODE	Oct 28, 2012
	5492897	Feb 20, 2013		U-689		
	5747472	Feb 20, 2013		U-695		
	5747472	Feb 20, 2013		U-696		
	5747472	Feb 20, 2013		U-689		
	5821236	Feb 20, 2013		U-695		
<u>NELFINAVIR MESYLATE - VIRACEPT</u>						
	N020778 001 5484926	Oct 07, 2013				
	5484926*PED	Apr 07, 2014				
	5952343	Oct 07, 2013		U-257		
	5952343*PED	Apr 07, 2014		U-257		
	6162812	Oct 07, 2013		U-248		
	6162812*PED	Apr 07, 2014		U-248		
<u>NELFINAVIR MESYLATE - VIRACEPT</u>						
	N020779 001 5484926	Oct 07, 2013				
	5484926*PED	Apr 07, 2014				
	5952343	Oct 07, 2013		U-257		
	5952343*PED	Apr 07, 2014		U-257		
	6162812	Oct 07, 2013		U-248		
	6162812*PED	Apr 07, 2014		U-248		
<u>NELFINAVIR MESYLATE - VIRACEPT</u>						
	N021503 001 5484926	Oct 07, 2013				
	5484926*PED	Apr 07, 2014				
	5952343	Oct 07, 2013		U-257		
	5952343*PED	Apr 07, 2014		U-257		
	6162812	Oct 07, 2013		U-248		
	6162812*PED	Apr 07, 2014		U-248		
<u>NEPAFENAC - NEVANAC</u>						
	N021862 001 5475034	Jun 06, 2014		U-100		
	7834059	Jan 31, 2027		U-1095		
	8071648	Dec 02, 2025	DP			
<u>NESIRITIDE RECOMBINANT - NATRECOR</u>						
	N020920 001 5114923	May 19, 2014	DS DP	U-855		
<u>NEVIRAPINE - VIRAMUNE</u>						
	N020636 001 5366972	Nov 22, 2011		U-167		
	5366972*PED	May 22, 2012				
<u>NEVIRAPINE - VIRAMUNE XR</u>						
	N020933 001 5366972	Nov 22, 2011	DS DP	U-167		
	5366972*PED	May 22, 2012				
<u>NIACIN - NIASPAN</u>						
	N020381 001 6080428	May 27, 2017		U-331		
	6129930	Sep 20, 2013		U-354		
	6406715	Sep 20, 2013		U-450		
	6746691	Sep 20, 2013		U-586		
	6818229	Feb 15, 2014	DP			
	7011848	Sep 20, 2013		U-712		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>NIACIN - NIASPIN</u>						
N020381 002	6080428	May 27, 2017	U-1138			
	6080428	May 27, 2017	U-1139			
	6080428	May 27, 2017	U-1140			
	6080428	May 27, 2017	U-331			
	6080428	May 27, 2017	U-1141			
	6129930	Sep 20, 2013	DP U-1141			
	6129930	Sep 20, 2013	DP U-1140			
	6129930	Sep 20, 2013	DP U-1138			
	6129930	Sep 20, 2013	DP U-354			
	6129930	Sep 20, 2013	DP U-1139			
	6406715	Sep 20, 2013	DP U-450			
	6469035	Mar 15, 2018	U-1142			
	6469035	Mar 15, 2018	U-1145			
	6469035	Mar 15, 2018	U-768			
	6469035	Mar 15, 2018	U-1144			
	6469035	Mar 15, 2018	U-1143			
	6676967	Sep 20, 2013	U-1139			
	6676967	Sep 20, 2013	U-1140			
	6676967	Sep 20, 2013	U-1138			
	6676967	Sep 20, 2013	U-548			
	6676967	Sep 20, 2013	U-1146			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1148			
	7011848	Sep 20, 2013	U-712			
	7011848	Sep 20, 2013	U-1147			
	7011848	Sep 20, 2013	U-1140			
	7011848	Sep 20, 2013	U-1141			
	7998506	Sep 20, 2013	U-1140			
	7998506	Sep 20, 2013	U-1138			
	7998506	Sep 20, 2013	U-1141			
	7998506	Sep 20, 2013	U-1139			
<u>NIACIN - NIASPIN</u>						
N020381 003	6080428	May 27, 2017	U-1140			
	6080428	May 27, 2017	U-1138			
	6080428	May 27, 2017	U-331			
	6080428	May 27, 2017	U-1139			
	6080428	May 27, 2017	U-1141			
	6129930	Sep 20, 2013	DP U-354			
	6129930	Sep 20, 2013	DP U-1138			
	6129930	Sep 20, 2013	DP U-1141			
	6129930	Sep 20, 2013	DP U-1139			
	6129930	Sep 20, 2013	DP U-1140			
	6406715	Sep 20, 2013	DP U-450			
	6469035	Mar 15, 2018	U-1144			
	6469035	Mar 15, 2018	U-768			
	6469035	Mar 15, 2018	U-1145			
	6469035	Mar 15, 2018	U-1143			
	6469035	Mar 15, 2018	U-1142			
	6676967	Sep 20, 2013	U-1140			
	6676967	Sep 20, 2013	U-1141			
	6676967	Sep 20, 2013	U-1146			
	6676967	Sep 20, 2013	U-1138			
	6676967	Sep 20, 2013	U-548			
	6676967	Sep 20, 2013	U-1139			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1148			
	7011848	Sep 20, 2013	U-712			
	7011848	Sep 20, 2013	U-1147			
	7011848	Sep 20, 2013	U-1140			
	7011848	Sep 20, 2013	U-1141			
	7998506	Sep 20, 2013	U-1138			
	7998506	Sep 20, 2013	U-1140			
	7998506	Sep 20, 2013	U-1139			
	7998506	Sep 20, 2013	U-1141			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>NIACIN - NIASPAN</u>						
N020381 004	6080428	May 27, 2017	U-1140			
	6080428	May 27, 2017	U-331			
	6080428	May 27, 2017	U-1141			
	6080428	May 27, 2017	U-1139			
	6080428	May 27, 2017	U-1138			
	6129930	Sep 20, 2013	DP U-1139			
	6129930	Sep 20, 2013	DP U-354			
	6129930	Sep 20, 2013	DP U-1138			
	6129930	Sep 20, 2013	DP U-1141			
	6129930	Sep 20, 2013	DP U-1140			
	6406715	Sep 20, 2013	DP U-450			
	6469035	Mar 15, 2018	U-1143			
	6469035	Mar 15, 2018	U-768			
	6469035	Mar 15, 2018	U-1144			
	6469035	Mar 15, 2018	U-1145			
	6469035	Mar 15, 2018	U-1142			
	6676967	Sep 20, 2013	U-1140			
	6676967	Sep 20, 2013	U-548			
	6676967	Sep 20, 2013	U-1141			
	6676967	Sep 20, 2013	U-1138			
	6676967	Sep 20, 2013	U-1139			
	6676967	Sep 20, 2013	U-1146			
	6746691	Sep 20, 2013	DP U-586			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1147			
	7011848	Sep 20, 2013	U-1148			
	7011848	Sep 20, 2013	U-1141			
	7011848	Sep 20, 2013	U-1140			
	7011848	Sep 20, 2013	U-712			
	7998506	Sep 20, 2013	U-1140			
	7998506	Sep 20, 2013	U-1139			
	7998506	Sep 20, 2013	U-1141			
	7998506	Sep 20, 2013	U-1138			
<u>NIACIN - NIASPAN TITRATION STARTER PACK</u>						
N020381 005	6080428	May 27, 2017	U-331			
	6129930	Sep 20, 2013	U-354			
	6406715	Sep 20, 2013	U-450			
	6746691	Sep 20, 2013	U-586			
	7011848	Sep 20, 2013	U-712			
<u>NIACIN; SIMVASTATIN - SIMCOR</u>						
N022078 001	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-1132			
	6080428	May 27, 2017	U-862			
	6129930	Sep 20, 2013	DP U-1132			
	6129930	Sep 20, 2013	DP U-862			
	6129930	Sep 20, 2013	DP U-1134			
	6406715	Sep 20, 2013	DP			
	6469035	Mar 15, 2018	U-1149			
	6469035	Mar 15, 2018	U-863			
	6469035	Mar 15, 2018	U-1129			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-862			
	6746691	Sep 20, 2013	DP			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-862			
	7011848	Sep 20, 2013	U-1150			
	7011848	Sep 20, 2013	U-1151			
	7998506	Sep 20, 2013	U-1151			
	7998506	Sep 20, 2013	U-1150			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>NIACIN; SIMVASTATIN - SIMCOR</u>						
N022078 002	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-1132			
	6080428	May 27, 2017	U-862			
	6129930	Sep 20, 2013	DP U-862			
	6129930	Sep 20, 2013	DP U-1134			
	6129930	Sep 20, 2013	DP U-1132			
	6406715	Sep 20, 2013	DP			
	6469035	Mar 15, 2018	U-1129			
	6469035	Mar 15, 2018	U-863			
	6469035	Mar 15, 2018	U-1149			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-862			
	6746691	Sep 20, 2013	DP			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-1150			
	7011848	Sep 20, 2013	U-862			
	7011848	Sep 20, 2013	U-1151			
	7998506	Sep 20, 2013	U-1150			
	7998506	Sep 20, 2013	U-1151			
<u>NIACIN; SIMVASTATIN - SIMCOR</u>						
N022078 003	6080428	May 27, 2017	U-862			
	6080428	May 27, 2017	U-1132			
	6080428	May 27, 2017	U-1134			
	6129930	Sep 20, 2013	DP U-1132			
	6129930	Sep 20, 2013	DP U-862			
	6129930	Sep 20, 2013	DP U-1134			
	6406715	Sep 20, 2013	DP			
	6469035	Mar 15, 2018	U-1149			
	6469035	Mar 15, 2018	U-1129			
	6469035	Mar 15, 2018	U-863			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-862			
	6746691	Sep 20, 2013	DP			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-862			
	7011848	Sep 20, 2013	U-1151			
	7011848	Sep 20, 2013	U-1150			
	7998506	Sep 20, 2013	U-1150			
	7998506	Sep 20, 2013	U-1151			
<u>NIACIN; SIMVASTATIN - SIMCOR</u>						
N022078 004	6080428	May 27, 2017	U-1132			
	6080428	May 27, 2017	U-862			
	6080428	May 27, 2017	U-1134			
	6129930	Sep 20, 2013	DP U-862			
	6406715	Sep 20, 2013	DP			
	6469035	Mar 15, 2018	U-863			
	6469035	Mar 15, 2018	U-1129			
	6469035	Mar 15, 2018	U-1149			
	6676967	Sep 20, 2013	U-1132			
	6676967	Sep 20, 2013	U-1134			
	6676967	Sep 20, 2013	U-862			
	6746691	Sep 20, 2013	DP			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013	U-862			
	7011848	Sep 20, 2013	U-1151			
	7011848	Sep 20, 2013	U-1150			
	7998506	Sep 20, 2013	U-1151			
	7998506	Sep 20, 2013	U-1150			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>NIACIN; SIMVASTATIN - SIMCOR</u>						
N022078 005	6080428	May 27, 2017	U-862			
	6080428	May 27, 2017	U-1134			
	6080428	May 27, 2017	U-1132			
	6129930	Sep 20, 2013	DP	U-862		
	6406715	Sep 20, 2013	DP			
	6469035	Mar 15, 2018		U-1149		
	6469035	Mar 15, 2018		U-863		
	6469035	Mar 15, 2018		U-1129		
	6676967	Sep 20, 2013		U-1134		
	6676967	Sep 20, 2013		U-1132		
	6676967	Sep 20, 2013		U-862		
	6746691	Sep 20, 2013	DP			
	6818229	Sep 20, 2013	DP			
	7011848	Sep 20, 2013		U-1150		
	7011848	Sep 20, 2013		U-862		
	7011848	Sep 20, 2013		U-1151		
	7998506	Sep 20, 2013		U-1151		
	7998506	Sep 20, 2013		U-1150		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N019734 004	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027		U-1029		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N019734 003	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027		U-1029		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u>						
N019734 002	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027		U-1029		
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER</u>						
N019734 005	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027		U-1029		
<u>NICOTINE - NICODERM CQ</u>						
N020165 004	5508038	Apr 16, 2013				
<u>NICOTINE - NICODERM CQ</u>						
N020165 005	5508038	Apr 16, 2013				
<u>NICOTINE - NICODERM CQ</u>						
N020165 006	5508038	Apr 16, 2013				
<u>NICOTINE - NICOTROL</u>						
N020385 001	5656255	Aug 12, 2014				
<u>NILOTINIB HYDROCHLORIDE MONOHYDRATE - TASIGNA</u>						
N022068 001	7169791	Jul 04, 2023	DS DP U-836		I-627	Jun 17, 2013
					NCE	Oct 29, 2012
					ODE	Oct 29, 2014
<u>NILOTINIB HYDROCHLORIDE MONOHYDRATE - TASIGNA</u>						
N022068 002	7169791	Jul 04, 2023	DS DP U-836		I-627	Jun 17, 2014
					NCE	Oct 29, 2012
					ODE	Oct 29, 2014
<u>NISOLDIPINE - SULAR</u>						
N020356 005	5422123	Jun 06, 2012	DP			
	5626874	Nov 30, 2014	DP			
<u>NISOLDIPINE - SULAR</u>						
N020356 006	5422123	Jun 06, 2012	DP			
	5626874	Nov 30, 2014	DP			
<u>NISOLDIPINE - SULAR</u>						
N020356 007	5422123	Jun 06, 2012	DP			
	5626874	Nov 30, 2014	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>NISOLDIPINE - SULAR</u>						
	N020356 008	5422123	Jun 06, 2012	DP		
		5626874	Nov 30, 2014	DP		
<u>NITAZOXANIDE - ALINIA</u>						
	N021497 001	5387598	Feb 07, 2012	DP	U-524	
		5578621	Nov 26, 2013	DP	U-525	
		5968961	May 07, 2017	DP		
		6020353	Sep 18, 2014	DS	DP	
<u>NITAZOXANIDE - ALINIA</u>						
	N021498 001	5387598	Feb 07, 2012		U-524	
		5578621	Sep 08, 2014		U-525	
		5965590	Jul 03, 2017		U-523	
		5968961	May 07, 2017			
		6020353	Sep 08, 2014			
		6117894	May 07, 2017			
<u>NITISINONE - ORFADIN</u>						
	N021232 001	5550165	Aug 27, 2013			
<u>NITISINONE - ORFADIN</u>						
	N021232 002	5550165	Aug 27, 2013			
<u>NITISINONE - ORFADIN</u>						
	N021232 003	5550165	Aug 27, 2013			
<u>NITRIC OXIDE - INOMAX</u>						
	N020845 002	5485827	Jan 23, 2013		U-297	
		5485827*PED	Jul 23, 2013			
		5873359	Jan 23, 2013		U-297	
		5873359*PED	Jul 23, 2013			
<u>NITRIC OXIDE - INOMAX</u>						
	N020845 003	5485827	Jan 23, 2013		U-297	
		5485827*PED	Jul 23, 2013			
		5873359	Jan 23, 2013		U-297	
		5873359*PED	Jul 23, 2013			
<u>NITROGLYCERIN - NITROMIST</u>						
	N021780 001	5869082	Apr 16, 2016	DP		
<u>NITROGLYCERIN - NITROSTAT</u>						
	N021134 001	6500456	Sep 16, 2018			
<u>NITROGLYCERIN - NITROSTAT</u>						
	N021134 002	6500456	Sep 16, 2018			
<u>NITROGLYCERIN - NITROSTAT</u>						
	N021134 003	6500456	Sep 16, 2018			
<u>NITROGLYCERIN - RECTIV</u>						
	N021359 001	7189761	May 27, 2014	DP	U-1198	
						NP
						Jun 21, 2014
<u>NIZATIDINE - AXID</u>						
	N021494 001	6930119	Jul 17, 2022	DP		
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>						
	N019667 001	5753618	May 19, 2015			
		5753618*PED	Nov 19, 2015			
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>						
	N019667 002	5753618	May 19, 2015			
		5753618*PED	Nov 19, 2015			
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>						
	N019667 003	5753618	May 19, 2015			
		5753618*PED	Nov 19, 2015			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>OCTREOTIDE ACETATE - SANDOSTATIN</u>						
N019667 004	5753618	May 19, 2015				
	5753618*PED	Nov 19, 2015				
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>						
N019667 005	5753618	May 19, 2015				
	5753618*PED	Nov 19, 2015				
<u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u>						
N021008 001	5538739	Jul 23, 2013		DP		
	5538739*PED	Jan 23, 2014				
	5639480	Jun 17, 2014		DP		
	5639480*PED	Dec 17, 2014				
	5688530	Nov 18, 2014		U-268		
	5688530*PED	May 18, 2015				
	5922338	Jul 13, 2016		DP		
	5922338*PED	Jan 13, 2017				
	5922682	Jul 13, 2016		DP		
	5922682*PED	Jan 13, 2017				
<u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u>						
N021008 002	5538739	Jul 23, 2013				
	5538739*PED	Jan 23, 2014		DP		
	5639480	Jun 17, 2014				
	5639480*PED	Dec 17, 2014				
	5688530	Nov 18, 2014		U-268		
	5688530*PED	May 18, 2015				
	5922338	Jul 13, 2016		DP		
	5922338*PED	Jan 13, 2017				
	5922682	Jul 13, 2016		DP		
	5922682*PED	Jan 13, 2017				
<u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u>						
N021008 003	5538739	Jul 23, 2013				
	5538739*PED	Jan 23, 2014		DP		
	5639480	Jun 17, 2014				
	5639480*PED	Dec 17, 2014				
	5688530	Nov 18, 2014		U-268		
	5688530*PED	May 18, 2015				
	5922338	Jul 13, 2016		DP		
	5922338*PED	Jan 13, 2017				
	5922682	Jul 13, 2016		DP		
	5922682*PED	Jan 13, 2017				
<u>OFLOXACIN - FLOXIN OTIC</u>						
N020799 001	5401741	Mar 27, 2012		U-407		
<u>OLANZAPINE - OLANZAPINE</u>						
A076000 001					PC	Apr 21, 2012
<u>OLANZAPINE - OLANZAPINE</u>						
A076000 002					PC	Apr 21, 2012
<u>OLANZAPINE - OLANZAPINE</u>						
A076000 003					PC	Apr 21, 2012
<u>OLANZAPINE - OLANZAPINE</u>						
A076000 004					PC	Apr 21, 2012
<u>OLANZAPINE - OLANZAPINE</u>						
A076000 005					PC	Apr 21, 2012
<u>OLANZAPINE - OLANZAPINE</u>						
A076133 002					PC	Apr 21, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>OLANZAPINE - ZYPREXA</u>						
N020592 001	5229382*PED	Oct 23, 2011			I-591	Mar 19, 2012
	5605897	Feb 25, 2014	U-176		NPP	Dec 04, 2012
	5605897*PED	Aug 25, 2014			NPP	Dec 04, 2012
	5627178*PED	Oct 23, 2011			PED	Jun 04, 2013
	5736541	Mar 24, 2015	U-307		PED	Jun 04, 2013
	5736541*PED	Sep 24, 2015				
	5817655*PED	Oct 23, 2011				
	5817656*PED	Oct 23, 2011				
	5817657*PED	Oct 23, 2011				
	5919485	Mar 24, 2015	U-308			
	5919485*PED	Sep 24, 2015				
	6251895	Sep 23, 2017				
	6251895*PED	Mar 23, 2018				
	6960577	Nov 01, 2017	U-963			
<u>OLANZAPINE - ZYPREXA</u>						
N020592 002	5229382*PED	Oct 23, 2011			I-591	Mar 19, 2012
	5605897	Feb 25, 2014	U-176		NPP	Dec 04, 2012
	5605897*PED	Aug 25, 2014			NPP	Dec 04, 2012
	5627178*PED	Oct 23, 2011			PED	Jun 04, 2013
	5736541	Mar 24, 2015	U-307		PED	Jun 04, 2013
	5736541*PED	Sep 24, 2015				
	5817655*PED	Oct 23, 2011				
	5817656*PED	Oct 23, 2011				
	5817657*PED	Oct 23, 2011				
	5919485	Mar 24, 2015	U-308			
	5919485*PED	Sep 24, 2015				
	6251895	Sep 23, 2017				
	6251895*PED	Mar 23, 2018				
	6960577	Nov 01, 2017	U-963			
<u>OLANZAPINE - ZYPREXA</u>						
N020592 003	5229382*PED	Oct 23, 2011			I-591	Mar 19, 2012
	5605897	Feb 25, 2014	U-176		NPP	Dec 04, 2012
	5605897*PED	Aug 25, 2014			NPP	Dec 04, 2012
	5627178*PED	Oct 23, 2011			PED	Jun 04, 2013
	5736541	Mar 24, 2015	U-307		PED	Jun 04, 2013
	5736541*PED	Sep 24, 2015				
	5817655*PED	Oct 23, 2011				
	5817656*PED	Oct 23, 2011				
	5817657*PED	Oct 23, 2011				
	5919485	Mar 24, 2015	U-308			
	5919485*PED	Sep 24, 2015				
	6251895	Sep 23, 2017				
	6251895*PED	Mar 23, 2018				
	6960577	Nov 01, 2017	U-963			
<u>OLANZAPINE - ZYPREXA</u>						
N020592 004	5229382*PED	Oct 23, 2011			I-591	Mar 19, 2012
	5605897	Feb 25, 2014	U-176		NPP	Dec 04, 2012
	5605897*PED	Aug 25, 2014			NPP	Dec 04, 2012
	5627178*PED	Oct 23, 2011			PED	Jun 04, 2013
	5736541	Mar 24, 2015	U-307		PED	Jun 04, 2013
	5736541*PED	Sep 24, 2015				
	5817655*PED	Oct 23, 2011				
	5817656*PED	Oct 23, 2011				
	5817657*PED	Oct 23, 2011				
	5919485	Mar 24, 2015	U-308			
	5919485*PED	Sep 24, 2015				
	6251895	Sep 23, 2017				
	6251895*PED	Mar 23, 2018				
	6960577	Nov 01, 2017	U-963			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>OLANZAPINE - ZYPREXA</u>						
N020592 005	5229382*PED 5605897 5605897*PED 5627178*PED 5736541 5736541*PED 5817655*PED 5817656*PED 5817657*PED 5919485 5919485*PED 6251895 6251895*PED 6960577	Oct 23, 2011 Feb 25, 2014 Aug 25, 2014 Oct 23, 2011 Mar 24, 2015 Sep 24, 2015 Oct 23, 2011 Oct 23, 2011 Oct 23, 2011 Mar 24, 2015 Sep 24, 2015 Oct 23, 2011 Oct 23, 2011 Oct 23, 2011 Nov 01, 2017	U-176 U-307 U-308 U-963	I-591 NPP NPP PED PED	Mar 19, 2012 Dec 04, 2012 Dec 04, 2012 Jun 04, 2013 Jun 04, 2013	
N020592 006	5229382*PED 5605897 5605897*PED 5627178*PED 5736541 5736541*PED 5817655*PED 5817656*PED 5817657*PED 5919485 5919485*PED 6251895 6251895*PED 6960577	Oct 23, 2011 Feb 25, 2014 Aug 25, 2014 Oct 23, 2011 Mar 24, 2015 Sep 24, 2015 Oct 23, 2011 Oct 23, 2011 Oct 23, 2011 Mar 24, 2015 Sep 24, 2015 Oct 23, 2011 Oct 23, 2011 Oct 23, 2011 Nov 01, 2017	U-176 U-307 U-308 U-963	I-591 NPP NPP PED PED	Mar 19, 2012 Dec 04, 2012 Dec 04, 2012 Jun 04, 2013 Jun 04, 2013	
<u>OLANZAPINE - ZYPREXA</u>						
N021253 001	5229382*PED	Oct 23, 2011				
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
N021086 001	5229382*PED 6960577	Oct 23, 2011 Nov 01, 2017	U-964	I-591 NPP NPP PED PED	Mar 19, 2012 Dec 04, 2012 Dec 04, 2012 Jun 04, 2013 Jun 04, 2013	
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
N021086 002	5229382*PED 6960577	Oct 23, 2011 Nov 01, 2017	U-964	I-591 NPP NPP PED PED	Mar 19, 2012 Dec 04, 2012 Dec 04, 2012 Jun 04, 2013 Jun 04, 2013	
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
N021086 003	5229382*PED 6960577	Oct 23, 2011 Nov 01, 2017	U-964	I-591 NPP NPP PED PED	Mar 19, 2012 Dec 04, 2012 Dec 04, 2012 Jun 04, 2013 Jun 04, 2013	
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
N021086 004	5229382*PED 6960577	Oct 23, 2011 Nov 01, 2017	U-964	I-591 NPP NPP PED PED	Mar 19, 2012 Dec 04, 2012 Dec 04, 2012 Jun 04, 2013 Jun 04, 2013	
<u>OLANZAPINE PAMOATE - ZYPREXA RELPREVV</u>						
N022173 001	5229382*PED 6169084	Oct 23, 2011 Sep 30, 2018	DS DP U-1026	NP	Dec 11, 2012	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>OLANZAPINE PAMOATE - ZYPREXA RELPREVV</u>							
N022173 002	5229382*PED 6169084	Oct 23, 2011 Sep 30, 2018		DS DP U-1026		NP	Dec 11, 2012
<u>OLANZAPINE PAMOATE - ZYPREXA RELPREVV</u>							
N022173 003	5229382*PED 6169084	Oct 23, 2011 Sep 30, 2018		DS DP U-1026		NP	Dec 11, 2012
<u>OLMESARTAN MEDOXOMIL - BENICAR</u>							
N021286 001	5616599 5616599*PED 6878703 6878703*PED	Apr 25, 2016 Oct 25, 2016 Nov 19, 2021 May 19, 2022		DS DP U-500 U-3	Y	NPP PED	Feb 04, 2013 Aug 04, 2013
<u>OLMESARTAN MEDOXOMIL - BENICAR</u>							
N021286 003	5616599 5616599*PED 6878703 6878703*PED	Apr 25, 2016 Oct 25, 2016 Nov 19, 2021 May 19, 2022		DS DP U-500 U-3	Y	NPP PED	Feb 04, 2013 Aug 04, 2013
<u>OLMESARTAN MEDOXOMIL - BENICAR</u>							
N021286 004	5616599 5616599*PED 6878703 6878703*PED	Apr 25, 2016 Oct 25, 2016 Nov 19, 2021 May 19, 2022		DS DP U-500 U-3	Y	NPP PED	Feb 04, 2013 Aug 04, 2013
<u>OLOPATADINE HYDROCHLORIDE - PATADAY</u>							
N021545 001	5641805 5641805*PED 6995186 6995186*PED 7402609 7402609*PED	Jun 06, 2015 Dec 06, 2015 Nov 12, 2023 May 12, 2024 Jun 19, 2022 Dec 19, 2022		U-765 DP U-765 DP			
<u>OLOPATADINE HYDROCHLORIDE - PATANASE</u>							
N021861 001	7977376 7977376*PED	Feb 02, 2023 Aug 02, 2023		DP DP		NPP PED PED	Dec 01, 2012 Jun 01, 2013 Oct 15, 2011
<u>OLOPATADINE HYDROCHLORIDE - PATANOL</u>							
N020688 001	5641805 5641805*PED	Jun 06, 2015 Dec 06, 2015		U-184			
<u>OMEGA-3-ACID ETHYL ESTERS - LOVAZA</u>							
N021654 001	5502077 5656667 7732488	Mar 26, 2013 Apr 10, 2017 Jan 30, 2025	DS DS DP DS DP	U-822 U-822		M-87	Sep 16, 2012
<u>OMEPRAZOLE - PRILOSEC</u>							
N019810 001	6147103 6147103*PED 6150380 6150380*PED 6166213 6166213*PED 6191148 6191148*PED	Oct 09, 2018 Apr 09, 2019 Nov 10, 2018 May 10, 2019 Oct 09, 2018 Apr 09, 2019 Oct 09, 2018 Apr 09, 2019					
<u>OMEPRAZOLE - PRILOSEC</u>							
N019810 002	6147103 6147103*PED 6150380 6150380*PED 6166213 6166213*PED 6191148 6191148*PED	Oct 09, 2018 Apr 09, 2019 Nov 10, 2018 May 10, 2019 Oct 09, 2018 Apr 09, 2019 Oct 09, 2018 Apr 09, 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
<u>OMEPRAZOLE - PRILOSEC</u>							
N019810 003	6147103	Oct	09, 2018				
	6147103*PED	Apr	09, 2019				
	6150380	Nov	10, 2018				
	6150380*PED	May	10, 2019				
	6166213	Oct	09, 2018				
	6166213*PED	Apr	09, 2019				
	6191148	Oct	09, 2018				
	6191148*PED	Apr	09, 2019				
<u>OMEPRAZOLE MAGNESIUM - PRILOSEC</u>							
N022056 001	5690960	Nov	25, 2014		DP	U-864	
	5900424	May	04, 2016	DS		U-864	
	6428810	Nov	03, 2019		DP	U-864	
<u>OMEPRAZOLE MAGNESIUM - PRILOSEC</u>							
N022056 002	5690960	Nov	25, 2014		DP	U-864	
	5900424	May	04, 2016	DS		U-864	
	6428810	Nov	03, 2019		DP	U-864	
<u>OMEPRAZOLE MAGNESIUM - PRILOSEC OTC</u>							
N021229 001	5690960	Nov	25, 2014				
	5753265	Jun	07, 2015				
	5817338	Oct	06, 2015				
	5900424	May	04, 2016				
	6403616	Nov	15, 2019				
	6428810	Nov	03, 2019				
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>							
N021636 001	5840737	Jul	16, 2016			U-588	
	6489346	Jul	16, 2016	DS	DP	U-588	
	6645988	Jul	16, 2016	DS	DP		
	6699885	Jul	16, 2016			U-588	
	6780882	Jul	16, 2016	DS	DP		
	7399772	Jul	16, 2016			U-588	
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>							
N021636 002	5840737	Jul	16, 2016			U-624	
	5840737	Jul	16, 2016			U-623	
	6489346	Jul	16, 2016	DS	DP	U-624	
	6489346	Jul	16, 2016	DS	DP	U-623	
	6645988	Jul	16, 2016	DS	DP		
	6699885	Jul	16, 2016			U-623	
	6699885	Jul	16, 2016			U-624	
	6780882	Jul	16, 2016	DS	DP		
	7399772	Jul	16, 2016			U-623	
	7399772	Jul	16, 2016			U-624	
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>							
N021849 001	6489346	Jul	16, 2016	DS	DP	U-588	
	6645988	Jul	16, 2016	DS	DP		
	6699885	Jul	16, 2016			U-588	
	7399772	Jul	16, 2016			U-588	
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>							
N021849 002	6489346	Jul	16, 2016	DS	DP	U-623	
	6645988	Jul	16, 2016	DS	DP		
	6699885	Jul	16, 2016			U-623	
	7399772	Jul	16, 2016			U-623	
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID OTC</u>							
N022281 001	6489346	Jul	15, 2016		DP	U-1025	
	6645988	Jul	15, 2016		DP		
	6699885	Jul	15, 2016		DP		
	7399772	Jul	15, 2016			U-1025	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ONDANSETRON - ZOFRAN ODT</u>							
N020781 001	5955488	Nov	14, 2015				
	5955488*PED	May	14, 2016				
	6063802	Nov	14, 2015				
	6063802*PED	May	14, 2016				
<u>ONDANSETRON - ZOFRAN ODT</u>							
N020781 002	5955488	Nov	14, 2015				
	5955488*PED	May	14, 2016				
	6063802	Nov	14, 2015				
	6063802*PED	May	14, 2016				
<u>ONDANSETRON HYDROCHLORIDE - ZOFRAN</u>							
N020103 001	5344658	Sep	06, 2011				
	5344658*PED	Mar	06, 2012				
<u>ONDANSETRON HYDROCHLORIDE - ZOFRAN</u>							
N020103 002	5344658	Sep	06, 2011				
	5344658*PED	Mar	06, 2012				
<u>ONDANSETRON HYDROCHLORIDE - ZOFRAN</u>							
N020103 003	5344658	Sep	06, 2011				
	5344658*PED	Mar	06, 2012				
<u>ONDANSETRON HYDROCHLORIDE - ZOFRAN</u>							
N020605 001	5854270	Nov	20, 2015		DP	U-44	
	5854270*PED	May	20, 2016				
<u>ORLISTAT - ALLI</u>							
N021887 001	6004996	Jan	06, 2018		DP		
<u>ORLISTAT - XENICAL</u>							
N020766 001	6004996	Jan	06, 2018				
	6004996*PED	Jul	06, 2018				
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>							
N021087 001	5763483	Dec	27, 2016	DS	U-1113		
	5763483*PED	Jun	27, 2017				
	5866601	Feb	02, 2016	DS	DP		
	5866601*PED	Aug	02, 2016				
	5952375	Feb	27, 2015	DS	DP		
	5952375*PED	Aug	27, 2015				
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>							
N021087 002	5763483	Dec	27, 2016	DS	U-1113		
	5763483*PED	Jun	27, 2017				
	5866601	Feb	02, 2016	DS	DP		
	5866601*PED	Aug	02, 2016				
	5952375	Feb	27, 2015	DS	DP		
	5952375*PED	Aug	27, 2015				
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>							
N021087 003	5763483	Dec	27, 2016	DS	U-1113		
	5763483*PED	Jun	27, 2017				
	5866601	Feb	02, 2016	DS	DP		
	5866601*PED	Aug	02, 2016				
	5952375	Feb	27, 2015	DS	DP		
	5952375*PED	Aug	27, 2015				
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>							
N021246 001	5763483	Dec	27, 2016	DS	U-376		
	5763483	Dec	27, 2016	DS	U-1113		
	5763483*PED	Jun	27, 2017				
	5866601	Feb	02, 2016	DS	DP		
	5866601*PED	Aug	02, 2016				
	5952375	Feb	27, 2015	DS	DP		
	5952375*PED	Aug	27, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
OSELTAMIVIR PHOSPHATE - TAMIFLU							
N021246 002	5763483	Dec	27, 2016	DS DP	U-1113		
	5763483*PED	Jun	27, 2017				
	5866601	Feb	02, 2016	DS DP			
	5866601*PED	Aug	02, 2016				
	5952375	Feb	27, 2015	DS DP			
	5952375*PED	Aug	27, 2015				
OXALIPLATIN - ELOXATIN							
N021492 001	5290961	Jan	12, 2013	DS			
	5290961*PED	Jul	12, 2013				
	5338874	Apr	07, 2013	DS			
	5338874*PED	Oct	07, 2013				
	5420319	Aug	09, 2016	DS			
	5420319*PED	Feb	09, 2017				
OXALIPLATIN - ELOXATIN							
N021492 002	5290961	Jan	12, 2013	DS			
	5290961*PED	Jul	12, 2013				
	5338874	Apr	07, 2013	DS			
	5338874*PED	Oct	07, 2013				
	5420319	Aug	09, 2016	DS			
	5420319*PED	Feb	09, 2017				
OXALIPLATIN - ELOXATIN							
N021759 001	5290961	Jan	12, 2013	DS			
	5290961*PED	Jul	12, 2013				
	5338874	Apr	07, 2013	DS			
	5338874*PED	Oct	07, 2013				
	5420319	Aug	09, 2016	DS			
	5420319*PED	Feb	09, 2017				
	5716988	Aug	07, 2015	DP			
	5716988*PED	Feb	07, 2016				
OXALIPLATIN - ELOXATIN							
N021759 002	5290961	Jan	12, 2013	DS			
	5290961*PED	Jul	12, 2013				
	5338874	Apr	07, 2013	DS			
	5338874*PED	Oct	07, 2013				
	5420319	Aug	09, 2016	DS			
	5420319*PED	Feb	09, 2017				
	5716988	Aug	07, 2015	DP			
	5716988*PED	Feb	07, 2016				
OXALIPLATIN - ELOXATIN							
N021759 003	5290961	Jan	12, 2013	DS			
	5290961*PED	Jul	12, 2013				
	5338874	Apr	07, 2013	DS			
	5338874*PED	Oct	07, 2013				
	5420319	Aug	09, 2016	DS			
	5420319*PED	Feb	09, 2017				
	5716988	Aug	07, 2015	DP			
	5716988*PED	Feb	07, 2016				
OXANDROLONE - OXANDRIN							
N013718 001	5872147	Dec	05, 2017		U-585		
	6090799	Jul	18, 2017		U-585		
	6576659	Dec	05, 2017		U-585		
	6670351	Oct	20, 2012		U-585		
	6828313	Dec	05, 2017		U-585		
OXANDROLONE - OXANDRIN							
N013718 002	5872147	Dec	05, 2017		U-585		
	6090799	Jul	18, 2017		U-585		
	6576659	Dec	05, 2017		U-585		
	6670351	Oct	20, 2012		U-585		
	6828313	Dec	05, 2017		U-585		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>OXCARBAZEPINE - TRILEPTAL</u>							
N021014 001	7037525	Feb	12, 2018			U-724	
	7037525*PED	Aug	12, 2018				
<u>OXCARBAZEPINE - TRILEPTAL</u>							
N021014 002	7037525	Feb	12, 2018			U-724	
	7037525*PED	Aug	12, 2018				
<u>OXCARBAZEPINE - TRILEPTAL</u>							
N021014 003	7037525	Feb	12, 2018			U-724	
	7037525*PED	Aug	12, 2018				
<u>OXYBUTYNIN - ANTUROL</u>							
N202513 001						NP	Dec 07, 2014
<u>OXYBUTYNIN - OXYTROL</u>							
N021351 002	5601839	Apr	26, 2015				
	5834010	Apr	26, 2015				
	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		
<u>OXYBUTYNIN CHLORIDE - DITROPAN XL</u>							
N020897 001	5674895	May	22, 2015				
	5674895*PED	Nov	22, 2015				
	5840754	May	22, 2015				
	5840754*PED	Nov	22, 2015				
	5912268	May	22, 2015				
	5912268*PED	Nov	22, 2015				
	6262115	May	22, 2015		U-393		
	6262115*PED	Nov	22, 2015		U-393		
<u>OXYBUTYNIN CHLORIDE - DITROPAN XL</u>							
N020897 002	5674895	May	22, 2015				
	5674895*PED	Nov	22, 2015				
	5840754	May	22, 2015				
	5840754*PED	Nov	22, 2015				
	5912268	May	22, 2015				
	5912268*PED	Nov	22, 2015				
	6262115	May	22, 2015		U-393		
	6262115*PED	Nov	22, 2015		U-393		
<u>OXYBUTYNIN CHLORIDE - DITROPAN XL</u>							
N020897 003	5674895	May	22, 2015				
	5674895*PED	Nov	22, 2015				
	5840754	May	22, 2015				
	5840754*PED	Nov	22, 2015				
	5912268	May	22, 2015				
	5912268*PED	Nov	22, 2015				
	6262115	May	22, 2015		U-393		
	6262115*PED	Nov	22, 2015		U-393		
<u>OXYBUTYNIN CHLORIDE - GELNIQUE</u>							
N022204 001	7029694	Apr	26, 2020	DP	U-318	NDF	Jan 27, 2012
	7179483	Apr	26, 2020		U-318		
<u>OXYCODONE HYDROCHLORIDE - OXECTA</u>							
N202080 001	7201920	Mar	16, 2025	DP			
	7510726	Nov	26, 2023	DP			
	7981439	Nov	26, 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
OXYCODONE HYDROCHLORIDE - OXECTA						
N0202080 002	7201920	Mar 16, 2025	DP			
	7510726	Nov 26, 2023	DP			
	7981439	Nov 26, 2023	DP			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 001	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 002	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 003	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 004	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 005	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 006	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 007	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N020553 008	5508042	Apr 16, 2013		U-443		
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N022272 001	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 2025	DP			
OXYCODONE HYDROCHLORIDE - OXYCONTIN						
N022272 002	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 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
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 003	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 2025	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 004	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 2025	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 005	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 2025	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 006	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 2025	DP			
<u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u>						
N022272 007	5508042	Apr 16, 2013		U-443		
	6488963	Jun 24, 2017	DP			
	7674799	Mar 30, 2025	DP			
	7674800	Mar 30, 2025	DS			
	7683072	Mar 30, 2025	DS			
	7776314	Apr 19, 2025	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 001	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	7276250	Feb 04, 2023	DP	U-826		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 002	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	7276250	Feb 04, 2023	DP	U-826		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 003	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	7276250	Feb 04, 2023	DP	U-826		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 004	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	7276250	Feb 04, 2023	DP	U-826		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 005	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	7276250	Feb 04, 2023	DP	U-826		
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 006	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	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
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N021610 007	5662933	Sep 09, 2013	DP			
	5958456	Sep 09, 2013	DP			
	7276250	Feb 04, 2023	DP U-826			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 001	8075872	Nov 20, 2023	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 002	8075872	Nov 20, 2023	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 003	8075872	Nov 20, 2023	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 004	8075872	Nov 20, 2023	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 005	8075872	Nov 20, 2023	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 006	8075872	Nov 20, 2023	DP			
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>						
N201655 007	8075872	Nov 20, 2023	DP			
<u>PACLITAXEL - ABRAXANE</u>						
N021660 001	5439686	Feb 22, 2013	DP			
	5498421	Mar 12, 2013	DP U-634			
	6096331	Feb 22, 2013	DP U-633			
	6506405	Feb 22, 2013	DP U-633			
	6537579	Feb 22, 2013	U-632			
	6749868	Feb 22, 2013	DP			
	6753006	Feb 22, 2013	DP			
	7820788	Mar 03, 2024	DP U-1092			
	7923536	Dec 09, 2023	U-1117			
	RE41884	Aug 14, 2016	U-1117			
<u>PALIPERIDONE - INVEGA</u>						
N021999 001				I-605	Jul 31, 2012	
				I-606	Jul 31, 2012	
				NPP	Apr 06, 2014	
				NCE	Dec 19, 2011	
				PED	Oct 06, 2014	
				PED	Jun 19, 2012	
				PED	Jan 31, 2013	
				PED	Jan 31, 2013	
<u>PALIPERIDONE - INVEGA</u>						
N021999 002				I-605	Jul 31, 2012	
				I-606	Jul 31, 2012	
				NPP	Apr 06, 2014	
				NCE	Dec 19, 2011	
				PED	Oct 06, 2014	
				PED	Jun 19, 2012	
				PED	Jan 31, 2013	
				PED	Jan 31, 2013	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PALIPERIDONE - INVEGA</u>						
	N021999 003				I-605 I-606 NPP NCE PED PED PED PED	Jul 31, 2012 Jul 31, 2012 Apr 06, 2014 Dec 19, 2011 Oct 06, 2014 Jun 19, 2012 Jan 31, 2013 Jan 31, 2013
<u>PALIPERIDONE - INVEGA</u>						
	N021999 004				NPP NCE PED PED	Apr 06, 2014 Dec 19, 2011 Oct 06, 2014 Jun 19, 2012
<u>PALIPERIDONE - INVEGA</u>						
	N021999 006				I-605 I-606 NPP NCE PED PED PED PED	Jul 31, 2012 Jul 31, 2012 Apr 06, 2014 Dec 19, 2011 Oct 06, 2014 Jun 19, 2012 Jan 31, 2013 Jan 31, 2013
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N022264 001	5254556 5254556*PED 5352459 5352459*PED 6077843 6077843*PED 6555544 6555544*PED	Oct 27, 2012 Apr 27, 2013 Dec 16, 2012 Jun 16, 2013 May 12, 2017 Nov 12, 2017 Nov 10, 2018 May 10, 2019	DS DP U-543 DP DP U-543 DP U-543		NDF NCE PED PED	Jul 31, 2012 Dec 19, 2011 Jun 19, 2012 Jan 31, 2013
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N022264 002	5254556 5254556*PED 5352459 5352459*PED 6077843 6077843*PED 6555544 6555544*PED	Oct 27, 2012 Apr 27, 2013 Dec 16, 2012 Jun 16, 2013 May 12, 2017 Nov 12, 2017 Nov 10, 2018 May 10, 2019	DS DP U-543 DP DP U-543 DP U-543		NDF NCE PED PED	Jul 31, 2012 Dec 19, 2011 Jun 19, 2012 Jan 31, 2013
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N022264 003	5254556 5254556*PED 5352459 5352459*PED 6077843 6077843*PED 6555544 6555544*PED	Oct 27, 2012 Apr 27, 2013 Dec 16, 2012 Jun 16, 2013 May 12, 2017 Nov 12, 2017 Nov 10, 2018 May 10, 2019	DS DP U-543 DP DP U-543 DP U-543		NDF NCE PED PED	Jul 31, 2012 Dec 19, 2011 Jun 19, 2012 Jan 31, 2013
<u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>						
N022264 004	5254556 5254556*PED 5352459 5352459*PED 6077843 6077843*PED 6555544 6555544*PED	Oct 27, 2012 Apr 27, 2013 Dec 16, 2012 Jun 16, 2013 May 12, 2017 Nov 12, 2017 Nov 10, 2018 May 10, 2019	DS DP U-543 DP DP U-543 DP U-543		NDF NCE PED PED	Jul 31, 2012 Dec 19, 2011 Jun 19, 2012 Jan 31, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u>								
N022264 005	5254556	Oct	27,	2012	DS	DP	U-543	NDF Jul 31, 2012
	5254556*PED	Apr	27,	2013				NCE Dec 19, 2011
	5352459	Dec	16,	2012	DP			PED Jun 19, 2012
	5352459*PED	Jun	16,	2013				PED Jan 31, 2013
	6077843	May	12,	2017	DP	U-543		
	6077843*PED	Nov	12,	2017				
	6555544	Nov	10,	2018	DP	U-543		
	6555544*PED	May	10,	2019				
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>								
N021372 001	5202333	Apr	13,	2015	DS	DP	U-528	
	7947724	Jan	30,	2024	DP			
	7947725	Jan	30,	2024	DP			
	7960424	Jan	30,	2024	DP			
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>								
N021372 002	5202333	Apr	13,	2015	DS	DP	U-901	
	7947724	Jan	30,	2024	DP			
	7947725	Jan	30,	2024	DP			
	7960424	Jan	30,	2024	DP			
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>								
N022233 001	5202333	Apr	13,	2015	DS	DP	U-528	NDF Aug 22, 2011
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>								
N020725 001							I-625 Apr 30, 2013	
							M-93 Jul 29, 2013	
							NCE Apr 30, 2014	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>								
N020725 002							I-625 Apr 30, 2013	
							M-93 Jul 29, 2013	
							NCE Apr 30, 2014	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u>								
N020725 003							I-625 Apr 30, 2013	
							M-93 Jul 29, 2013	
							NCE Apr 30, 2014	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u>								
N022523 001							NCE Apr 12, 2015	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u>								
N022523 002							NCE Apr 12, 2015	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u>								
N022523 003							NCE Apr 12, 2015	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u>								
N022523 004							NCE Apr 12, 2015	
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>								
N022210 001	7658918	Feb	20,	2028	DP			NCE Aug 27, 2014
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>								
N022210 002	7658918	Feb	20,	2028	DP			NCE Aug 27, 2014
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>								
N022210 003	7658918	Feb	20,	2028	DP			NCE Aug 27, 2014
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>								
N022210 004	7658918	Feb	20,	2028	DP			NCE Aug 27, 2014
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>								
N020987 001	5997903	Dec	07,	2016			I-614 Nov 12, 2012	
	5997903*PED	Jun	07,	2017			M-54 Nov 12, 2012	
							PED May 12, 2013	
							PED May 12, 2013	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
PANTOPRAZOLE SODIUM - PROTONIX							
N020987 002	5997903	Dec	07, 2016			I-614	Nov 12, 2012
	5997903*PED	Jun	07, 2017			M-54	Nov 12, 2012
						PED	May 12, 2013
						PED	May 12, 2013
PANTOPRAZOLE SODIUM - PROTONIX							
N022020 001	7544370	Jun	07, 2026	DP		I-614	Nov 12, 2012
	7544370*PED	Dec	07, 2026			M-54	Nov 12, 2012
	7550153	Sep	30, 2024	U-859		PED	May 12, 2013
	7550153*PED	Mar	30, 2025	U-859		PED	May 12, 2013
	7553498	Sep	30, 2024				
	7553498*PED	Mar	30, 2025				
	7838027	Sep	30, 2024	DP	U-859		
	7838027*PED	Mar	30, 2025				
PANTOPRAZOLE SODIUM - PROTONIX IV							
N020988 001	6780881	Nov	17, 2021	DP			
	6780881*PED	May	17, 2022				
	7351723	Nov	17, 2021	DP			
	7351723*PED	May	17, 2022				
PARICALCITOL - ZEMPLAR							
N020819 001	5246925	Apr	17, 2012		U-314		
	5246925*PED	Oct	17, 2012				
	5587497	Dec	24, 2013				
	5587497*PED	Jun	24, 2014				
	5597815	Jul	13, 2015	U-1195			
	5597815*PED	Jan	13, 2016				
	6136799	Apr	08, 2018				
	6136799*PED	Oct	08, 2018				
	6361758	Apr	08, 2018	DP			
	6361758*PED	Oct	08, 2018				
PARICALCITOL - ZEMPLAR							
N020819 002	5246925	Apr	17, 2012		U-314		
	5246925*PED	Oct	17, 2012				
	5587497	Dec	24, 2013				
	5587497*PED	Jun	24, 2014				
	5597815	Jul	13, 2015	U-1195			
	5597815*PED	Jan	13, 2016				
PARICALCITOL - ZEMPLAR							
N021606 001	5246925	Apr	17, 2012		U-671		
	5246925*PED	Oct	17, 2012			I-599	Jun 29, 2012
	5587497	Dec	24, 2013	DS			
	5587497*PED	Jun	24, 2014				
	5597815	Jul	13, 2015				
	5597815*PED	Jan	13, 2016	U-1195			
PARICALCITOL - ZEMPLAR							
N021606 002	5246925	Apr	17, 2012		U-671		
	5246925*PED	Oct	17, 2012			I-599	Jun 29, 2012
	5587497	Dec	24, 2013	DS			
	5587497*PED	Jun	24, 2014				
	5597815	Jul	13, 2015				
	5597815*PED	Jan	13, 2016	U-1195			
PARICALCITOL - ZEMPLAR							
N021606 003	5246925	Apr	17, 2012		U-671		
	5246925*PED	Oct	17, 2012			I-599	Jun 29, 2012
	5587497	Dec	24, 2013	DS			
	5587497*PED	Jun	24, 2014				
	5597815	Jul	13, 2015				
	5597815*PED	Jan	13, 2016	U-1195			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PAROXETINE HYDROCHLORIDE - PAROXETINE HYDROCHLORIDE</u>						
	A091427 001				PC	Nov 01, 2011
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020031 001	5872132 5872132*PED 5900423 5900423*PED 6113944 6113944*PED 6121291 6121291 6121291*PED 6121291*PED 6133289 6133289*PED	May 19, 2015 Nov 19, 2015 May 19, 2015 Nov 19, 2015 Dec 14, 2014 Jun 14, 2015 Mar 17, 2017 Mar 17, 2017 Sep 17, 2017 Sep 17, 2017 May 19, 2015 Nov 19, 2015		U-286 U-431 U-431 U-286 U-358 U-358		
N020031 002	5872132 5872132*PED 5900423 5900423*PED 6113944 6113944*PED 6121291 6121291 6121291*PED 6121291*PED 6133289 6133289*PED	May 19, 2015 Nov 19, 2015 May 19, 2015 Nov 19, 2015 Dec 14, 2014 Jun 14, 2015 Mar 17, 2017 Mar 17, 2017 Sep 17, 2017 Sep 17, 2017 May 19, 2015 Nov 19, 2015		U-286 U-431 U-431 U-286 U-358 U-358		
N020031 003	5872132 5872132*PED 5900423 5900423*PED 6113944 6113944*PED 6121291 6121291 6121291*PED 6121291*PED 6133289 6133289*PED	May 19, 2015 Nov 19, 2015 May 19, 2015 Nov 19, 2015 Dec 14, 2014 Jun 14, 2015 Mar 17, 2017 Mar 17, 2017 Sep 17, 2017 Sep 17, 2017 May 19, 2015 Nov 19, 2015		U-286 U-431 U-431 U-286 U-358 U-358		
N020031 004	5872132 5872132*PED 5900423 5900423*PED 6113944 6113944*PED 6121291 6121291 6121291*PED 6121291*PED 6133289 6133289*PED	May 19, 2015 Nov 19, 2015 May 19, 2015 Nov 19, 2015 Dec 14, 2014 Jun 14, 2015 Mar 17, 2017 Mar 17, 2017 Sep 17, 2017 Sep 17, 2017 May 19, 2015 Nov 19, 2015		U-431 U-286 U-431 U-286 U-358 U-358		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020031 005	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6113944	Dec 14, 2014				
	6113944*PED	Jun 14, 2015				
	6121291	Mar 17, 2017		U-431		
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-431		
	6133289	May 19, 2015		U-358		
	6133289*PED	Nov 19, 2015		U-358		
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020710 001	5811436	Sep 22, 2015				
	5811436*PED	Mar 22, 2016				
	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-286		
	6121291	Mar 17, 2017		U-431		
	6121291*PED	Sep 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-431		
	6133289	May 19, 2015		U-358		
	6133289*PED	Nov 19, 2015		U-358		
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020885 001	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6063927	Apr 23, 2019				
	6063927*PED	Oct 23, 2019				
	6080759	May 19, 2015				
	6080759*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-286		
	6121291	Mar 17, 2017		U-431		
	6121291*PED	Sep 17, 2017		U-431		
	6121291*PED	Sep 17, 2017		U-286		
	6133289	May 19, 2015		U-358		
	6133289*PED	Nov 19, 2015		U-358		
	6172233	Jan 15, 2018				
	6172233*PED	Jul 15, 2018				
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020885 002	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6063927	Apr 23, 2019				
	6063927*PED	Oct 23, 2019				
	6080759	May 19, 2015				
	6080759*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-431		
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-431		
	6121291*PED	Sep 17, 2017		U-286		
	6133289	May 19, 2015		U-358		
	6133289*PED	Nov 19, 2015		U-358		
	6172233	Jan 15, 2018				
	6172233*PED	Jul 15, 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>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020885 003	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6063927	Apr 23, 2019				
	6063927*PED	Oct 23, 2019				
	6080759	May 19, 2015				
	6080759*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-431		
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-431		
	6133289	May 19, 2015		U-358		
	6133289*PED	Nov 19, 2015		U-358		
	6172233	Jan 15, 2018				
	6172233*PED	Jul 15, 2018				
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>						
N020885 004	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6062927	Apr 23, 2019				
	6063927*PED	Oct 23, 2019				
	6080759	May 19, 2015				
	6080759*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-431		
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-431		
	6133289	May 19, 2015		U-358		
	6133289*PED	Nov 19, 2015		U-358		
	6172233	Jan 15, 2018				
	6172233*PED	Jul 15, 2018				
<u>PAROXETINE HYDROCHLORIDE - PAXIL CR</u>						
N020936 001	5422123	Jun 06, 2012				
	5422123*PED	Dec 06, 2012				
	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-286		
	6133289	May 19, 2015		U-286		
	6133289*PED	Nov 19, 2015		U-286		
	6548084	Jul 19, 2016				
	6548084*PED	Jan 19, 2017				
	7229640	Jul 19, 2016	DP	U-816		
<u>PAROXETINE HYDROCHLORIDE - PAXIL CR</u>						
N020936 002	5422123	Jun 06, 2012				
	5422123*PED	Dec 06, 2012				
	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-286		
	6133289	May 19, 2015		U-286		
	6133289*PED	Nov 19, 2015		U-286		
	6548084	Jul 19, 2016				
	6548084*PED	Jan 19, 2017				
	7229640	Jul 19, 2016	DP	U-816		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PAROXETINE HYDROCHLORIDE - PAXIL CR</u>						
N020936 003	5422123	Jun 06, 2012				
	5422123*PED	Dec 06, 2012				
	5872132	May 19, 2015				
	5872132*PED	Nov 19, 2015				
	5900423	May 19, 2015				
	5900423*PED	Nov 19, 2015				
	6121291	Mar 17, 2017		U-286		
	6121291*PED	Sep 17, 2017		U-286		
	6133289	May 19, 2015		U-286		
	6133289*PED	Nov 19, 2015		U-286		
	6548084	Jul 19, 2016				
	6548084*PED	Jan 19, 2017				
	7229640	Jul 19, 2016	DP	U-816		
<u>PAROXETINE MESYLATE - PEDEXVA</u>						
N021299 001	5874447	Jun 10, 2017		U-286		
	5874447	Jun 10, 2017		U-518		
	5874447	Jun 10, 2017		U-46		
	6703408	Oct 21, 2022	DP			
	7598271	Feb 11, 2023	DS	DP		
<u>PAROXETINE MESYLATE - PEDEXVA</u>						
N021299 002	5874447	Jun 10, 2017		U-518		
	5874447	Jun 10, 2017		U-46		
	5874447	Jun 10, 2017		U-286		
	6703408	Oct 21, 2022	DP			
	7598271	Feb 11, 2023	DS	DP		
<u>PAROXETINE MESYLATE - PEDEXVA</u>						
N021299 003	5874447	Jun 10, 2017		U-518		
	5874447	Jun 10, 2017		U-46		
	5874447	Jun 10, 2017		U-286		
	6703408	Oct 21, 2022	DP			
	7598271	Feb 11, 2023	DS	DP		
<u>PAROXETINE MESYLATE - PEDEXVA</u>						
N021299 004	5874447	Jun 10, 2017		U-46		
	5874447	Jun 10, 2017		U-518		
	5874447	Jun 10, 2017		U-286		
	6703408	Oct 21, 2022	DP			
	7598271	Feb 11, 2023	DS	DP		
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N022465 001	7105530	Dec 19, 2021	DS	DP		
	7262203	Dec 19, 2021	DS	DP	NCE	Oct 19, 2014
<u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u>						
N022465 002	7105530	Dec 19, 2021	DS	DP		
	7262203	Dec 19, 2021	DS	DP	NCE	Oct 19, 2014
<u>PEGAPTANIB SODIUM - MACUGEN</u>						
N021756 001	5919455	Oct 27, 2013	DS			
	5932462	Aug 03, 2016	DS			
	6011020	Jan 04, 2017	DS			
	6051698	May 19, 2015	DS	U-622		
	6113906	Oct 27, 2013	DS			
<u>PEGVISOMANT - SOMAVERT</u>						
N021106 001	5350836	Sep 27, 2011		U-507		
	5681809	Sep 27, 2011		U-507		
	5849535	Mar 25, 2017	DS			
	5958879	Sep 27, 2011		U-507		
	6057292	Sep 21, 2015		U-507		
	6583115	Sep 27, 2011		U-507		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PEGVISOMANT - SOMAVERT</u>						
N021106 002	5350836	Sep 27, 2011	U-507			
	5681809	Sep 27, 2011	U-507			
	5849535	Mar 25, 2017	DS			
	5958879	Sep 27, 2011	U-507			
	6057292	Sep 21, 2015	U-507			
	6583115	Sep 27, 2011	U-507			
<u>PEGVISOMANT - SOMAVERT</u>						
N021106 003	5350836	Sep 27, 2011	U-507			
	5681809	Sep 27, 2011	U-507			
	5849535	Mar 25, 2017	DS			
	5958879	Sep 27, 2011	U-507			
	6057292	Sep 21, 2015	U-507			
	6583115	Sep 27, 2011	U-507			
<u>PEMETREXED DISODIUM - ALIMTA</u>						
N021462 001	5217974*PED	Sep 29, 2011			I-601	Jul 02, 2012
	5344932	Jul 24, 2016	DS DP		I-571	Sep 26, 2011
	5344932*PED	Jan 24, 2017			M-61	Mar 17, 2014
	7772209	Nov 24, 2021		U-1077	PED	Sep 17, 2014
	7772209*PED	May 24, 2022			PED	Aug 04, 2011
					PED	Mar 26, 2012
					PED	Jan 02, 2013
<u>PEMETREXED DISODIUM - ALIMTA</u>						
N021462 002	5217974*PED	Sep 29, 2011			I-601	Jul 02, 2012
	5344932	Jul 24, 2016	DS DP		I-571	Sep 26, 2011
	5344932*PED	Jan 24, 2017			M-61	Mar 17, 2014
	7772209	Nov 24, 2021		U-1077	PED	Sep 17, 2014
	7772209*PED	May 24, 2022			PED	Aug 04, 2011
					PED	Mar 26, 2012
					PED	Jan 02, 2013
<u>PEMIROLAST POTASSIUM - ALAMAST</u>						
N021079 001	5034230*PED	Jul 02, 2011		U-184		
<u>PENCICLOVIR SODIUM - DENAVIR</u>						
N020629 001	5840763	Sep 01, 2015	U-501			
	5866581	Oct 04, 2014	U-501			
	5916893	Sep 01, 2015	U-501			
	6124304	Oct 04, 2014	U-501			
	6469015	Oct 22, 2019	U-501			
	6579981	Jun 17, 2020	U-501			
<u>PENTETATE CALCIUM TRISODIUM - PENTETATE CALCIUM TRISODIUM</u>						
N021749 001					ODE	Aug 11, 2011
<u>PENTETATE ZINC TRISODIUM - PENTETATE ZINC TRISODIUM</u>						
N021751 001					ODE	Aug 11, 2011
<u>PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE - SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS</u>						
N021084 001	5607979	May 30, 2015				
<u>PERFLUTREN - DEFINITY</u>						
N021064 001	5527521	Feb 22, 2015	DP U-665			
	5585112	Dec 17, 2013	DP			
	6033645	Jun 19, 2016	U-665			
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N202088 001	6149938	Jul 23, 2018	DP			
<u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u>						
N202088 002	6149938	Jul 23, 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>PHENTOLAMINE MESYLATE - ORAVERSE</u>						
N022159 001	6764678	May 11, 2021		U-967		
	6872390	May 11, 2021	DP			
	7229630	Jun 20, 2023	DP			
	7569230	Oct 17, 2023		U-967		
	7575757	Apr 21, 2025	DP			
<u>PIMECROLIMUS - ELIDEL</u>						
N021302 001	5912238	Jun 15, 2016				
	5912238*PED	Dec 15, 2016				
	6352998	Oct 26, 2015				
	6352998*PED	Apr 26, 2016				
	6423722	Jun 26, 2018				
	6423722*PED	Dec 26, 2018				
<u>PIOGLITAZONE HYDROCHLORIDE - ACTOS</u>						
N021073 001	5965584	Jun 19, 2016		U-753		
	6150383	Jun 19, 2016		U-418		
	6150384	Jun 19, 2016		U-419		
	6166042	Jun 19, 2016		U-414		
	6166043	Jun 19, 2016		U-415		
	6172090	Jun 19, 2016		U-416		
	6211205	Jun 19, 2016		U-410		
	6271243	Jun 19, 2016		U-411		
	6303640	Aug 09, 2016		U-425		
	6329404	Jun 19, 2016		U-753		
<u>PIOGLITAZONE HYDROCHLORIDE - ACTOS</u>						
N021073 002	5965584	Jun 19, 2016		U-753		
	6150383	Jun 19, 2016		U-418		
	6150384	Jun 19, 2016		U-419		
	6166042	Jun 19, 2016		U-414		
	6166043	Jun 19, 2016		U-415		
	6172090	Jun 19, 2016		U-416		
	6211205	Jun 19, 2016		U-410		
	6271243	Jun 19, 2016		U-411		
	6303640	Aug 09, 2016		U-425		
	6329404	Jun 19, 2016		U-753		
<u>PIOGLITAZONE HYDROCHLORIDE - ACTOS</u>						
N021073 003	5965584	Jun 19, 2016		U-753		
	6150383	Jun 19, 2016		U-418		
	6150384	Jun 19, 2016		U-419		
	6166042	Jun 19, 2016		U-414		
	6166043	Jun 19, 2016		U-415		
	6172090	Jun 19, 2016		U-416		
	6211205	Jun 19, 2016		U-410		
	6271243	Jun 19, 2016		U-411		
	6303640	Aug 09, 2016		U-425		
	6329404	Jun 19, 2016		U-753		
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 001	6900184	Apr 14, 2023	DP	U-282		
	7915229	Apr 14, 2023	DP			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 002	6900184	Apr 14, 2023	DP	U-282		
	7915229	Apr 14, 2023	DP			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 003	6900184	Apr 14, 2023	DP	U-282		
	7915229	Apr 14, 2023	DP			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u>						
N050684 004	6900184	Apr 14, 2023	DP	U-282		
	7915229	Apr 14, 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
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>							
N050750 001	6207661	Feb	22, 2019	DP			
	6900184	Apr	14, 2023	DP	U-282		
	7915229	Apr	14, 2023	DP			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>							
N050750 002	6207661	Feb	22, 2019	DP			
	6900184	Apr	14, 2023	DP	U-282		
	7915229	Apr	14, 2023	DP			
<u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u>							
N050750 003	6207661	Feb	22, 2019	DP			
	6900184	Apr	14, 2023	DP	U-282		
	7915229	Apr	14, 2023	DP			
<u>PITAVASTATIN CALCIUM - LIVALO</u>							
N022363 001	5753675	May	19, 2015	DS	DP	U-998	
	5854259	Dec	29, 2015	DP			
	5856336	Jan	05, 2016	DS		U-998	
	6465477	Dec	20, 2016	DP			
	7022713	Feb	19, 2024			U-998	
<u>PITAVASTATIN CALCIUM - LIVALO</u>							
N022363 002	5753675	May	19, 2015	DS	DP	U-998	
	5854259	Dec	29, 2015	DP			
	5856336	Jan	05, 2016	DS		U-998	
	6465477	Dec	20, 2016	DP			
	7022713	Feb	19, 2024			U-998	
<u>PITAVASTATIN CALCIUM - LIVALO</u>							
N022363 003	5753675	May	19, 2015	DS	DP	U-998	
	5854259	Dec	29, 2015	DP			
	5856336	Jan	05, 2016	DS		U-998	
	6465477	Dec	20, 2016	DP			
	7022713	Feb	19, 2024			U-998	
<u>PLERIXAFOR - MOZOBIL</u>							
N022311 001	6987102	Jul	22, 2023			U-936	
	7897590	Jul	22, 2023			U-936	
	RE42152	Dec	10, 2013	DP			
<u>POLIDOCANOL - ASCLERA</u>							
N021201 001							NCE
							Mar 30, 2015
<u>POLIDOCANOL - ASCLERA</u>							
N021201 002							NCE
							Mar 30, 2015
<u>PORFIMER SODIUM - PHOTOFRIN</u>							
N020451 001	5438071	Aug	01, 2012				
<u>POSACONAZOLE - NOXAFIL</u>							
N022003 001	5661151	Jul	19, 2019	DS	DP	U-760	
	5703079	Aug	26, 2014	DS	DP	U-760	
	6958337	Oct	05, 2018	DS	DP	U-760	
<u>PRALATREXATE - FOLOTYN</u>							
N022468 001	6028071	Jul	16, 2017	DS	DP	U-1004	
	7622470	May	31, 2025			U-1015	
							OCE
							Sep 24, 2016
<u>PRALATREXATE - FOLOTYN</u>							
N022468 002	6028071	Jul	16, 2017	DS	DP	U-1004	
	7622470	May	31, 2025			U-1015	
							OCE
							Sep 24, 2016
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>							
N020667 001	6001861	Jan	16, 2018			U-784	
	6194445	Jan	16, 2018			U-784	
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>							
N020667 002	6001861	Jan	16, 2018			U-784	
	6194445	Jan	16, 2018			U-784	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>						
N020667 003	6001861	Jan 16, 2018	U-784		M-104	May 13, 2014
	6194445	Jan 16, 2018	U-784			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>						
N020667 005	6001861	Jan 16, 2018	U-784		M-104	May 13, 2014
	6194445	Jan 16, 2018	U-784			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>						
N020667 006	6001861	Jan 16, 2018	U-784		M-104	May 13, 2014
	6194445	Jan 16, 2018	U-784			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>						
N020667 007	6001861	Jan 16, 2018	U-784		M-104	May 13, 2014
	6194445	Jan 16, 2018	U-784			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 001	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 002	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 003	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 004	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 005	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 006	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421 007	7695734	Apr 26, 2028	DP		I-623	Mar 19, 2013
					NDF	Feb 19, 2013
<u>PRAMLINTIDE ACETATE - SYMLIN</u>						
N021332 001	5686411	Mar 16, 2019	DS	DP	U-638	
	5814600	Sep 29, 2015			U-639	
	6114304	Sep 05, 2017			U-640	
	6608029	Sep 07, 2013			U-641	
<u>PRAMLINTIDE ACETATE - SYMLIN</u>						
N021332 002	5686411	Mar 16, 2019	DS	DP	U-638	
	5814600	Sep 29, 2015			U-639	
	5814600	Sep 29, 2015			U-638	
	5814600	Sep 29, 2015			U-637	
	6114304	Sep 05, 2017			U-640	
	6114304	Sep 05, 2017			U-637	
	6608029	Sep 07, 2013			U-641	
	6608029	Sep 07, 2013			U-640	
	6608029	Sep 07, 2013			U-637	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE			PATENT DELIST REQUESTED		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PRAMINTIDE ACETATE - SYMLIN</u>								
N021332 003	5686411	Mar 16, 2019	DS	DP	U-638			
	5814600	Sep 29, 2015			U-638			
	5814600	Sep 29, 2015			U-637			
	5814600	Sep 29, 2015			U-639			
	6114304	Sep 05, 2017			U-640			
	6114304	Sep 05, 2017			U-637			
	6608029	Sep 07, 2013			U-641			
	6608029	Sep 07, 2013			U-640			
	6608029	Sep 07, 2013			U-637			
<u>PRASUGREL HYDROCHLORIDE - EFFIENT</u>								
N022307 001	5288726	Apr 14, 2017	DS	DP	U-991		NCE	
	6693115	Jul 03, 2021	DS	DP	U-991			Jul 10, 2014
<u>PRASUGREL HYDROCHLORIDE - EFFIENT</u>								
N022307 002	5288726	Apr 14, 2017	DS	DP	U-991		NCE	
	6693115	Jul 03, 2021	DS	DP	U-991			Jul 10, 2014
<u>PRAVASTATIN SODIUM - PRAVACHOL</u>								
N019898 002	5622985	Apr 22, 2014			U-335			
	5622985*PED	Oct 22, 2014			U-335			
<u>PRAVASTATIN SODIUM - PRAVACHOL</u>								
N019898 003	5622985	Apr 22, 2014			U-335			
	5622985*PED	Oct 22, 2014			U-335			
<u>PRAVASTATIN SODIUM - PRAVACHOL</u>								
N019898 004	5622985	Apr 22, 2014			U-335			
	5622985*PED	Oct 22, 2014			U-335			
<u>PRAVASTATIN SODIUM - PRAVACHOL</u>								
N019898 008	5622985	Apr 22, 2014			U-335			
	5622985*PED	Oct 22, 2014			U-335			
<u>PREDNISOLONE ACETATE - FLO-PRED</u>								
N022067 001	5881926	Mar 16, 2016			DP			
	6071523	Jun 03, 2018			DP			
	6102254	Mar 11, 2013			DP			
	6399079	Jun 03, 2018			DP			
	6656482	Jun 03, 2018			DP			
<u>PREDNISOLONE ACETATE - FLO-PRED</u>								
N022067 002	5881926	Mar 16, 2016			DP			
	6071523	Jun 03, 2018			DP			
	6102254	Mar 11, 2013			DP			
	6399079	Jun 03, 2018			DP			
	6656482	Jun 03, 2018			DP			
<u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u>								
N021959 001	6024981	Apr 09, 2018			DP		Y	
	6221392	Apr 09, 2018			DP		Y	
	6740341	Nov 24, 2019			DP			
<u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u>								
N021959 002	6024981	Apr 09, 2018			DP		Y	
	6221392	Apr 09, 2018			DP		Y	
	6740341	Nov 24, 2019			DP			
<u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u>								
N021959 003	6024981	Apr 09, 2018			DP		Y	
	6221392	Apr 09, 2018			DP		Y	
	6740341	Nov 24, 2019			DP			
<u>PREGABALIN - LYRICA</u>								
N021446 001	5563175	Oct 08, 2013			U-661			
	6001876	Dec 30, 2018			U-55			
	6001876	Dec 30, 2018			U-819			
	6197819	Dec 30, 2018	DS	DP				
	RE41920	Dec 30, 2018			U-1099			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PREGABALIN - LYRICA</u>						
N021446 002	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N021446 003	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N021446 004	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N021446 005	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-819	
	6001876	Dec 30, 2018			U-55	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N021446 006	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N021446 007	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N021446 008	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PREGABALIN - LYRICA</u>						
N022488 001	5563175	Oct 08, 2013			U-661	
	6001876	Dec 30, 2018			U-55	
	6001876	Dec 30, 2018			U-819	
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018			U-1099	
<u>PROCAINAMIDE HYDROCHLORIDE - PROCANBID</u>						
N020545 001	5656296	Aug 12, 2014				
<u>PROCAINAMIDE HYDROCHLORIDE - PROCANBID</u>						
N020545 002	5656296	Aug 12, 2014				
<u>PROGESTERONE - CRINONE</u>						
N020701 001	5543150	Sep 15, 2013			U-209	
<u>PROGESTERONE - CRINONE</u>						
N020701 002	5543150	Sep 15, 2013			U-209	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>PROGESTERONE - ENDOMETRIN</u>						
N022057 001	7300664	Nov 17, 2019		U-856		
	7320800	Nov 17, 2019		U-856		
	7393543	Nov 17, 2019	DP	U-880		
<u>PROPafenone Hydrochloride - RYTHMOL SR</u>						
N021416 001	5681588	Oct 28, 2014				
<u>PROPafenone Hydrochloride - RYTHMOL SR</u>						
N021416 002	5681588	Oct 28, 2014				
<u>PROPafenone Hydrochloride - RYTHMOL SR</u>						
N021416 003	5681588	Oct 28, 2014				
<u>PROPOFOL - DIPRIVAN</u>						
N019627 002	5714520	Mar 22, 2015				
	5714520*PED	Sep 22, 2015				
	5731355	Mar 22, 2015		U-217		
	5731355*PED	Sep 22, 2015		U-217		
	5731356	Mar 22, 2015		U-218		
	5731356*PED	Sep 22, 2015		U-218		
	5908869	Mar 22, 2015		U-270		
	5908869*PED	Sep 22, 2015		U-270		
<u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u>						
N021438 001	6500454	Dec 31, 2022				
<u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u>						
N021438 002	6500454	Dec 31, 2022				
<u>QUAZEPAM - DORAL</u>						
N018708 001	7608616	Jun 03, 2028		U-1012		
<u>QUAZEPAM - DORAL</u>						
N018708 003	7608616	Jun 03, 2028		U-1012		
<u>QUETIAPINE FUMARATE - SEROQUEL</u>						
N020639 001	4879288	Sep 26, 2011	DS DP	U-550	NPP	Dec 02, 2012
	4879288*PED	Mar 26, 2012			NPP	Dec 02, 2012
					PED	Jun 02, 2013
					PED	Jun 02, 2013
					PED	Nov 13, 2011
<u>QUETIAPINE FUMARATE - SEROQUEL</u>						
N020639 002	4879288	Sep 26, 2011	DS DP	U-550	NPP	Dec 02, 2012
	4879288*PED	Mar 26, 2012			NPP	Dec 02, 2012
					PED	Jun 02, 2013
					PED	Jun 02, 2013
					PED	Nov 13, 2011
<u>QUETIAPINE FUMARATE - SEROQUEL</u>						
N020639 003	4879288	Sep 26, 2011	DS DP	U-550	NPP	Dec 02, 2012
	4879288*PED	Mar 26, 2012			NPP	Dec 02, 2012
					PED	Jun 02, 2013
					PED	Jun 02, 2013
					PED	Nov 13, 2011
<u>QUETIAPINE FUMARATE - SEROQUEL</u>						
N020639 004	4879288	Sep 26, 2011	DS DP	U-550	NPP	Dec 02, 2012
	4879288*PED	Mar 26, 2012			NPP	Dec 02, 2012
					PED	Jun 02, 2013
					PED	Jun 02, 2013
					PED	Nov 13, 2011

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>QUETIAPINE FUMARATE - SEROQUEL</u>								
N020639 005	4879288	Sep	26,	2011	DS DP U-550		NPP	Dec 02, 2012
	4879288*PED	Mar	26,	2012			NPP	Dec 02, 2012
							PED	Jun 02, 2013
							PED	Jun 02, 2013
							PED	Nov 13, 2011
<u>QUETIAPINE FUMARATE - SEROQUEL</u>								
N020639 006	4879288	Sep	26,	2011	DS DP U-550		NPP	Dec 02, 2012
	4879288*PED	Mar	26,	2012			NPP	Dec 02, 2012
							PED	Jun 02, 2013
							PED	Jun 02, 2013
							PED	Nov 13, 2011
<u>QUETIAPINE FUMARATE - SEROQUEL</u>								
N020639 007	4879288	Sep	26,	2011	DS DP U-550		NPP	Dec 02, 2012
	4879288*PED	Mar	26,	2012			NPP	Dec 02, 2012
							PED	Jun 02, 2013
							PED	Jun 02, 2013
							PED	Nov 13, 2011
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>								
N022047 001	4879288	Sep	26,	2011	DS DP U-601		D-117	Oct 08, 2011
	4879288	Sep	26,	2011	DS DP U-814		I-618	Dec 02, 2012
	4879288	Sep	26,	2011	DS DP U-839		I-576	Oct 08, 2011
	4879288*PED	Mar	26,	2012			I-575	Oct 08, 2011
	5948437	May	28,	2017	DP U-839		I-574	Oct 08, 2011
	5948437	May	28,	2017	DP U-814		PED	Apr 08, 2012
	5948437	May	28,	2017	DP U-601		PED	Apr 08, 2012
	5948437*PED	Nov	28,	2017			PED	Apr 08, 2012
							PED	Apr 08, 2012
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>								
N022047 002	4879288	Sep	26,	2011	DS DP U-601		D-117	Oct 08, 2011
	4879288	Sep	26,	2011	DS DP U-814		I-618	Dec 02, 2012
	4879288	Sep	26,	2011	DS DP U-839		I-576	Oct 08, 2011
	4879288*PED	Mar	26,	2012			I-575	Oct 08, 2011
	5948437	May	28,	2017	DP U-814		I-574	Oct 08, 2011
	5948437	May	28,	2017	DP U-601		PED	Apr 08, 2012
	5948437	May	28,	2017	DP U-839		PED	Apr 08, 2012
	5948437*PED	Nov	28,	2017			PED	Apr 08, 2012
							PED	Apr 08, 2012
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>								
N022047 003	4879288	Sep	26,	2011	DS DP U-839		D-117	Oct 08, 2011
	4879288	Sep	26,	2011	DS DP U-601		I-618	Dec 02, 2012
	4879288	Sep	26,	2011	DS DP U-814		I-576	Oct 08, 2011
	4879288*PED	Mar	26,	2012			I-575	Oct 08, 2011
	5948437	May	28,	2017	DP U-601		I-574	Oct 08, 2011
	5948437	May	28,	2017	DP U-814		PED	Apr 08, 2012
	5948437	May	28,	2017	DP U-839		PED	Apr 08, 2012
	5948437*PED	Nov	28,	2017			PED	Apr 08, 2012
							PED	Apr 08, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE			PATENT DELIST REQUESTED			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>											
N022047 004	4879288	Sep	26, 2011	DS	DP	U-814		D-117	Oct	08, 2011	
	4879288	Sep	26, 2011	DS	DP	U-601		I-618	Dec	02, 2012	
	4879288	Sep	26, 2011	DS	DP	U-839		I-576	Oct	08, 2011	
	4879288*PED	Mar	26, 2012					I-575	Oct	08, 2011	
	5948437	May	28, 2017		DP	U-814		I-574	Oct	08, 2011	
	5948437	May	28, 2017		DP	U-601		PED	Apr	08, 2012	
	5948437	May	28, 2017		DP	U-839		PED	Apr	08, 2012	
	5948437*PED	Nov	28, 2017					PED	Apr	08, 2012	
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>											
N022047 005	4879288	Sep	26, 2011	DS	DP	U-814		D-117	Oct	08, 2011	
	4879288	Sep	26, 2011	DS	DP	U-601		I-618	Dec	02, 2012	
	4879288	Sep	26, 2011	DS	DP	U-839		I-576	Oct	08, 2011	
	4879288*PED	Mar	26, 2012					I-575	Oct	08, 2011	
	5948437	May	28, 2017		DP	U-839		I-574	Oct	08, 2011	
	5948437	May	28, 2017		DP	U-814		PED	Apr	08, 2012	
	5948437	May	28, 2017		DP	U-601		PED	Apr	08, 2012	
	5948437*PED	Nov	28, 2017					PED	Apr	08, 2012	
<u>QUINAPRIL HYDROCHLORIDE - ACCUPRIL</u>											
N019885 001	5684016	Nov	04, 2014			U-210					
	5684016*PED	May	04, 2015			U-210					
<u>QUINAPRIL HYDROCHLORIDE - ACCUPRIL</u>											
N019885 002	5684016	Nov	04, 2014			U-210					
	5684016*PED	May	04, 2015			U-210					
<u>QUINAPRIL HYDROCHLORIDE - ACCUPRIL</u>											
N019885 003	5684016	Nov	04, 2014			U-210					
	5684016*PED	May	04, 2015			U-210					
<u>QUINAPRIL HYDROCHLORIDE - ACCUPRIL</u>											
N019885 004	5684016	Nov	04, 2014			U-210					
	5684016*PED	May	04, 2015			U-210					
<u>QUININE SULFATE - QUALAQUIN</u>											
N021799 001								ODE	Aug	12, 2012	
<u>RABEPRAZOLE SODIUM - ACIPHEX</u>											
N020973 001	5045552	May	08, 2013			U-385					
<u>RABEPRAZOLE SODIUM - ACIPHEX</u>											
N020973 002	5045552	May	08, 2013			U-385					
<u>RALOXIFENE HYDROCHLORIDE - EVISTA</u>											
N020815 001	5393763	Jul	28, 2012			U-114	Y				
	5457117	Jul	28, 2012			U-114	Y				
	5478847	Mar	02, 2014			U-114	Y				
	5811120	Mar	02, 2014								
	5972383	Mar	02, 2014			U-287					
	6458811	Mar	10, 2017	DS	DP	U-825					
	6797719	Mar	10, 2017			DP					
	6894064	Mar	10, 2017		DP	U-657					
	6906086	Jul	28, 2012			U-662					
	6906086	Jul	28, 2012			U-657					
	8030330	Mar	10, 2017	DP							
	RE38968	Jul	28, 2012			U-662					
	RE38968	Jul	28, 2012			U-657					
	RE39049	Jul	28, 2012			U-662					
	RE39049	Jul	28, 2012			U-657					
	RE39050	Mar	02, 2014			U-657					
	RE39050	Mar	02, 2014			U-662					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>RALTEGRAVIR POTASSIUM - ISENTRESS</u>							
N022145 001	7169780	Oct	03, 2023	DS DP		NPP	Dec 21, 2014
	7217713	Oct	21, 2022		U-257	NCE	Oct 12, 2012
	7435734	Oct	21, 2022		U-900		
	7435734	Oct	21, 2022		U-257		
	7754731	Mar	11, 2029	DS DP	U-257		
<u>RALTEGRAVIR POTASSIUM - ISENTRESS</u>							
N203045 001						NDF	Dec 21, 2014
<u>RAMELTEON - ROZEREM</u>							
N021782 001	6034239	Jul	22, 2019	DS DP	U-674	M-82	Oct 20, 2011
<u>RAMIPRIL - ALTACE</u>							
N019901 001	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N019901 002	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N019901 003	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N019901 004	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N022021 001	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N022021 002	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N022021 003	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RAMIPRIL - ALTACE</u>							
N022021 004	5403856	Apr	04, 2012		U-71		
	7368469	Aug	30, 2020		U-871		
<u>RANITIDINE BISMUTH CITRATE - TRITEC</u>							
N020559 001	5403830	Apr	04, 2012		U-200		
	5407688	Apr	04, 2012		U-201		
	5456925	Oct	10, 2012				
	5601848	Feb	11, 2014		U-202		
	5629297	May	13, 2014		U-186		
<u>RANOLAZINE - RANEXA</u>							
N021526 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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>RANOLAZINE - RANEXA</u>							
N021526 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		
<u>RAPACURONIUM BROMIDE - RAPLON</u>							
N020984 001	5418226	Apr	14, 2013				
<u>RAPACURONIUM BROMIDE - RAPLON</u>							
N020984 002	5418226	Apr	14, 2013				
<u>RASAGILINE MESYLATE - AZILECT</u>							
N021641 001	5387612	Feb	07, 2012		U-219		
	5453446	Feb	07, 2017		U-219		
	5457133	Feb	07, 2012	DS	DP		
	5532415	Jul	02, 2013	DS			
	5786390	Feb	07, 2012		DP		
	6126968	Sep	18, 2016		DP		
	7572834	Dec	05, 2026		DP		
	7815942	Aug	27, 2027	DS	DP	U-219	
<u>RASAGILINE MESYLATE - AZILECT</u>							
N021641 002	5387612	Feb	07, 2012		U-219		
	5453446	Feb	07, 2017		U-219		
	5457133	Feb	07, 2012	DS	DP		
	5532415	Jul	02, 2013	DS			
	5786390	Feb	07, 2012		DP		
	6126968	Sep	18, 2016		DP		
	7572834	Dec	05, 2026		DP		
	7815942	Aug	27, 2027	DS	DP	U-219	
<u>REGADENOSON - LEXISCAN</u>							
N022161 001	6403567	Apr	10, 2022	DS	DP	U-869	
	6642210	Jun	22, 2019	DS	DP	U-869	NCE
	7144872	Jun	22, 2019	DS	DP	U-870	Apr 10, 2013
	7144872	Jun	22, 2019	DS	DP	U-869	
	7144872	Jun	22, 2019	DS	DP	U-116	
	7183264	Jun	22, 2019		DP	U-870	
	7183264	Jun	22, 2019		DP	U-869	
	7183264	Jun	22, 2019		DP	U-116	
	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	
<u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u>							
N020630 001	5866591	Sep	10, 2017		DP		
	5866591*PED	Mar	10, 2018				
<u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u>							
N020630 002	5866591	Sep	10, 2017		DP		
	5866591*PED	Mar	10, 2018				
<u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u>							
N020630 003	5866591	Sep	10, 2017		DP		
	5866591*PED	Mar	10, 2018				
<u>REPAGLINIDE - PRANDIN</u>							
N020741 001	6677358	Jun	12, 2018	DS	DP	U-968	
<u>REPAGLINIDE - PRANDIN</u>							
N020741 002	6677358	Jun	12, 2018	DS	DP	U-968	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>REPAGLINIDE - PRANDIN</u>						
N020741 003	6677358	Jun 12, 2018	DS DP U-968			
<u>RETAPAMULIN - ALTABAX</u>						
N022055 001	7875630	Feb 14, 2027	DS		NCE	Apr 12, 2012
	RE39128	Apr 12, 2021	DS DP U-805			
<u>RIBAVIRIN - COPEGUS</u>						
N021511 001					NPP	Aug 09, 2014
<u>RIBAVIRIN - REBETOL</u>						
N020903 001	6172046	Sep 21, 2017	U-1014			
	6172046	Sep 21, 2017	U-377			
	6172046*PED	Mar 21, 2018	U-377			
	6177074	Nov 01, 2016	U-1013			
	6177074	Nov 01, 2016	U-454			
	6177074*PED	May 01, 2017	U-454			
	6461605	Nov 01, 2016	U-478			
	6461605	Nov 01, 2016	U-1013			
	6461605*PED	May 01, 2017	U-478			
	6472373	Sep 21, 2017	U-479			
	6472373	Sep 21, 2017	U-1014			
	6472373*PED	Mar 21, 2018	U-479			
	6524570	Nov 01, 2016	U-499			
	6524570	Nov 01, 2016	U-1013			
	6524570*PED	May 01, 2017	U-499			
<u>RIBAVIRIN - REBETOL</u>						
N020903 002	6172046	Sep 21, 2017	U-377			
	6172046	Sep 21, 2017	U-1014			
	6172046*PED	Mar 21, 2018	U-377			
	6177074	Nov 01, 2016	U-1013			
	6177074	Nov 01, 2016	U-454			
	6177074*PED	May 01, 2017	U-454			
	6461605	Nov 01, 2016	U-1013			
	6461605	Nov 01, 2016	U-478			
	6461605*PED	May 01, 2017	U-478			
	6472373	Sep 21, 2017	U-1014			
	6472373	Sep 21, 2017	U-479			
	6472373*PED	Mar 21, 2018	U-479			
	6524570	Nov 01, 2016	U-499			
	6524570	Nov 01, 2016	U-1013			
	6524570*PED	May 01, 2017	U-499			
<u>RIBAVIRIN - REBETOL</u>						
N021546 001	6172046	Sep 21, 2017	U-521			
	6172046	Sep 21, 2017	U-1014			
	6172046*PED	Mar 21, 2018	U-521			
	6177074	Nov 01, 2016	U-1013			
	6177074*PED	May 01, 2017				
	6461605	Nov 01, 2016	U-521			
	6461605	Nov 01, 2016	U-1013			
	6461605*PED	May 01, 2017	U-521			
	6472373	Sep 21, 2017	U-521			
	6472373*PED	Mar 21, 2018	U-521			
	6524570	Nov 01, 2016	U-1013			
	6524570*PED	May 01, 2017				
	6790837	Apr 05, 2023	DP			
	6790837*PED	Oct 05, 2023	U-1014			
<u>RIBAVIRIN - VIRAZOLE</u>						
N018859 001	6150337	Nov 21, 2017	U-400			
<u>RIFAXIMIN - XIFAXAN</u>						
N021361 001	7045620	Jun 19, 2024	DS DP			
	7612199	Jun 19, 2024	DS DP			
	7902206	Jun 19, 2024	DS DP			
	7906542	Jun 01, 2025	DS DP			
	7928115	Jul 24, 2029	U-1121			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>RIFAXIMIN - XIFAXAN</u>							
N022554 001	7045620	Jun	19, 2024	DS		NP	Mar 24, 2013
	7612199	Jun	19, 2024	DS	DP	ODE	Mar 24, 2017
	7902206	Jun	19, 2024	DS	DP		
	7906542	Jun	01, 2025	DS	DP		
<u>RILPIVIRINE HYDROCHLORIDE - EDURANT</u>							
N202022 001	6838464	Feb	26, 2021	DS	DP	NCE	May 20, 2016
	7067522	Dec	20, 2019	DS	DP		
	7125879	Apr	14, 2023	DS	DP	U-1153	
	7638522	Apr	14, 2023		DP		
<u>RILUZOLE - RILUTEK</u>							
N020599 001	5527814	Jun	18, 2013				
<u>RISEDRONATE SODIUM - ACTONEL</u>							
N020835 001	5583122	Dec	10, 2013	U-222		M-61	Jul 23, 2012
	5583122*PED	Jun	10, 2014			PED	Jan 23, 2013
	6096342	Nov	22, 2011				
	6096342*PED	May	22, 2012				
	6165513	Jun	10, 2018				
	6165513*PED	Dec	10, 2018				
<u>RISEDRONATE SODIUM - ACTONEL</u>							
N020835 002	5583122	Dec	10, 2013	U-222		M-61	Jul 23, 2012
	5583122*PED	Jun	10, 2014			PED	Jan 23, 2013
	6096342	Nov	22, 2011				
	6096342*PED	May	22, 2012				
	6165513	Jun	10, 2018				
	6165513*PED	Dec	10, 2018				
<u>RISEDRONATE SODIUM - ACTONEL</u>							
N020835 003	5583122	Dec	10, 2013	DS	DP	U-756	M-61 Jul 23, 2012
	5583122	Dec	10, 2013	DS	DP	U-222	PED Jan 23, 2013
	5583122*PED	Jun	10, 2014				
	5994329	Jul	17, 2018	U-353			
	5994329*PED	Jan	17, 2019				
	6015801	Jul	17, 2018	U-353			
	6015801*PED	Jan	17, 2019				
	6096342	Nov	22, 2011	DP			
	6096342*PED	May	22, 2012				
	6165513	Jun	10, 2018	DP			
	6165513*PED	Dec	10, 2018				
	6432932	Jul	17, 2018	U-595			
	6432932*PED	Jan	17, 2019				
	6465443	Aug	14, 2018	DP			
	6465443*PED	Feb	14, 2019				
<u>RISEDRONATE SODIUM - ACTONEL</u>							
N020835 004	5583122	Dec	10, 2013	DS	DP	U-353	M-61 Jul 23, 2012
	5583122*PED	Jun	10, 2014				
	6096342	Nov	22, 2011	DP	U-353		PED Jan 23, 2013
	6096342*PED	May	22, 2012				
	6165513	Jun	10, 2018	DP			
	6165513*PED	Dec	10, 2018				
<u>RISEDRONATE SODIUM - ACTONEL</u>							
N020835 005	5583122	Dec	10, 2013	DS	DP	U-353	M-61 Jul 23, 2012
	5583122*PED	Jun	10, 2014				
	6165513	Jun	10, 2018	DP			
	6165513*PED	Dec	10, 2018				
	7192938	May	06, 2023	U-353			
	7192938*PED	Nov	06, 2023				
	7718634	May	06, 2023	U-662			
	7718634*PED	Nov	06, 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
RISEDRONATE SODIUM - ATELVIA								
N022560 001	5583122	Dec	10, 2013	DS	DP U-662		NDF	Oct 08, 2013
	5583122*PED	Jun	10, 2014					
	5622721	Apr	21, 2014		DP U-662			
	7645459	Jan	09, 2028		DP U-662			
	7645460	Jan	09, 2028		DP U-662			
RISPERIDONE - RISPERDAL								
N020588 001	5453425	Jul	11, 2014		DP			
	5453425*PED	Jan	11, 2015					
	5616587	Jul	11, 2014				Y	
	5616587*PED	Jan	11, 2015					
	RE39181	Jul	11, 2014		DP			
	RE39181*PED	Jan	11, 2015					
RISPERIDONE - RISPERDAL								
N021444 001	5648093	Jul	15, 2014		DP			
	5648093*PED	Jan	15, 2015					
	6224905	Jun	10, 2017		DP			
	6224905*PED	Dec	10, 2017					
RISPERIDONE - RISPERDAL								
N021444 002	5648093	Jul	15, 2014		DP			
	5648093*PED	Jan	15, 2015					
	6224905	Jun	10, 2017		DP			
	6224905*PED	Dec	10, 2017					
RISPERIDONE - RISPERDAL								
N021444 003	5648093	Jul	15, 2014		DP			
	5648093*PED	Jan	15, 2015					
	6224905	Jun	10, 2017		DP			
	6224905*PED	Dec	10, 2017					
RISPERIDONE - RISPERDAL								
N021444 004	5648093	Jul	15, 2014		DP			
	5648093*PED	Jan	15, 2015					
	6224905	Jun	10, 2017		DP			
	6224905*PED	Dec	10, 2017					
RISPERIDONE - RISPERDAL								
N021444 005	5648093	Jul	15, 2014		DP			
	5648093*PED	Jan	15, 2015					
	6224905	Jun	10, 2017		DP			
	6224905*PED	Dec	10, 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
RISPERIDONE - RISPERDAL CONSTA						
N021346 001	5688801	Nov 18, 2014	U-543		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-972		I-596	May 15, 2012
	5688801*PED	May 18, 2015				
	5792477	May 02, 2017	DP			
	5792477*PED	Nov 02, 2017				
	5916598	May 02, 2017	DP			
	5916598*PED	Nov 02, 2017				
	5965168	Nov 19, 2013	DP			
	5965168*PED	May 19, 2014				
	6110921	Nov 19, 2013	U-543			
	6110921*PED	May 19, 2014				
	6194006	Dec 30, 2018	DP			
	6194006*PED	Jun 30, 2019				
	6368632	Nov 19, 2013	U-543			
	6368632*PED	May 19, 2014				
	6379703	Dec 30, 2018	DP			
	6379703*PED	Jun 30, 2019				
	6403114	May 02, 2017	DP			
	6403114*PED	Nov 02, 2017				
	6596316	Dec 30, 2018	DP			
	6596316*PED	Jun 30, 2019				
	6667061	May 25, 2020	DP			
	6667061*PED	Nov 25, 2020				
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				
RISPERIDONE - RISPERDAL CONSTA						
N021346 002	5688801	Nov 18, 2014	U-543		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-972		I-596	May 15, 2012
	5688801*PED	May 18, 2015				
	5792477	May 02, 2017	DP			
	5792477*PED	Nov 02, 2017				
	5916598	May 02, 2017	DP			
	5916598*PED	Nov 02, 2017				
	5965168	Nov 19, 2013	DP			
	5965168*PED	May 19, 2014				
	6110921	Nov 19, 2013	U-543			
	6110921*PED	May 19, 2014				
	6194006	Dec 30, 2018	DP			
	6194006*PED	Jun 30, 2019				
	6368632	Nov 19, 2013	U-543			
	6368632*PED	May 19, 2014				
	6379703	Dec 30, 2018	DP			
	6379703*PED	Jun 30, 2019				
	6403114	May 02, 2017	DP			
	6403114*PED	Nov 02, 2017				
	6596316	Dec 30, 2018	DP			
	6596316*PED	Jun 30, 2019				
	6667061	May 25, 2020	DP			
	6667061*PED	Nov 25, 2020				
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
RISPERIDONE - RISPERDAL CONSTA						
N021346 003	5688801	Nov 18, 2014	U-543		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-972		I-596	May 15, 2012
	5688801*PED	May 18, 2015				
	5792477	May 02, 2017	DP			
	5792477*PED	Nov 02, 2017				
	5916598	May 02, 2017	DP			
	5916598*PED	Nov 02, 2017				
	5965168	Nov 19, 2013	DP			
	5965168*PED	May 19, 2014				
	6110921	Nov 19, 2013	U-543			
	6110921*PED	May 19, 2014				
	6194006	Dec 30, 2018	DP			
	6194006*PED	Jun 30, 2019				
	6368632	Nov 19, 2013	U-543			
	6368632*PED	May 19, 2014				
	6379703	Dec 30, 2018	DP			
	6379703*PED	Jun 30, 2019				
	6403114	May 02, 2017	DP			
	6403114*PED	Nov 02, 2017				
	6596316	Dec 30, 2018	DP			
	6596316*PED	Jun 30, 2019				
	6667061	May 25, 2020	DP			
	6667061*PED	Nov 25, 2020				
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				
RISPERIDONE - RISPERDAL CONSTA						
N021346 004	5688801	Nov 18, 2014	U-972		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-543		I-596	May 15, 2012
	5688801*PED	May 18, 2015				
	5792477	May 02, 2017	DP			
	5792477*PED	Nov 02, 2017				
	5916598	May 02, 2017	DP			
	5916598*PED	Nov 02, 2017				
	5965168	Nov 19, 2013	DP			
	5965168*PED	May 19, 2014				
	6110503	May 02, 2017	DP			
	6110503*PED	Nov 02, 2017				
	6110921	Nov 19, 2013	U-543			
	6110921*PED	May 19, 2014				
	6194006	Dec 30, 2018	DP			
	6194006*PED	Jun 30, 2019				
	6368632	Nov 19, 2013	U-543			
	6368632*PED	May 19, 2014				
	6379703	Dec 30, 2018	DP			
	6379703*PED	Jun 30, 2019				
	6403114	May 02, 2017	DP			
	6403114*PED	Nov 02, 2017				
	6596316	Dec 30, 2018	DP			
	6596316*PED	Jun 30, 2019				
	6667061	May 25, 2020	DP			
	6667061*PED	Nov 25, 2020				
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
RITONAVIR - NORVIR						
N020659 001	5484801	Jan 28, 2014				
	5484801*PED	Jul 28, 2014				
	5541206	Jul 30, 2013		U-140		
	5541206*PED	Jan 30, 2014				
	5635523	Jul 30, 2013		U-190		
	5635523*PED	Jan 30, 2014				
	5648497	Jul 15, 2014				
	5648497*PED	Jan 15, 2015				
	5674882	Jul 30, 2013		U-688		
	5674882*PED	Jan 30, 2014				
	5948436	Sep 13, 2013		DP		
	5948436*PED	Mar 13, 2014				
	6037157	Jun 26, 2016				
	6037157*PED	Dec 26, 2016				
	6703403	Jun 26, 2016		U-564		
	6703403*PED	Dec 26, 2016				
RITONAVIR - NORVIR						
N020680 001	5541206	Jul 30, 2013		U-140		
	5541206*PED	Jan 30, 2014				
	5635523	Jun 03, 2014		U-190		
	5635523*PED	Dec 03, 2014				
	5648497	Jul 15, 2014				
	5648497*PED	Jan 15, 2015				
	5948436	Sep 13, 2013				
	5948436*PED	Mar 13, 2014				
RITONAVIR - NORVIR						
N020945 001	5541206	Jul 30, 2013		U-348		
	5541206*PED	Jan 30, 2014				
	5635523	Jul 30, 2013		U-347		
	5635523*PED	Jan 30, 2014				
	5648497	Jul 15, 2014				
	5648497*PED	Jan 15, 2015				
	5674882	Jul 30, 2013		U-895		
	5674882*PED	Jan 30, 2014				
	5948436	Sep 13, 2013		DP		
	5948436*PED	Mar 13, 2014				
	6037157	Jun 26, 2016		U-895		
	6037157*PED	Dec 26, 2016				
	6232333	Nov 07, 2017				
	6232333*PED	May 07, 2018				
	6703403	Jun 26, 2016		U-564		
	6703403*PED	Dec 26, 2016				
	7141593	May 22, 2020		DP		
	7141593*PED	Nov 22, 2020				
	7432294	May 22, 2020		DP		
	7432294*PED	Nov 22, 2020				
RITONAVIR - NORVIR						
N022417 001	5541206	Jul 30, 2013	DS	DP	U-688	
	5541206*PED	Jan 30, 2014				
	5635523	Jul 30, 2013		U-688		
	5635523*PED	Jan 30, 2014				
	5648497	Jul 15, 2014	DS			
	5648497*PED	Jan 15, 2015				
	5674882	Jul 30, 2013		U-688		
	5674882*PED	Jan 30, 2014				
	6037157	Jun 26, 2016		U-688		
	6037157*PED	Dec 26, 2016				
	6703403	Jun 26, 2016		U-688		
	6703403*PED	Dec 26, 2016				
	7148359	Jul 19, 2019		DP		
	7148359*PED	Jan 19, 2020				
	7364752	Nov 10, 2020		DP	U-688	
	7364752*PED	May 10, 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
RIVAROXABAN - XARELTO							
N022406 001	7157456	Feb 08, 2021	DS	DP		NCE	Jul 01, 2016
	7585860	Dec 11, 2020	DS				
	7592339	Dec 11, 2020		U-1167			
RIVAROXABAN - XARELTO							
N202439 001	7157456	Feb 08, 2021	DS	DP		I-643	Nov 04, 2014
	7585860	Dec 11, 2020	DS			NCE	Jul 01, 2016
	7592339	Dec 11, 2020		U-1167			
	7592339	Dec 11, 2020		U-1200			
RIVAROXABAN - XARELTO							
N202439 002	7157456	Feb 08, 2021	DS	DP		I-643	Nov 04, 2014
	7585860	Dec 11, 2020	DS			NCE	Jul 01, 2016
	7592339	Dec 11, 2020		U-1167			
	7592339	Dec 11, 2020		U-1200			
RIVASTIGMINE - EXELON							
N022083 001	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014	DS	DP U-322			
	6316023	Jan 08, 2019		DP			
	6335031	Jan 08, 2019		DP			
RIVASTIGMINE - EXELON							
N022083 002	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014	DS	DP U-322			
	6316023	Jan 08, 2019		DP			
	6335031	Jan 08, 2019		DP			
RIVASTIGMINE TARTRATE - EXELON							
N020823 003	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014		U-322			
RIVASTIGMINE TARTRATE - EXELON							
N020823 004	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014		U-322			
RIVASTIGMINE TARTRATE - EXELON							
N020823 005	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014		U-322			
RIVASTIGMINE TARTRATE - EXELON							
N020823 006	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014		U-322			
RIVASTIGMINE TARTRATE - EXELON							
N021025 001	4948807	Aug 14, 2012	DS	U-322			
	5602176	Feb 11, 2014		U-322			
RIZATRIPTAN BENZOATE - MAXALT							
N020864 001	5298520	Jun 29, 2012	DS	DP U-240		NPP	Dec 15, 2014
	5298520*PED	Dec 29, 2012				PED	Jun 15, 2015
	5602162	Feb 11, 2014					
	5602162*PED	Aug 11, 2014			Y		
RIZATRIPTAN BENZOATE - MAXALT							
N020864 002	5298520	Jun 29, 2012	DS	DP U-240		NPP	Dec 15, 2014
	5298520*PED	Dec 29, 2012				PED	Jun 15, 2015
	5602162	Feb 11, 2014					
	5602162*PED	Aug 11, 2014			Y		
RIZATRIPTAN BENZOATE - MAXALT-MLT							
N020865 001	5298520	Jun 29, 2012	DS	DP U-240		NPP	Dec 15, 2014
	5298520*PED	Dec 29, 2012				PED	Jun 15, 2015
	5457895	Oct 01, 2013		DP			
	5457895*PED	Apr 01, 2014					
	5602162	Feb 11, 2014		U-240			
	5602162*PED	Aug 11, 2014			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
<u>RIZATRIPTAN BENZOATE - MAXALT-MLT</u>								
N020865 002	5298520	Jun	29,	2012	DS DP U-240		NPP	Dec 15, 2014
	5298520*PED	Dec	29,	2012			PED	Jun 15, 2015
	5457895	Oct	01,	2013	DP			
	5457895*PED	Apr	01,	2014				
	5602162	Feb	11,	2014	U-240	Y		
	5602162*PED	Aug	11,	2014				
<u>ROCURONIUM BROMIDE - ZEMURON</u>								
N020214 001							NPP	Aug 28, 2011
							PED	Feb 28, 2012
<u>ROCURONIUM BROMIDE - ZEMURON</u>								
N020214 002							NPP	Aug 28, 2011
							PED	Feb 28, 2012
<u>ROCURONIUM BROMIDE - ZEMURON</u>								
N020214 003							NPP	Aug 28, 2011
							PED	Feb 28, 2012
<u>ROFECOXIB - VIOXX</u>								
N021042 001	5474995	Jun	24,	2013	DS DP U-602			
	5474995*PED	Dec	24,	2013				
	5691374	May	18,	2015				
	5691374*PED	Nov	18,	2015				
	6063811	May	06,	2017	U-602			
	6063811*PED	Nov	06,	2017				
	6239173	Jun	24,	2013	DS DP U-602			
	6239173*PED	Dec	24,	2013				
<u>ROFECOXIB - VIOXX</u>								
N021042 002	5474995	Jun	24,	2013	DS DP U-602			
	5474995*PED	Dec	24,	2013				
	5691374	May	18,	2015				
	5691374*PED	Nov	18,	2015				
	6063811	May	06,	2017	U-602			
	6063811*PED	Nov	06,	2017				
	6239173	Jun	24,	2013	DS DP U-602			
	6239173*PED	Dec	24,	2013				
<u>ROFECOXIB - VIOXX</u>								
N021042 003	5474995	Jun	24,	2013	DS DP U-602			
	5474995*PED	Dec	24,	2013				
	5691374	May	18,	2015				
	5691374*PED	Nov	18,	2015				
	6063811	May	06,	2017	U-602			
	6063811*PED	Nov	06,	2017				
	6239173	Jun	24,	2013	DS DP U-602			
	6239173*PED	Dec	24,	2013				
<u>ROFECOXIB - VIOXX</u>								
N021052 001	5474995	Jun	24,	2013	U-266			
	5474995*PED	Dec	24,	2013				
	5691374	May	18,	2015				
	5691374*PED	Nov	18,	2015				
	6063811	May	06,	2017	U-266			
	6063811*PED	Nov	06,	2017				
	6239173	Jun	24,	2013	DS DP U-602			
	6239173*PED	Dec	24,	2013				
<u>ROFECOXIB - VIOXX</u>								
N021052 002	5474995	Jun	24,	2013	U-266			
	5474995*PED	Dec	24,	2013				
	5691374	May	18,	2015				
	5691374*PED	Nov	18,	2015				
	6063811	May	06,	2017	U-266			
	6063811*PED	Nov	06,	2017				
	6239173	Jun	24,	2013	DS DP U-602			
	6239173*PED	Dec	24,	2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ROFLUMILAST - DALIRESP</u>						
N022522 001	5712298	Jan 27, 2015	DS DP U-1115		NCE	Feb 28, 2016
<u>ROMIDEPSIN - ISTODAX</u>						
N022393 001	4977138	Jul 06, 2012	DS DP		NCE	Nov 05, 2014
	7608280	Aug 22, 2021	DS		ODE	Jun 16, 2018
	7611724	Aug 22, 2021	DS		ODE	Nov 05, 2016
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N022008 001	5422123	Jun 06, 2012	DP			
	7927624	Dec 02, 2021	DP U-20			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N022008 002	5422123	Jun 06, 2012	DP			
	7927624	Dec 02, 2021	DP U-20			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N022008 003	5422123	Jun 06, 2012	DP			
	7927624	Dec 02, 2021	DP U-20			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N022008 004	5422123	Jun 06, 2012	DP			
	7927624	Dec 02, 2021	DP U-20			
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
N022008 005	5422123	Jun 06, 2012	DP			
	7927624	Dec 02, 2021	DP U-20			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE - NAROPIN</u>						
N020533 001	5670524	May 26, 2014	DS DP U-833			
	5834489	May 26, 2014	DS DP U-838			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE - NAROPIN</u>						
N020533 003	5670524	May 26, 2014	DS DP U-833			
	5834489	May 26, 2014	DS DP U-838			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE - NAROPIN</u>						
N020533 004	5670524	May 26, 2014	DS DP U-833			
	5834489	May 26, 2014	DS DP U-838			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE - NAROPIN</u>						
N020533 005	5670524	May 26, 2014	DS DP U-833			
	5834489	May 26, 2014	DS DP U-838			
<u>ROSIGLITAZONE MALEATE - AVANDIA</u>						
N021071 002	5002953	Sep 17, 2011	DS DP U-840			
	5002953	Sep 17, 2011	DS DP U-329			
	5002953	Sep 17, 2011	DS DP U-628			
	5002953*PED	Mar 17, 2012				
	5741803	Apr 21, 2015	DS DP U-628	Y		
	5741803	Apr 21, 2015	DS DP U-329	Y		
	5741803*PED	Oct 21, 2015				
	6288095	Feb 11, 2017	U-420	Y		
	6288095*PED	Aug 11, 2017				
	7358366	Apr 19, 2020	DS			
	7358366*PED	Oct 19, 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>ROSIGLITAZONE MALEATE - AVANDIA</u>								
N021071 003	5002953	Sep	17,	2011	DS DP U-840			
	5002953	Sep	17,	2011	DS DP U-329			
	5002953	Sep	17,	2011	DS DP U-628			
	5002953*PED	Mar		17, 2012				
	5741803	Apr	21,	2015	DS DP U-628	Y		
	5741803	Apr	21,	2015	DS DP U-329	Y		
	5741803*PED	Oct		21, 2015				
	6288095	Feb	11,	2017	U-420	Y		
	6288095*PED	Aug	11,	2017				
	7358366	Apr	19,	2020	DS			
	7358366*PED	Oct	19,	2020				
<u>ROSIGLITAZONE MALEATE - AVANDIA</u>								
N021071 004	5002953	Sep	17,	2011	DS DP U-329			
	5002953	Sep	17,	2011	DS DP U-628			
	5002953	Sep	17,	2011	DS DP U-840			
	5002953*PED	Mar		17, 2012				
	5741803	Apr	21,	2015	DS DP U-628	Y		
	5741803	Apr	21,	2015	DS DP U-329	Y		
	5741803*PED	Oct		21, 2015				
	6288095	Feb	11,	2017	U-420	Y		
	6288095*PED	Aug	11,	2017				
	7358366	Apr	19,	2020	DS			
	7358366*PED	Oct	19,	2020				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>								
N021366 002	6316460	Aug	04,	2020	DP		I-621	Feb 08, 2013
	6316460*PED	Feb	04,	2021			I-611	Oct 16, 2012
	6858618	Dec	17,	2021	U-618		I-573	Nov 06, 2011
	6858618*PED	Jun	17,	2022			PED	Apr 16, 2013
	7030152	Apr	02,	2018	U-1032		PED	May 06, 2012
	7030152*PED	Oct	02,	2018				
	7964614	Apr	02,	2018	U-1032			
	7964614*PED	Oct	02,	2018				
	RE37314	Jan	08,	2016	DS			
	RE37314*PED	Jul	08,	2016				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>								
N021366 003	6316460	Aug	04,	2020	DP		I-621	Feb 08, 2013
	6316460*PED	Feb	04,	2021			I-611	Oct 16, 2012
	6858618	Dec	17,	2021	U-618		I-573	Nov 06, 2011
	6858618*PED	Jun	17,	2022			PED	Apr 16, 2013
	7030152	Apr	02,	2018	U-1032		PED	May 06, 2012
	7030152*PED	Oct	02,	2018				
	7964614	Apr	02,	2018	U-1032			
	7964614*PED	Oct	02,	2018				
	RE37314	Jan	08,	2016	DS			
	RE37314*PED	Jul	08,	2016				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>								
N021366 004	6316460	Aug	04,	2020	DP		I-621	Feb 08, 2013
	6316460*PED	Feb	04,	2021			I-611	Oct 16, 2012
	6858618	Dec	17,	2021	U-618		I-573	Nov 06, 2011
	6858618*PED	Jun	17,	2022			PED	Apr 16, 2013
	7030152	Apr	02,	2018	U-1032		PED	May 06, 2012
	7030152*PED	Oct	02,	2018				
	7964614	Apr	02,	2018	U-1032			
	7964614*PED	Oct	02,	2018				
	RE37314	Jan	08,	2016	DS			
	RE37314*PED	Jul	08,	2016				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>ROSUVASTATIN CALCIUM - CRESTOR</u>								
N021366 005	6316460	Aug	04, 2020	DP		I-621	Feb	08, 2013
	6316460*PED	Feb	04, 2021			I-611	Oct	16, 2012
	6858618	Dec	17, 2021	U-618		I-573	Nov	06, 2011
	6858618*PED	Jun	17, 2022			PED	Apr	16, 2013
	7030152	Apr	02, 2018	U-1032		PED	May	06, 2012
	7030152*PED	Oct	02, 2018					
	7964614	Apr	02, 2018	U-1032				
	7964614*PED	Oct	02, 2018					
	RE37314	Jan	08, 2016	DS				
	RE37314*PED	Jul	08, 2016					
<u>ROTIGOTINE - NEUPRO</u>								
N021829 001	6699498	Nov	27, 2020	DP		NCE	May	09, 2012
	6884434	Mar	30, 2021	DP				
	7413747	Mar	18, 2019	DP				
<u>ROTIGOTINE - NEUPRO</u>								
N021829 002	6699498	Nov	27, 2020	DP		NCE	May	09, 2012
	6884434	Mar	30, 2021	DP				
	7413747	Mar	18, 2019	DP				
<u>ROTIGOTINE - NEUPRO</u>								
N021829 003	6699498	Nov	27, 2020	DP		NCE	May	09, 2012
	6884434	Mar	30, 2021	DP				
	7413747	Mar	18, 2019	DP				
<u>RUFINAMIDE - BANZEL</u>								
N021911 001	6740669	Nov	14, 2022	DS	DP	NCE	Nov	14, 2013
	8076362	Jun	08, 2018		DP	ODE	Nov	14, 2015
<u>RUFINAMIDE - BANZEL</u>								
N021911 002	6740669	Nov	14, 2022	DS	DP	NCE	Nov	14, 2013
	8076362	Jun	08, 2018		DP	ODE	Nov	14, 2015
<u>RUFINAMIDE - BANZEL</u>								
N021911 003	6740669	Nov	14, 2022	DS	DP	NCE	Nov	14, 2013
	8076362	Jun	08, 2018		DP	ODE	Nov	14, 2015
<u>RUFINAMIDE - BANZEL</u>								
N201367 001	6740669	Aug	17, 2020	DS	DP	NCE	Nov	14, 2013
						ODE	Nov	14, 2015
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>								
N202192 001	7598257	Dec	24, 2027	DS	DP	U-1201	NCE	Nov 16, 2016
							ODE	Nov 16, 2018
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>								
N202192 002	7598257	Dec	24, 2027	DS	DP	U-1201	NCE	Nov 16, 2016
							ODE	Nov 16, 2018
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>								
N202192 003	7598257	Dec	24, 2027	DS	DP	U-1201	NCE	Nov 16, 2016
							ODE	Nov 16, 2018
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>								
N202192 004	7598257	Dec	24, 2027	DS	DP	U-1201	NCE	Nov 16, 2016
							ODE	Nov 16, 2018
<u>RUXOLITINIB PHOSPHATE - JAKAFI</u>								
N202192 005	7598257	Dec	24, 2027	DS	DP	U-1201	NCE	Nov 16, 2016
							ODE	Nov 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
SALMETEROL XINAFOATE - SEREVENT							
N020692 001	5590645*PED	Sep	01, 2011				
	5860419*PED	Sep	01, 2011				
	5873360	Feb	23, 2016	DP			
	5873360*PED	Aug	23, 2016				
	6032666*PED	Sep	01, 2011				
	6378519*PED	Sep	01, 2011				
	6536427*PED	Sep	01, 2011				
	6792945*PED	Sep	01, 2011				
	7225808*PED	Sep	01, 2011				
	7389775*PED	Sep	01, 2011				
SAPROPTERIN DIHYDROCHLORIDE - KUVAN							
N022181 001	7566462	Nov	16, 2025	DP		NCE	Dec 13, 2012
	7566714	Nov	17, 2024		U-989	ODE	Dec 13, 2014
	7612073	Nov	17, 2024		U-1010		
	7727987	Nov	17, 2024	DP			
	7947681	Nov	17, 2024		U-1156		
	8003126	Nov	16, 2025				
	8067416	Nov	17, 2024		U-989		
SAQUINAVIR - FORTOVASE							
N020828 001	6008228	Jun	06, 2015				
	6008228*PED	Dec	06, 2015				
	6352717	Nov	16, 2019				
	6352717*PED	May	16, 2020				
SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA							
N022350 001	6395767	Feb	16, 2021	DS	DP U-995	M-108	Dec 16, 2013
	7951400	Nov	30, 2028	DP		NCE	Jul 31, 2014
SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA							
N022350 002	6395767	Feb	16, 2021	DS	DP U-995	M-108	Dec 16, 2013
	7951400	Nov	30, 2028	DP		NCE	Jul 31, 2014
SELEGILINE - EMSAM							
N021336 001	7070808	May	10, 2018	DS	DP		
	7150881	Jun	12, 2018	DS	DP		
	7638140	May	10, 2018		DP		
SELEGILINE - EMSAM							
N021336 002	7070808	May	10, 2018	DS	DP		
	7150881	Jun	12, 2018	DS	DP		
	7638140	May	10, 2018		DP		
SELEGILINE - EMSAM							
N021336 003	7070808	May	10, 2018	DS	DP		
	7150881	Jun	12, 2018	DS	DP		
	7638140	May	10, 2018		DP		
SELEGILINE HYDROCHLORIDE - ZELAPAR							
N021479 001	5648093	Jul	15, 2014		DP		
	6423342	Mar	01, 2016		DP		
SERTACONAZOLE NITRATE - ERTACZO							
N021385 001	5135943	May	31, 2014	DS	DP U-786		
SERTRALINE HYDROCHLORIDE - ZOLOFT							
N019839 001	5248699	Aug	13, 2012				
	5248699*PED	Feb	13, 2013				
SERTRALINE HYDROCHLORIDE - ZOLOFT							
N019839 002	5248699	Aug	13, 2012				
	5248699*PED	Feb	13, 2013				
SERTRALINE HYDROCHLORIDE - ZOLOFT							
N019839 003	5248699	Aug	13, 2012				
	5248699*PED	Feb	13, 2013				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>SERTRALINE HYDROCHLORIDE - ZOLOFT</u>						
N019839 004	5248699	Aug 13, 2012				
	5248699*PED	Feb 13, 2013				
<u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u>						
N019839 005	5248699	Aug 13, 2012				
	5248699*PED	Feb 13, 2013				
<u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u>						
N020990 001	5248699	Aug 13, 2012				
	5248699*PED	Feb 13, 2013				
	6727283	Oct 11, 2019	DP	U-580		
	6727283*PED	Apr 11, 2020				
	7067555	Nov 10, 2019	DP			
	7067555*PED	May 10, 2020				
<u>SEVELAMER CARBONATE - RENVELA</u>						
N022127 001	5496545	Aug 11, 2013	DP	U-246		
	5667775	Sep 16, 2014		U-246		
	6509013	Aug 11, 2013	DP			
	6858203	Aug 11, 2013	DP	U-246		
	7014846	Aug 11, 2013	DP	U-246		
	7459151	Aug 11, 2013		U-246		
	7985418	Oct 27, 2025	DP			
<u>SEVELAMER CARBONATE - RENVELA</u>						
N022318 001	5496545	Aug 11, 2013	DP	U-246	NDF	
	5667775	Sep 16, 2014		U-246		Aug 12, 2012
	6509013	Aug 11, 2013	DP			
	6858203	Aug 11, 2013	DP	U-246		
	7014846	Aug 11, 2013	DP	U-246		
	7459151	Aug 11, 2013		U-246		
<u>SEVELAMER CARBONATE - RENVELA</u>						
N022318 002	5496545	Aug 11, 2013	DP	U-246	NDF	
	5667775	Sep 16, 2014		U-246		Aug 12, 2012
	6509013	Aug 11, 2013	DP			
	6858203	Aug 11, 2013	DP	U-246		
	7014846	Aug 11, 2013	DP	U-246		
	7459151	Aug 11, 2013		U-246		
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>						
N020926 001	5496545	Aug 11, 2013		U-246		
	5667775	Sep 16, 2014		U-246		
	6509013	Aug 11, 2013				
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>						
N021179 001	5496545	Aug 11, 2013		U-246		
	5667775	Sep 16, 2014		U-246		
	6509013	Aug 11, 2013				
	6733780	Oct 18, 2020	DP			
	7014846	Aug 11, 2013	DP	U-246		
	7459151	Aug 11, 2013		U-246		
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>						
N021179 002	5496545	Aug 11, 2013		U-246		
	5667775	Sep 16, 2014		U-246		
	6509013	Aug 11, 2013				
	6733780	Oct 18, 2020	DP			
	7014846	Aug 11, 2013	DP	U-246		
	7459151	Aug 11, 2013		U-246		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>SEVOFLURANE - ULTANE</u>							
N020478 001	5990176	Jan	27, 2017				
	5990176*PED	Jul	27, 2017				
	6074668	Jan	09, 2018				
	6074668*PED	Jul	09, 2018				
	6288127	Jan	27, 2017				
	6288127*PED	Jul	27, 2017				
	6444859	Jan	27, 2017				
	6444859*PED	Jul	27, 2017				
<u>SIBUTRAMINE HYDROCHLORIDE - MERIDIA</u>							
N020632 001	5436272	Jul	25, 2012		U-439		
	5436272*PED	Jan	25, 2013				
<u>SIBUTRAMINE HYDROCHLORIDE - MERIDIA</u>							
N020632 002	5436272	Jul	25, 2012		U-439		
	5436272*PED	Jan	25, 2013				
<u>SIBUTRAMINE HYDROCHLORIDE - MERIDIA</u>							
N020632 003	5436272	Jul	25, 2012		U-439		
	5436272*PED	Jan	25, 2013				
<u>SILDENAFIL CITRATE - REVATIO</u>							
N021845 001	5250534	Mar	27, 2012	DS DP		I-598	May 07, 2012
<u>SILDENAFIL CITRATE - REVATIO</u>							
N022473 001	5250534	Mar	27, 2012	DS DP		NDF	Nov 20, 2012
<u>SILDENAFIL CITRATE - VIAGRA</u>							
N020895 001	5250534	Mar	27, 2012		U-155		
	6469012	Oct	22, 2019				
<u>SILDENAFIL CITRATE - VIAGRA</u>							
N020895 002	5250534	Mar	27, 2012		U-155		
	6469012	Oct	22, 2019				
<u>SILDENAFIL CITRATE - VIAGRA</u>							
N020895 003	5250534	Mar	27, 2012		U-155		
	6469012	Oct	22, 2019				
<u>SILODOSIN - RAPAFLO</u>							
N022206 001	5387603	Dec	01, 2018	DS DP		NCE	Oct 08, 2013
	5403847	Nov	13, 2012	U-902			
	5780485	Nov	13, 2012	U-902			
	6015819	Nov	13, 2012	U-902			
<u>SILODOSIN - RAPAFLO</u>							
N022206 002	5387603	Dec	01, 2018	DS DP		NCE	Oct 08, 2013
	5403847	Nov	13, 2012	U-902			
	5780485	Nov	13, 2012	U-902			
	6015819	Nov	13, 2012	U-902			
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>							
N202343 001	6303661	Apr	24, 2017		U-1188		
	6699871	Jul	26, 2022	DS DP			
	6890898	Feb	02, 2019		U-1190		
	6890898	Feb	02, 2019		U-1189		
	6890898	Feb	02, 2019		U-1191		
	7078381	Feb	02, 2019		U-1188		
	7125873	Jul	26, 2022	DP	U-1190		
	7125873	Jul	26, 2022	DP	U-1189		
	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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>							
N202343 002	6303661	Apr	24, 2017		U-1188		
	6699871	Jul	26, 2022	DS DP			
	6890898	Feb	02, 2019		U-1190		
	6890898	Feb	02, 2019		U-1189		
	6890898	Feb	02, 2019		U-1191		
	7078381	Feb	02, 2019		U-1188		
	7125873	Jul	26, 2022	DP	U-1193		
	7125873	Jul	26, 2022	DP	U-1190		
	7125873	Jul	26, 2022	DP	U-1192		
	7125873	Jul	26, 2022	DP	U-1189		
	7326708	Apr	11, 2026	DS DP	U-1188		
	7459428	Feb	02, 2019		U-1189		
<u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u>							
N202343 003	6303661	Apr	24, 2017		U-1188		
	6699871	Jul	26, 2022	DS DP			
	6890898	Feb	02, 2019		U-1190		
	6890898	Feb	02, 2019		U-1191		
	6890898	Feb	02, 2019		U-1189		
	7078381	Feb	02, 2019		U-1188		
	7125873	Jul	26, 2022	DP	U-1192		
	7125873	Jul	26, 2022	DP	U-1189		
	7125873	Jul	26, 2022	DP	U-1193		
	7125873	Jul	26, 2022	DP	U-1190		
	7326708	Apr	11, 2026	DS DP	U-1188		
	7459428	Feb	02, 2019		U-1189		
<u>SINCALIDE - KINEVAC</u>							
N017697 001	6803046	Aug	16, 2022	DP			
<u>SINECATECHINS - VEREGEN</u>							
N021902 001	5795911	Oct	31, 2020		U-172		
	5968973	Apr	10, 2017		U-172		
	7858662	Oct	02, 2026	DP	U-172		
<u>SIROLIMUS - RAPAMUNE</u>							
N021083 001	5100899	Jul	07, 2013		U-290		
	5100899*PED	Jan	07, 2014				
	5403833	Apr	04, 2012		U-293		
	5403833*PED	Oct	04, 2012				
	5536729	Sep	30, 2013	DP			
	5536729*PED	Mar	30, 2014				
<u>SIROLIMUS - RAPAMUNE</u>							
N021110 001	5100899	Jul	07, 2013		U-290		
	5100899*PED	Jan	07, 2014				
	5145684*PED	Jul	25, 2011				
	5403833	Apr	04, 2012		U-293		
	5403833*PED	Oct	04, 2012				
	5989591	Mar	11, 2018	DP			
	5989591*PED	Sep	11, 2018				
<u>SIROLIMUS - RAPAMUNE</u>							
N021110 002	5100899	Jul	07, 2013		U-290		
	5100899*PED	Jan	07, 2014				
	5145684*PED	Jul	25, 2011				
	5403833	Apr	04, 2012		U-293		
	5403833*PED	Oct	04, 2012				
	5989591	Mar	11, 2018	DP			
	5989591*PED	Sep	11, 2018				
<u>SIROLIMUS - RAPAMUNE</u>							
N021110 003	5100899	Jul	07, 2013		U-290		
	5100899*PED	Jan	07, 2014				
	5145684*PED	Jul	25, 2011				
	5403833	Apr	04, 2012		U-293		
	5403833*PED	Oct	04, 2012				
	5989591	Mar	11, 2018	DP			
	5989591*PED	Sep	11, 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
SIROLIMUS - RAPAMUNE							
N021110 004	5100899	Jul	07, 2013		U-290		
	5100899*PED	Jan	07, 2014				
	5145684*PED	Jul	25, 2011				
	5403833	Apr	04, 2012		U-293		
	5403833*PED	Oct	04, 2012				
	5989591	Mar	11, 2018	DP			
	5989591*PED	Sep	11, 2018				
SITAGLIPTIN PHOSPHATE - JANUVIA							
N021995 001	6303661	Apr	24, 2017		U-774		
	6699871	Jul	26, 2022	DS DP	U-774	NCE	Oct 16, 2011
	6890898	Feb	02, 2019		U-775		
	6890898	Feb	02, 2019		U-1039		
	6890898	Feb	02, 2019		U-1036		
	7078381	Feb	02, 2019		U-775		
	7078381	Feb	02, 2019		U-1036		
	7078381	Feb	02, 2019		U-1037		
	7078381	Feb	02, 2019		U-1038		
	7125873	Jul	26, 2022		U-775		
	7125873	Jul	26, 2022		U-1036		
	7125873	Jul	26, 2022		U-1037		
	7125873	Jul	26, 2022		U-1038		
	7326708	Apr	11, 2026	DS DP	U-802		
SITAGLIPTIN PHOSPHATE - JANUVIA							
N021995 002	6303661	Apr	24, 2017		U-774		
	6699871	Jul	26, 2022	DS DP	U-774	NCE	Oct 16, 2011
	6890898	Feb	02, 2019		U-1039		
	6890898	Feb	02, 2019		U-1036		
	6890898	Feb	02, 2019		U-775		
	7078381	Feb	02, 2019		U-775		
	7078381	Feb	02, 2019		U-1038		
	7078381	Feb	02, 2019		U-1037		
	7078381	Feb	02, 2019		U-1036		
	7125873	Jul	26, 2022		U-1038		
	7125873	Jul	26, 2022		U-1036		
	7125873	Jul	26, 2022		U-1037		
	7125873	Jul	26, 2022		U-775		
	7326708	Apr	11, 2026	DS DP	U-802		
SITAGLIPTIN PHOSPHATE - JANUVIA							
N021995 003	6303661	Apr	24, 2017		U-774		
	6699871	Jul	26, 2022	DS DP	U-774	NCE	Oct 16, 2011
	6890898	Feb	02, 2019		U-1039		
	6890898	Feb	02, 2019		U-1036		
	6890898	Feb	02, 2019		U-775		
	7078381	Feb	02, 2019		U-1037		
	7078381	Feb	02, 2019		U-1036		
	7078381	Feb	02, 2019		U-1038		
	7078381	Feb	02, 2019		U-775		
	7125873	Jul	26, 2022		U-1037		
	7125873	Jul	26, 2022		U-1038		
	7125873	Jul	26, 2022		U-1036		
	7125873	Jul	26, 2022		U-775		
	7326708	Apr	11, 2026	DS DP	U-802		
SODIUM BENZOATE; SODIUM PHENYLACETATE - AMMONUL							
N020645 001						ODE	Feb 17, 2012
SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOTE							
N201444 001						ODE	Jan 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
<u>SODIUM OXYBATE - XYREM</u>							
N021196 001	6780889	Jul	04, 2020	DP		ODE	Nov 18, 2012
	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-1102		
	7851506	Dec	22, 2019		U-1101		
	7895059	Dec	17, 2022		U-1110		
<u>SODIUM PHOSPHATE, DIBASIC ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - VISICOL</u>							
N021097 001	5616346	May	18, 2013		U-359		
<u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u>							
N021892 001	5616346	May	18, 2013	DP	U-715		
	7687075	Jun	22, 2028	DS	DP		
<u>SOLIFENACIN SUCCINATE - VESICARE</u>							
N021518 001	6017927	Nov	19, 2018	DS	DP		
<u>SOLIFENACIN SUCCINATE - VESICARE</u>							
N021518 002	6017927	Nov	19, 2018	DS	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 001	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
	6152897	Nov	20, 2018	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 002	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
	6152897	Nov	20, 2018	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 003	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
	6152897	Nov	20, 2018	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 005	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
	6152897	Nov	20, 2018	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 008	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
	6152897	Nov	20, 2018	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 009	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
	6152897	Nov	20, 2018	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 010	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	DP			
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>							
N020280 011	5435076	Apr	16, 2013	DP			
	5501673	Apr	16, 2013	DP			
	5716338	Feb	10, 2015	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
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE						
N020280 012	5435076	Apr 16, 2013	DP			
	5501673	Apr 16, 2013	DP			
	5716338	Feb 10, 2015	DP			
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE						
N020280 013	5435076	Apr 16, 2013	DP			
	5501673	Apr 16, 2013	DP			
	5716338	Feb 10, 2015	DP			
SOMATROPIN RECOMBINANT - HUMATROPE						
N019640 001				I-585	Mar 12, 2012	
				ODE	Nov 01, 2013	
SOMATROPIN RECOMBINANT - HUMATROPE						
N019640 004				I-585	Mar 12, 2012	
				ODE	Nov 01, 2013	
SOMATROPIN RECOMBINANT - HUMATROPE						
N019640 005				I-585	Mar 12, 2012	
				ODE	Nov 01, 2013	
SOMATROPIN RECOMBINANT - HUMATROPE						
N019640 006				I-585	Mar 12, 2012	
				ODE	Nov 01, 2013	
SOMATROPIN RECOMBINANT - HUMATROPE						
N019640 007				I-585	Mar 12, 2012	
				ODE	Nov 01, 2013	
SOMATROPIN RECOMBINANT - NORDITROPIN						
N021148 001	5849700	Dec 15, 2015	U-340		I-572	Oct 31, 2011
	5849704	Dec 15, 2015			ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN						
N021148 002	5849700	Dec 15, 2015	U-340		I-572	Oct 31, 2011
	5849704	Dec 15, 2015			ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN						
N021148 003	5633352	May 27, 2014			I-572	Oct 31, 2011
	5849700	Dec 15, 2015	U-340		ODE	May 31, 2014
	5849704	Dec 15, 2015				
SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO						
N021148 008	5849700	Dec 15, 2015	U-1041		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-1041		ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO						
N021148 009	5849700	Dec 15, 2015	U-1041		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-1041		ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO						
N021148 010	5849700	Dec 15, 2015	U-1041		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-1041		ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX						
N021148 004	5849700	Dec 15, 2015	U-340		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-340		ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX						
N021148 005	5849700	Dec 15, 2015	U-340		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-340		ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX						
N021148 006	5849700	Dec 15, 2015	U-340		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-340		ODE	May 31, 2014

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX						
N021148 007	5849700	Dec 15, 2015		U-340	I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP	U-340	ODE	May 31, 2014
SOMATROPIN RECOMBINANT - NUTROPIN AQ						
N020522 001	5763394	Jun 09, 2015		DP		
SOMATROPIN RECOMBINANT - NUTROPIN AQ PEN						
N020522 002	5763394	Jun 09, 2015		DP		
SOMATROPIN RECOMBINANT - NUTROPIN DEPOT						
N021075 001	5654010	Aug 05, 2014				
	5656297	Jul 25, 2014				
	5912015	Mar 12, 2012				
	6051259	Dec 02, 2012		U-340		
SOMATROPIN RECOMBINANT - NUTROPIN DEPOT						
N021075 002	5654010	Aug 05, 2014				
	5656297	Jul 25, 2014				
	5912015	Mar 12, 2012				
	6051259	Dec 02, 2012		U-340		
SOMATROPIN RECOMBINANT - NUTROPIN DEPOT						
N021075 003	5654010	Aug 05, 2014				
	5656297	Jul 25, 2014				
	5912015	Mar 12, 2012				
	6051259	Dec 02, 2012		U-340		
SOMATROPIN RECOMBINANT - SAIZEN						
N019764 002	5898030	Apr 27, 2016		DP		
SOMATROPIN RECOMBINANT - SAIZEN						
N019764 003	5898030	Apr 27, 2016		DP		
SOMATROPIN RECOMBINANT - SEROSTIM						
N020604 001	5898030	Apr 27, 2016		DP		
SOMATROPIN RECOMBINANT - SEROSTIM						
N020604 002	5898030	Apr 27, 2016		DP		
SOMATROPIN RECOMBINANT - SEROSTIM						
N020604 003	5898030	Apr 27, 2016		DP		
SOMATROPIN RECOMBINANT - SEROSTIM						
N020604 004	5898030	Apr 27, 2016		DP		
SOMATROPIN RECOMBINANT - ZORBTIVE						
N021597 004	5288703	Oct 07, 2011		U-898		
	5898030	Apr 27, 2016	DP			
SORAFENIB TOSYLATE - NEXAVAR						
N021923 001	7235576	Jan 12, 2020	DS	DP	ODE	Nov 16, 2014
	7351834	Jan 12, 2020	DS		ODE	Dec 20, 2012
	7897623	Jan 12, 2020		DP		
SOTALOL HYDROCHLORIDE - SOTALOL HYDROCHLORIDE						
N022306 001					ODE	Jul 02, 2016
SPINOSAD - NATROBA						
N022408 001	5496931	Mar 05, 2013	DS	U-1105		
	6063771	Jun 22, 2019	DP	U-1105	NCE	Jan 18, 2016
	6342482	Jun 22, 2019	DP	U-1105		
	7030095	Jul 02, 2021	DP	U-1105		
STAVUDINE - ZERIT XR						
N021453 001	7135465	Feb 18, 2023	DP	U-167		
	7135465*PED	Aug 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
STAVUDINE - ZERIT XR							
N021453 002	7135465	Feb	18, 2023	DP	U-167		
	7135465*PED	Aug	18, 2023				
STAVUDINE - ZERIT XR							
N021453 003	7135465	Feb	18, 2023	DP	U-167		
	7135465*PED	Aug	18, 2023				
STAVUDINE - ZERIT XR							
N021453 004	7135465	Feb	18, 2023	DP	U-167		
	7135465*PED	Aug	18, 2023				
SUMATRIPTAN - IMITREX							
N020626 001	5307953	Dec	02, 2012				
	5307953*PED	Jun	02, 2013				
	5554639	Sep	10, 2013	U-232			
	5554639*PED	Mar	10, 2014				
	5705520	Dec	10, 2011	U-232			
	5705520*PED	Jun	10, 2012				
SUMATRIPTAN - IMITREX							
N020626 002	5307953	Dec	02, 2012				
	5307953*PED	Jun	02, 2013				
	5554639	Sep	10, 2013	U-232			
	5554639*PED	Mar	10, 2014				
	5705520	Dec	10, 2011	U-232			
	5705520*PED	Jun	10, 2012				
SUMATRIPTAN - IMITREX							
N020626 003	5307953	Dec	02, 2012				
	5307953*PED	Jun	02, 2013				
	5554639	Sep	10, 2013	U-232			
	5554639*PED	Mar	10, 2014				
	5705520	Dec	10, 2011	U-232			
	5705520*PED	Jun	10, 2012				
SUMATRIPTAN SUCCINATE - ALSUMA							
N022377 001	7811254	Aug	26, 2027	DP	U-1083		
SUMATRIPTAN SUCCINATE - SUMAVEL DOSEPRO							
N022239 001	5891086	Jul	27, 2014	DP			
	5957886	Mar	08, 2016	DP			
	6135979	Mar	21, 2017	DP			
	7776007	Nov	28, 2026	DP			
	7901385	Jul	31, 2026	DP			
SUNITINIB MALATE - SUTENT							
N021938 001	6573293	Feb	15, 2021	DS	DP	U-1154	I-639 May 20, 2014
	7125905	Feb	15, 2021	DS	DP		
	7211600	Dec	22, 2020			U-883	
SUNITINIB MALATE - SUTENT							
N021938 002	6573293	Feb	15, 2021	DS	DP	U-1154	I-639 May 20, 2014
	7125905	Feb	15, 2021	DS	DP		
	7211600	Dec	22, 2020			U-883	
SUNITINIB MALATE - SUTENT							
N021938 003	6573293	Feb	15, 2021	DS	DP	U-1154	I-639 May 20, 2014
	7125905	Feb	15, 2021	DS	DP		
	7211600	Dec	22, 2020			U-883	
SUNITINIB MALATE - SUTENT							
N021938 004	6573293	Feb	15, 2021	DS	DP	U-1154	I-639 May 20, 2014
	7125905	Feb	15, 2021	DS	DP		
	7211600	Dec	22, 2020			U-883	
TACROLIMUS - PROGRAF							
N050708 001						ODE	Mar 29, 2013
TACROLIMUS - PROGRAF							
N050708 002						ODE	Mar 29, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TACROLIMUS - PROGRAF</u>						
	N050708 003				ODE	Mar 29, 2013
<u>TACROLIMUS - PROGRAF</u>						
	N050709 001				ODE	Mar 29, 2013
<u>TACROLIMUS - PROTOPIC</u>						
N050777 001	5385907 5665727	Jan 31, 2012 Sep 09, 2014	DP U-919			
<u>TACROLIMUS - PROTOPIC</u>						
N050777 002	5385907 5665727	Jan 31, 2012 Sep 09, 2014	DP U-919			
<u>TADALAFIL - ADCIRCA</u>						
N022332 001	5859006 6821975 7182958	Nov 21, 2017 Nov 19, 2020 Apr 26, 2020	DS DP U-975 DS DP DP		NP ODE	May 22, 2012 May 22, 2016
<u>TADALAFIL - CIALIS</u>						
N021368 001	5859006 6140329 6140329 6821975 6821975 6821975 6943166 6943166 6943166 7182958 7182958	Nov 21, 2017 Jul 11, 2016 Jul 11, 2016 Nov 19, 2020 Nov 19, 2020 Nov 19, 2020 Apr 26, 2020 Apr 26, 2020 Apr 26, 2020 Apr 26, 2020	DS DP DP U-155 DP U-1184 DS DP U-533 DS DP U-614 DS DP U-1184 U-155 U-1184 U-614 DP U-1184 DP U-155		I-642 I-641	Oct 07, 2014 Oct 07, 2014
<u>TADALAFIL - CIALIS</u>						
N021368 002	5859006 6140329 6821975 6821975 6943166 6943166 7182958	Nov 21, 2017 Jul 11, 2016 Nov 19, 2020 Nov 19, 2020 Apr 26, 2020 Apr 26, 2020 Apr 26, 2020	DS DP DP U-155 DS DP U-614 DS DP U-533 U-155 U-614 DP U-155		I-642 I-641	Oct 07, 2014 Oct 07, 2014
<u>TADALAFIL - CIALIS</u>						
N021368 003	5859006 6140329 6821975 6821975 6943166 7182958	Nov 21, 2017 Jul 11, 2016 Nov 19, 2020 Nov 19, 2020 Apr 26, 2020 Apr 26, 2020	DS DP DP U-155 DS DP U-533 DS DP U-614 U-155 DP U-155		I-642 I-641	Oct 07, 2014 Oct 07, 2014
<u>TADALAFIL - CIALIS</u>						
N021368 004	5859006 6140329 6821975 6821975 6943166 7182958	Nov 21, 2017 Jul 11, 2016 Nov 19, 2020 Nov 19, 2020 Apr 26, 2020 Apr 26, 2020	DS DP DP U-155 DS DP U-614 DS DP U-533 U-155 DP U-155		I-642 I-641	Oct 07, 2014 Oct 07, 2014
<u>TAMOXIFEN CITRATE - SOLTAMOX</u>						
N021807 001	6127425	Jun 26, 2018	DP			
<u>TAMSULOSIN HYDROCHLORIDE - FLOMAX</u>						
N020579 001					M-54 PED	Dec 22, 2012 Jun 22, 2013
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>						
N022304 001	6071970 7994364 RE39593	Jun 06, 2017 Jun 27, 2025 Aug 05, 2022	U-931 DS DP U-931 DS DP U-931		NCE	Nov 20, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>							
N022304 002	6071970	Jun	06, 2017		U-931		
	7994364	Jun	27, 2025	DS DP	U-931		
	RE39593	Aug	05, 2022	DS DP	U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA</u>							
N022304 003	6071970	Jun	06, 2017		U-931		
	7994364	Jun	27, 2025	DS DP	U-931		
	RE39593	Aug	05, 2022	DS DP	U-931		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>							
N200533 001	6071970	Jun	06, 2017		U-1178		
	7994364	Jun	27, 2025	DS DP	U-1178		
	8075872	Nov	20, 2023	DP			
	RE39593	Aug	05, 2022	DS DP	U-1178		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>							
N200533 002	6071970	Jun	06, 2017		U-1178		
	7994364	Jun	27, 2025	DS DP	U-1178		
	8075872	Nov	20, 2023	DP			
	RE39593	Aug	05, 2022	DS DP	U-1178		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>							
N200533 003	6071970	Jun	06, 2017		U-1178		
	7994364	Jun	27, 2025	DS DP	U-1178		
	8075872	Nov	20, 2023	DP			
	RE39593	Aug	05, 2022	DS DP	U-1178		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>							
N200533 004	6071970	Jun	06, 2017		U-1178		
	7994364	Jun	27, 2025	DS DP	U-1178		
	8075872	Nov	20, 2023	DP			
	RE39593	Aug	05, 2022	DS DP	U-1178		
<u>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</u>							
N200533 005	6071970	Jun	06, 2017		U-1178		
	7994364	Jun	27, 2025	DS DP	U-1178		
	8075872	Nov	20, 2023	DP			
	RE39593	Aug	05, 2022	DS DP	U-1178		
<u>TAZAROTENE - TAZORAC</u>							
N020600 001	5914334	Jun	07, 2014		U-517		
	6258830	Jun	07, 2014		U-517		
<u>TAZAROTENE - TAZORAC</u>							
N020600 002	5914334	Jun	07, 2014		U-517		
	6258830	Jun	07, 2014		U-517		
<u>TECHNETIUM TC-99M APCITIDE - ACUTECT</u>							
N020887 001	5443815	Aug	22, 2012				
	5508020	Apr	16, 2013				
	5645815	Jul	08, 2014				
<u>TECHNETIUM TC-99M BICISATE KIT - NEUROLITE</u>							
N020256 001	5431900	Jul	11, 2012		U-336		
<u>TECHNETIUM TC-99M MERTIATIDE KIT - TECHNESCAN MAG3</u>							
N019882 001	5573748	Nov	12, 2013	DP			
<u>TECHNETIUM TC-99M SESTAMIBI KIT - CARDIOLITE</u>							
N019785 001						PED	Oct 30, 2011
<u>TECHNETIUM TC-99M TEBOROXIME KIT - CARDIOTEC</u>							
N019928 001	6056941	Jul	28, 2019	DP			
<u>TEGASEROD MALEATE - ZELNORM</u>							
N021200 001	5510353	Apr	26, 2013		U-466		
<u>TEGASEROD MALEATE - ZELNORM</u>							
N021200 002	5510353	Apr	26, 2013		U-466		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TELAPREVIR - INCIVEK</u>								
N0201917 001	7820671	Feb	25, 2025	DS	DP		NCE	May 23, 2016
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>								
N022110 001	6635618	Sep	22, 2021	DS	DP	U-728	NCE	Sep 11, 2014
	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		
<u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u>								
N022110 002	6635618	Sep	22, 2021	DS	DP	U-728	NCE	Sep 11, 2014
	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		
<u>TELBIVUDINE - TYZEKA</u>								
N022011 001	6395716	Aug	10, 2019			U-782	NCE	Oct 25, 2011
	6444652	Aug	10, 2019			U-782		
	6566344	Aug	10, 2019			U-782		
	6569837	Oct	25, 2020			U-999		
	6569837	Oct	25, 2020			U-782		
	7589079	Sep	11, 2023	DS	DP	U-999		
	7795238	Aug	10, 2019			U-999		
	7858594	Sep	11, 2023	DS	DP	U-999		
<u>TELBIVUDINE - TYZEKA</u>								
N022154 001	6395716	Aug	10, 2019			U-999	NCE	Oct 25, 2011
	6444652	Aug	10, 2019			U-999		
	6566344	Aug	10, 2019			U-999		
	6569837	Oct	25, 2020			U-999		
	7795238	Aug	10, 2019			U-999		
<u>TELITHROMYCIN - KETEK</u>								
N021144 001	5635485	Apr	01, 2018	DS	DP	U-578		
	D459798	Sep	24, 2015			DP		
<u>TELITHROMYCIN - KETEK</u>								
N021144 002	5635485	Apr	01, 2018	DS	DP	U-578		
	D459798	Sep	24, 2015			DP		
<u>TELMISARTAN - MICARDIS</u>								
N020850 001	5591762	Jan	07, 2014			U-3		
	6358986	Jan	10, 2020					
<u>TELMISARTAN - MICARDIS</u>								
N020850 002	5591762	Jan	07, 2014	DS	DP	U-3	I-612	Oct 16, 2012
	6358986	Jan	10, 2020					
	7998953	Jun	06, 2020			U-1177		
	8003679	Oct	06, 2022			U-1176		
<u>TELMISARTAN - MICARDIS</u>								
N020850 003	5591762	Jan	07, 2014			U-3		
	6358986	Jan	10, 2020					
<u>TEMOZOLOMIDE - TEMODAR</u>								
N021029 001	5260291	Aug	11, 2013	DS	DP	U-619	ODE	Mar 15, 2012
	5260291*PED	Feb	11, 2014					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TEMOZOLOMIDE - TEMODAR</u>										
N021029 002	5260291	Aug	11, 2013	DS	DP	U-619			ODE	Mar 15, 2012
	5260291*PED	Feb	11, 2014							
<u>TEMOZOLOMIDE - TEMODAR</u>										
N021029 003	5260291	Aug	11, 2013	DS	DP	U-619			ODE	Mar 15, 2012
	5260291*PED	Feb	11, 2014							
<u>TEMOZOLOMIDE - TEMODAR</u>										
N021029 004	5260291	Aug	11, 2013	DS	DP	U-619			ODE	Mar 15, 2012
	5260291*PED	Feb	11, 2014							
<u>TEMOZOLOMIDE - TEMODAR</u>										
N021029 005	5260291	Aug	11, 2013	DS	DP	U-619			ODE	Mar 15, 2012
	5260291*PED	Feb	11, 2014							
<u>TEMOZOLOMIDE - TEMODAR</u>										
N021029 006	5260291	Aug	11, 2013	DS	DP	U-619			ODE	Mar 15, 2012
	5260291*PED	Feb	11, 2014							
<u>TEMOZOLOMIDE - TEMODAR</u>										
N022277 001	5260291	Aug	11, 2013	DS	DP	U-619				
	5260291*PED	Feb	11, 2014							
	6987108	Sep	08, 2023		DP					
	7786118	Feb	21, 2023		DP					
<u>TEMSIROLIMUS - TORISEL</u>										
N022088 001	5362718	Apr	18, 2014	DS	DP				M-92	Jul 09, 2013
	8026276	Jan	20, 2026		DP				M-91	Apr 26, 2013
									NCE	May 30, 2012
									ODE	May 30, 2014
<u>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</u>										
N021356 001	5922695	Jul	25, 2017	DS		U-999			I-569	Aug 11, 2011
	5922695	Jul	25, 2017	DS		U-248			M-95	Oct 01, 2013
	5922695	Jul	25, 2017	DS		U-250			NPP	Mar 24, 2013
	5922695	Jul	25, 2017	DS		U-256			ODE	Mar 24, 2017
	5922695*PED	Jan	25, 2018						PED	Feb 11, 2012
	5935946	Jul	25, 2017	DS	DP	U-250			PED	Apr 01, 2014
	5935946	Jul	25, 2017	DS	DP	U-999			PED	Sep 24, 2013
	5935946	Jul	25, 2017	DS	DP	U-256			PED	Sep 24, 2017
	5935946	Jul	25, 2017	DS	DP	U-248				
	5935946*PED	Jan	25, 2018							
	5977089	Jul	25, 2017	DS	DP	U-250				
	5977089	Jul	25, 2017	DS	DP	U-248				
	5977089	Jul	25, 2017	DS	DP	U-256				
	5977089	Jul	25, 2017	DS	DP	U-999				
	5977089*PED	Jan	25, 2018							
	6043230	Jul	25, 2017			U-250				
	6043230	Jul	25, 2017			U-256				
	6043230	Jul	25, 2017			U-999				
	6043230	Jul	25, 2017			U-248				
	6043230*PED	Jan	25, 2018							
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>										
N019057 001	5294615	Apr	29, 2013			U-3				
	5294615	Apr	29, 2013			U-165				
	5412095	Apr	29, 2013							
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>										
N019057 002	5294615	Apr	29, 2013			U-165				
	5294615	Apr	29, 2013			U-3				
	5412095	Apr	29, 2013							
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>										
N019057 003	5294615	Apr	29, 2013			U-165				
	5294615	Apr	29, 2013			U-3				
	5412095	Apr	29, 2013							

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>						
N019057 004	5294615	Apr 29, 2013		U-3		
	5294615	Apr 29, 2013		U-165		
	5412095	Apr 29, 2013				
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>						
N020347 001	5294615	Apr 29, 2013		U-165		
	5294615	Apr 29, 2013		U-3		
	5412095	Apr 29, 2013				
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>						
N020347 002	5294615	Apr 29, 2013		U-165		
	5294615	Apr 29, 2013		U-3		
	5412095	Apr 29, 2013				
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>						
N020347 003	5294615	Apr 29, 2013		U-3		
	5294615	Apr 29, 2013		U-165		
	5412095	Apr 29, 2013				
<u>TERAZOSIN HYDROCHLORIDE - HYTRIN</u>						
N020347 004	5294615	Apr 29, 2013		U-3		
	5294615	Apr 29, 2013		U-165		
	5412095	Apr 29, 2013				
<u>TERBINAFINE - LAMISIL</u>						
N020846 001	5681849	Oct 28, 2014	DP			
	5681849*PED	Apr 28, 2015				
	5856355	May 18, 2012	DP	U-502		
	5856355	May 18, 2012	DP	U-504		
	5856355	May 18, 2012	DP	U-540		
	5856355*PED	Nov 18, 2012				
	6005001	May 18, 2012	DP	U-502		
	6005001	May 18, 2012	DP	U-540		
	6005001	May 18, 2012	DP	U-504		
	6005001*PED	Nov 18, 2012				
<u>TERBINAFINE - LAMISIL AT</u>						
N021958 001	5681849	Oct 28, 2014	DP			
	5681849*PED	Apr 28, 2015				
	5856355	May 18, 2012	DP	U-540		
	5856355	May 18, 2012	DP	U-504		
	5856355*PED	Nov 18, 2012				
<u>TERBINAFINE HYDROCHLORIDE - LAMISIL AT</u>						
N021124 001	5681849	Oct 28, 2014				
	5681849*PED	Apr 28, 2015				
<u>TERBINAFINE HYDROCHLORIDE - LAMISIL AT</u>						
N021124 002	5681849	Oct 28, 2014				
	5681849*PED	Apr 28, 2015				
<u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u>						
N021318 001	6770623	Dec 08, 2018	DP	U-597	I-602	Jul 22, 2012
	6977077	Aug 19, 2019		U-597		
	7144861	Dec 08, 2018	DP			
	7163684	Aug 19, 2019		U-790		
	7351414	Aug 19, 2019		U-865		
	7550434	Dec 08, 2018	DP	U-982		
<u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u>						
N021318 002	6770623	Dec 08, 2018	DP	U-982	I-602	Jul 22, 2012
	6977077	Aug 19, 2019		U-994		
	6977077	Aug 19, 2019		U-982		
	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		
	7550434	Dec 08, 2018	DP	U-982		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE			PATENT DELIST REQUESTED		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TESAMORELIN ACETATE - EGRIFTA</u>								
N022505 001	5861379	May 26, 2015	DS	DP	U-1100		NCE	Nov 10, 2015
	6020311	May 26, 2015	DS	DP	U-1100			
	7144577	Jul 14, 2020			U-1100			
	7316997	Aug 14, 2023			U-1100			
<u>TESTOSTERONE - ANDRODERM</u>								
N020489 003							NS	Oct 20, 2014
<u>TESTOSTERONE - ANDRODERM</u>								
N020489 004							NS	Oct 20, 2014
<u>TESTOSTERONE - ANDROGEL</u>								
N021015 001	6503894	Aug 30, 2020			U-490			
	6503894*PED	Mar 01, 2021						
<u>TESTOSTERONE - ANDROGEL</u>								
N021015 002	6503894	Aug 30, 2020			U-490			
	6503894*PED	Mar 01, 2021						
<u>TESTOSTERONE - ANDROGEL</u>								
N021015 003	6503894	Aug 30, 2020			U-490			
	6503894*PED	Mar 01, 2021						
<u>TESTOSTERONE - ANDROGEL</u>								
N022309 001	6503894	Aug 30, 2020			U-1103		NP	Apr 29, 2014
<u>TESTOSTERONE - AXIRON</u>								
N022504 001	6299900	Feb 19, 2017	DP	U-1103			NP	Nov 23, 2013
	6818226	Feb 19, 2017	DP	U-1103				
	6923983	Feb 19, 2017	DP	U-1103				
	8071075	Feb 19, 2017	DP	U-1103				
<u>TESTOSTERONE - FORTESTA</u>								
N021463 001	6319913	Nov 09, 2018		U-490			NP	Dec 29, 2013
	6579865	Nov 09, 2018	DP					
<u>TESTOSTERONE - STRIANT</u>								
N021543 001	6248358	Aug 23, 2019		U-527				
<u>TESTOSTERONE - TESTIM</u>								
N021454 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				
<u>TESTOSTERONE - TESTODERM</u>								
N019762 001	5840327	Aug 15, 2016						
<u>TESTOSTERONE - TESTODERM</u>								
N019762 002	5840327	Aug 15, 2016						
<u>TESTOSTERONE - TESTODERM TTS</u>								
N020791 001	6348210	Nov 10, 2019		U-440				
<u>TETRABENAZINE - XENAZINE</u>								
N021894 001							NCE	Aug 15, 2013
							ODE	Aug 15, 2015
<u>TETRABENAZINE - XENAZINE</u>								
N021894 002							NCE	Aug 15, 2013
							ODE	Aug 15, 2015

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>THALIDOMIDE - THALOMID</u>						
N020785 001	5629327	May 13, 2014	U-731		ODE	May 25, 2013
	6045501	Aug 28, 2018	U-731			
	6045501	Aug 28, 2018	U-371			
	6235756	Mar 01, 2013	U-731			
	6315720	Oct 23, 2020	U-731			
	6315720	Oct 23, 2020	U-442			
	6561976	Aug 28, 2018	U-731			
	6561976	Aug 28, 2018	U-371			
	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-731			
	6908432	Aug 28, 2018	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-733			
	7141018	Oct 23, 2020	U-732			
	7230012	Dec 09, 2023	DP			
	7435745	Nov 03, 2017	U-899			
	7723361	Mar 01, 2013	U-1058			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-442			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7959566	Oct 23, 2020	U-1155			
<u>THALIDOMIDE - THALOMID</u>						
N020785 002	5629327	May 13, 2014	U-731		ODE	May 25, 2013
	6045501	Aug 28, 2018	U-371			
	6045501	Aug 28, 2018	U-731			
	6235756	Mar 01, 2013	U-731			
	6315720	Oct 23, 2020	U-731			
	6315720	Oct 23, 2020	U-442			
	6561976	Aug 28, 2018	U-731			
	6561976	Aug 28, 2018	U-371			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-733			
	6908432	Aug 28, 2018	U-371			
	6908432	Aug 28, 2018	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-371			
	7230012	Dec 09, 2023	DP			
	7435745	Nov 03, 2017	U-899			
	7723361	Mar 01, 2013	U-1058			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7874984	Aug 28, 2018	U-442			
	7959566	Oct 23, 2020	U-1155			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>THALIDOMIDE - THALOMID</u>						
N020785 003	5629327	May 13, 2014	U-731		ODE	May 25, 2013
	6045501	Aug 28, 2018	U-731			
	6045501	Aug 28, 2018	U-371			
	6235756	Mar 01, 2013	U-731			
	6315720	Oct 23, 2020	U-731			
	6315720	Oct 23, 2020	U-442			
	6561976	Aug 28, 2018	U-371			
	6561976	Aug 28, 2018	U-731			
	6561977	Oct 23, 2020	U-731			
	6561977	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	6869399	Oct 23, 2020	U-371			
	6908432	Aug 28, 2018	U-371			
	6908432	Aug 28, 2018	U-731			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023	DP			
	7435745	Nov 03, 2017	U-899			
	7723361	Mar 01, 2013	U-1058			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-442			
	7874984	Aug 28, 2018	U-733			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-1109			
	7959566	Oct 23, 2020	U-1155			
<u>THALIDOMIDE - THALOMID</u>						
N020785 004	5629327	May 13, 2014	U-731		ODE	May 23, 2013
	6045501	Aug 28, 2018	U-731			
	6235756	Mar 01, 2013	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			
	7723361	Mar 01, 2013	U-1058			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-442			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7959566	Oct 23, 2020	U-1155			
<u>THYROTROPIN ALFA - THYROGEN</u>						
N020898 001	5840566	Nov 24, 2015			ODE	
	6365127	Nov 24, 2015	DS DP U-556			Dec 14, 2014
<u>TIAGABINE HYDROCHLORIDE - GABITRIL</u>						
N020646 001	5010090	Sep 30, 2011				
	5354760	Mar 24, 2012				
	5866590	Apr 29, 2016				
	5958951	Jun 10, 2017				
<u>TIAGABINE HYDROCHLORIDE - GABITRIL</u>						
N020646 002	5010090	Sep 30, 2011				
	5354760	Mar 24, 2012				
	5866590	Apr 29, 2016				
	5958951	Jun 10, 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
<u>TIAGABINE HYDROCHLORIDE - GABITRIL</u>							
N020646 003	5010090	Sep	30, 2011				
	5354760	Mar	24, 2012				
	5866590	Apr	29, 2016				
	5958951	Jun	10, 2017				
<u>TIAGABINE HYDROCHLORIDE - GABITRIL</u>							
N020646 004	5010090	Sep	30, 2011				
	5354760	Mar	24, 2012				
	5866590	Apr	29, 2016				
	5958951	Jun	10, 2017				
<u>TIAGABINE HYDROCHLORIDE - GABITRIL</u>							
N020646 005	5010090	Sep	30, 2011				
	5354760	Mar	24, 2012				
	5866590	Apr	29, 2016				
	5958951	Jun	10, 2017				
<u>TICAGRELOR - BRILINTA</u>							
N022433 001	6251910	Jul	15, 2018	DS			
	6525060	Dec	02, 2019	DS	DP	U-1171	
	7250419	Dec	02, 2019	DS	DP	U-1171	
	7265124	Jul	09, 2021	DS	DP	U-1171	
<u>TIGECYCLINE - TYGACIL</u>							
N021821 001	7879828	Feb	05, 2029				
	RE40086	Jun	25, 2013				
	RE40183	Apr	09, 2016	DS	DP	U-282	
I-588	Mar	20, 2012					
I-587	Mar	20, 2012					
I-586	Mar	20, 2012					
<u>TIMOLOL MALEATE - ISTALOL</u>							
N021516 001	6335335	Nov	02, 2018				
<u>TIMOLOL MALEATE - TIMOLOL MALEATE</u>							
N020963 001	6174524	Mar	26, 2019				
	6174524*PED	Sep	26, 2019				
<u>TIMOLOL MALEATE - TIMOLOL MALEATE</u>							
N020963 002	6174524	Mar	26, 2019				
	6174524*PED	Sep	26, 2019				
<u>TIOTROPIUM BROMIDE MONOHYDRATE - SPIRIVA</u>							
N021395 001	5478578	Dec	26, 2012				
	6777423	Sep	24, 2021	DS	DP		
	6908928	Sep	24, 2021	DS	DP	U-762	
	6908928	Sep	24, 2021	DS	DP	U-566	
	7070800	Jan	22, 2022				
	7309707	Sep	24, 2021	DS	DP		
	7642268	Sep	24, 2021	DS	DP		
	7694676	Mar	12, 2027				
	8022082	Jan	19, 2026				
	RE38912	Oct	11, 2021				
	RE39820	Jan	30, 2018	DS	DP	U-1186	
M-89	Dec	17, 2012					
<u>TIPRANAVIR - APTIVUS</u>							
N021814 001	5852195	Jun	22, 2019	DS			
	5852195*PED	Dec	22, 2019				
	6147095	Oct	29, 2019				
	6147095*PED	Apr	29, 2020				
	6169181	May	06, 2014	DS			
	6169181*PED	Nov	06, 2014				
	6231887	Jul	27, 2018				
	6231887*PED	Jan	27, 2019				
PED	Dec	23, 2011					
<u>TIPRANAVIR - APTIVUS</u>							
N022292 001	5852195	Jun	22, 2019	DS			
	5852195*PED	Dec	22, 2019				
	6147095	Oct	29, 2019				
	6147095*PED	Apr	29, 2020				
	6169181	May	06, 2014	DS			
	6169181*PED	Nov	06, 2014				
PED	Dec	23, 2011					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N020912 001	5292756	May 14, 2012		U-230		
	5733919	Oct 23, 2016				
	5965581	Oct 23, 2016				
	5972967	Oct 23, 2016				
	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N020913 001	5292756	May 14, 2012		U-230		
	5733919	Oct 23, 2016				
	5965581	Oct 23, 2016				
	5972967	Oct 23, 2016				
	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N020913 002	5292756	May 14, 2012		U-230		
	5733919	Oct 23, 2016				
	5965581	Oct 23, 2016				
	5972967	Oct 23, 2016				
	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
<u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u>						
N020913 003	5292756	May 14, 2012		U-230		
	5733919	Oct 23, 2016				
	5965581	Oct 23, 2016				
	5972967	Oct 23, 2016				
	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
<u>TIZANIDINE HYDROCHLORIDE - ZANAFLEX</u>						
N021447 001	6455557	Nov 28, 2021				
<u>TIZANIDINE HYDROCHLORIDE - ZANAFLEX</u>						
N021447 002	6455557	Nov 28, 2021				
<u>TIZANIDINE HYDROCHLORIDE - ZANAFLEX</u>						
N021447 003	6455557	Nov 28, 2021				
<u>TOBRAMYCIN - TOBI</u>						
N050753 001	5508269	Oct 19, 2014		DP U-909		
<u>TOLCAPONE - TASMAR</u>						
N020697 001	5236952	Jan 29, 2012				
	5476875	Dec 19, 2012		U-219		
<u>TOLCAPONE - TASMAR</u>						
N020697 002	5236952	Jan 29, 2012				
	5476875	Dec 19, 2012		U-219		
<u>TOLTERODINE TARTRATE - DETROL</u>						
N020771 001	5382600	Mar 25, 2012				
	5382600*PED	Sep 25, 2012				
	5559269	Nov 05, 2013		U-318	Y	
	5559269*PED	May 05, 2014				
<u>TOLTERODINE TARTRATE - DETROL</u>						
N020771 002	5382600	Mar 25, 2012				
	5382600*PED	Sep 25, 2012				
	5559269	Nov 05, 2013		U-318	Y	
	5559269*PED	May 05, 2014				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
TOLTERODINE TARTRATE - DETROL LA							
N021228 001	5382600	Mar	25, 2012				
	5382600*PED	Sep	25, 2012				
	6630162	Nov	11, 2019	DP	U-544		
	6630162*PED	May	11, 2020				
	6770295	Aug	26, 2019	DP	U-544		
	6770295*PED	Feb	26, 2020				
	6911217	Aug	26, 2019	DP	U-544		
	6911217*PED	Feb	26, 2020	DP	U-544		
TOLTERODINE TARTRATE - DETROL LA							
N021228 002	5382600	Mar	25, 2012				
	5382600*PED	Sep	25, 2012				
	6630162	Nov	11, 2019	DP	U-544		
	6630162*PED	May	11, 2020				
	6770295	Aug	26, 2019	DP	U-544		
	6770295*PED	Feb	26, 2020				
	6911217	Aug	26, 2019	DP	U-544		
	6911217*PED	Feb	26, 2020	DP	U-544		
TOLVAPTAN - SAMSCA							
N022275 001	5753677	May	19, 2015	U-978		NCE	May 19, 2014
TOLVAPTAN - SAMSCA							
N022275 002	5753677	May	19, 2015	U-978		NCE	May 19, 2014
TOLVAPTAN - SAMSCA							
N022275 003	5753677	May	19, 2015	U-978		NCE	May 19, 2014
TOPIRAMATE - TOPAMAX							
N020505 001	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
TOPIRAMATE - TOPAMAX							
N020505 002	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
TOPIRAMATE - TOPAMAX							
N020505 003	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
TOPIRAMATE - TOPAMAX							
N020505 004	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TOPIRAMATE - TOPAMAX</u>							
N020505 005	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>							
N020505 006	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>							
N020844 001	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7125560	Mar	01, 2019	U-766			
	7125560*PED	Sep	01, 2019				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
<u>TOPIRAMATE - TOPAMAX SPRINKLE</u>							
N020844 002	5998380	Oct	13, 2015	U-598		M-54	Dec 22, 2012
	5998380*PED	Apr	13, 2016			PED	Jun 22, 2013
	6503884	Oct	13, 2015	U-598			
	6503884*PED	Apr	13, 2016				
	7018983	Oct	13, 2015	U-723			
	7018983*PED	Apr	13, 2016				
	7125560	Mar	01, 2019	U-766			
	7125560*PED	Sep	01, 2019				
	7498311	Oct	13, 2015	U-955			
	7498311*PED	Apr	13, 2016				
<u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u>							
N020671 001	5674872	Oct	07, 2014	U-910			
	5674872*PED	Apr	07, 2015				
<u>TRAMADOL HYDROCHLORIDE - CONZIP</u>							
N022370 001	7858118	Apr	11, 2022	DP	U-1104		
<u>TRAMADOL HYDROCHLORIDE - CONZIP</u>							
N022370 002	7858118	Apr	11, 2022	DP	U-1104		
<u>TRAMADOL HYDROCHLORIDE - CONZIP</u>							
N022370 003	7858118	Apr	11, 2022	DP	U-1104		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
TRAMADOL HYDROCHLORIDE - RYZOLT							
N021745 001	5591452	May	10, 2014	DP		NP	Dec 30, 2011
	6254887	May	10, 2014	DP			
	6607748	Jun	29, 2020	DP			
	7988998	Oct	27, 2023	DP			
TRAMADOL HYDROCHLORIDE - RYZOLT							
N021745 002	5591452	May	10, 2014	DP		NP	Dec 30, 2011
	6254887	May	10, 2014	DP			
	6607748	Jun	29, 2020	DP			
	7988998	Oct	27, 2023	DP			
TRAMADOL HYDROCHLORIDE - RYZOLT							
N021745 003	5591452	May	10, 2014	DP		NP	Dec 30, 2011
	6254887	May	10, 2014	DP			
	6607748	Jun	29, 2020	DP			
	7988998	Oct	27, 2023	DP			
TRAMADOL HYDROCHLORIDE - TRAMADOL HYDROCHLORIDE							
N021693 001	5464632	Mar	22, 2013	DP			
	6106861	Dec	05, 2017	DP			
TRAMADOL HYDROCHLORIDE - ULTRAM							
N020281 001	6339105	Oct	12, 2019	U-435			
	6339105*PED	Apr	12, 2020	U-435			
TRAMADOL HYDROCHLORIDE - ULTRAM							
N020281 002	6339105	Oct	12, 2019	U-435			
	6339105*PED	Apr	12, 2020	U-435			
TRANDOLAPRIL - MAVIK							
N020528 001	5744496	Apr	28, 2015	U-229			
TRANDOLAPRIL - MAVIK							
N020528 002	5744496	Apr	28, 2015	U-229			
TRANDOLAPRIL - MAVIK							
N020528 003	5744496	Apr	28, 2015	U-229			
TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE - TARKA							
N020591 001	5721244	Feb	24, 2015				
TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE - TARKA							
N020591 002	5721244	Feb	24, 2015				
TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE - TARKA							
N020591 003	5721244	Feb	24, 2015				
TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE - TARKA							
N020591 004	5721244	Feb	24, 2015				
TRANEXAMIC ACID - LYSTEDA							
N022430 001	7947739	Mar	04, 2025	DP		NDF	Nov 13, 2012
	8022106	Mar	04, 2025	U-1182			
TRAVOPROST - TRAVATAN							
N021257 001	5510383	Aug	03, 2013	DP	U-383		
	5631287	Dec	22, 2014	DP	U-382		
	5849792	Dec	22, 2014	DP	U-383		
	5889052	Dec	02, 2014	DP	U-383		
	6011062	Dec	22, 2014	DP	U-383		
TRAVOPROST - TRAVATAN Z							
N021994 001	5510383	Aug	03, 2013	DP	U-383		
	5889052	Dec	02, 2014	DP	U-383		
	6503497	May	06, 2012	DP			
	6849253	May	06, 2012	DP			
TRAZODONE HYDROCHLORIDE - OLEPTRO							
N022411 001	6607748	Jun	29, 2020	DP		NDF	Feb 02, 2013
	7829120	Mar	27, 2027	DP	U-796		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>							
N022411 002	6607748	Jun	29, 2020	DP		NDF	Feb 02, 2013
	7829120	Mar	27, 2027	DP	U-796		
<u>TREPROSTINIL SODIUM - REMODULIN</u>							
N021272 001	5153222	Oct	06, 2014		U-455		
	6765117	Oct	24, 2017	DS			
	7999007	Mar	29, 2029	DP	U-1174		
<u>TREPROSTINIL SODIUM - REMODULIN</u>							
N021272 002	5153222	Oct	06, 2014		U-455		
	6765117	Oct	24, 2017	DS			
	7999007	Mar	29, 2029	DP	U-1174		
<u>TREPROSTINIL SODIUM - REMODULIN</u>							
N021272 003	5153222	Oct	06, 2014		U-455		
	6765117	Oct	24, 2017	DS			
	7999007	Mar	29, 2029	DP	U-1174		
<u>TREPROSTINIL SODIUM - REMODULIN</u>							
N021272 004	5153222	Oct	06, 2014		U-455		
	6765117	Oct	24, 2017	DS			
	7999007	Mar	29, 2029	DP	U-1174		
<u>TREPROSTINIL SODIUM - TYVASO</u>							
N022387 001	5153222	Oct	16, 2014		U-1019	NDF	Jul 30, 2012
	6521212	Nov	13, 2018		U-1018	ODE	Jul 30, 2016
	6756033	Nov	13, 2018		U-1018		
	6765117	Oct	24, 2017	DS			
<u>TRETINOIN - ATRALIN</u>							
N022070 001	5670547	Sep	23, 2014	DP			
<u>TRETINOIN - RENOVA</u>							
N021108 001	6531141	Mar	07, 2020				
<u>TRETINOIN - RETIN-A MICRO</u>							
N020475 001	5955109	Sep	21, 2016	DP	U-134		
<u>TRETINOIN - RETIN-A MICRO</u>							
N020475 002	5955109	Sep	21, 2016		U-134		
<u>TRIAMCINOLONE ACETONIDE - NASACORT AQ</u>							
N020468 001	5976573	Jul	03, 2016	DP	U-295	NPP	Sep 19, 2011
	5976573	Jul	03, 2016	DP	U-896		
	6143329	Jul	03, 2016	DP	U-896		
	7977045	Jul	03, 2016	DP	U-896		
<u>TRIAMCINOLONE ACETONIDE - TRIESENCE</u>							
N022048 001	6395294	Jan	13, 2020	DP	U-846		
<u>TRIMETHOPRIM HYDROCHLORIDE - PRIMSOL</u>							
A074973 001	5763449	Aug	07, 2016				
	5962461	Aug	07, 2016				
<u>TRIMETREXATE GLUCURONATE - NEUTREXIN</u>							
N020326 001	6017922	May	18, 2018				
<u>TRIMETREXATE GLUCURONATE - NEUTREXIN</u>							
N020326 002	6017922	May	18, 2018				
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>							
N020715 001	5776885	Jul	07, 2015				
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>							
N021288 001	5776885	Jul	07, 2015	DP			
<u>TRIPTORELIN PAMOATE - TRELSTAR</u>							
N022437 001	5776885	Jul	07, 2015	DP		NP	Mar 10, 2013

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
TROGLITAZONE - PRELAY						
N020719 001	5602133	Sep 15, 2013	U-173			
	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
	6046202	Sep 15, 2013	U-317			
TROGLITAZONE - PRELAY						
N020719 002	5602133	Sep 15, 2013	U-173			
	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
	6046202	Sep 15, 2013	U-317			
TROGLITAZONE - PRELAY						
N020719 003	5602133	Sep 15, 2013	U-173			
	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
	6046202	Sep 15, 2013	U-317			
TROGLITAZONE - REZULIN						
N020720 001	5602133	Sep 15, 2013	U-173			
	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
	6046202	Sep 15, 2013	U-317			
TROGLITAZONE - REZULIN						
N020720 002	5602133	Sep 15, 2013	U-173			
	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
	6046202	Sep 15, 2013	U-317			
TROGLITAZONE - REZULIN						
N020720 003	5602133	Sep 15, 2013	U-173			
	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
	6046202	Sep 15, 2013	U-317			
TROSPiUM CHLORIDE - SANCTURA XR						
N022103 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		
TROVAFLOXACIN MESYLATE - TROVAN						
N020759 001	5164402	Dec 18, 2011	U-282			
	5763454	Jun 15, 2015	U-282			
	6187341	Jan 20, 2019				
TROVAFLOXACIN MESYLATE - TROVAN						
N020759 002	5164402	Dec 18, 2011	U-282			
	5763454	Jun 15, 2015	U-282			
	6187341	Jan 20, 2019				
TRYPAN BLUE - MEMBRANEBLUE						
N022278 001					ODE	Dec 16, 2011
ULIPRISTAL ACETATE - ELLA						
N022474 001					NCE	Aug 13, 2015
UNOPROSTONE ISOPROPYL - RESCULA						
N021214 001	5221763	Jul 15, 2012	DS			
	6458836	Jul 09, 2021		U-333		
UROFOLLITROPIN - FERTINEX						
N019415 004	5767067	Jun 16, 2015				
UROFOLLITROPIN - FERTINEX						
N019415 005	5767067	Jun 16, 2015				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>VALACYCLOVIR HYDROCHLORIDE - VALTREX</u>							
N020487 001	5879706	Jan	19, 2016	DP	U-530	I-570	Sep 02, 2011
	5879706	Jan	19, 2016	DP	U-894	NPP	Sep 02, 2011
	5879706*PED	Jul	19, 2016			PED	Mar 02, 2012
	6107302	Jan	19, 2016	DS	U-894	PED	Mar 02, 2012
	6107302	Jan	19, 2016	DS	U-530		
	6107302*PED	Jul	19, 2016				
<u>VALACYCLOVIR HYDROCHLORIDE - VALTREX</u>							
N020487 002	5879706	Jan	19, 2016	DP	U-894	I-570	Sep 02, 2011
	5879706	Jan	19, 2016	DP	U-530	NPP	Sep 02, 2011
	5879706*PED	Jul	19, 2016			PED	Mar 02, 2012
	6107302	Jan	19, 2016	DS	U-530	PED	Mar 02, 2012
	6107302	Jan	19, 2016	DS	U-894		
	6107302*PED	Jul	19, 2016				
<u>VALDECOXIB - BEXTRA</u>							
N021341 002	5633272	Feb	13, 2015		U-462		
<u>VALDECOXIB - BEXTRA</u>							
N021341 003	5633272	Feb	13, 2015		U-462		
<u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u>							
N021304 001	6083953	Mar	29, 2015	DS	DP U-384	D-125	Aug 05, 2013
	6083953	Mar	29, 2015	DS	DP U-854	I-604	Aug 28, 2012
	6083953*PED	Sep	29, 2015			PED	Feb 28, 2013
<u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u>							
N022257 001	6083953	Mar	29, 2015	DS	DP U-384	D-125	Aug 05, 2013
	6083953	Mar	29, 2015	DS	DP U-854	NDF	Aug 28, 2012
						PED	Feb 28, 2013
<u>VALSARTAN - DIOVAN</u>							
N020665 001	5399578	Mar	21, 2012		U-3		
	5399578*PED	Sep	21, 2012				
<u>VALSARTAN - DIOVAN</u>							
N020665 002	5399578	Mar	21, 2012		U-3		
	5399578*PED	Sep	21, 2012				
<u>VALSARTAN - DIOVAN</u>							
N021283 001	5399578	Mar	21, 2012				
	5399578*PED	Sep	21, 2012				
	5972990	Oct	26, 2016		U-692		
	5972990*PED	Apr	26, 2017				
	6294197	Jun	18, 2017		U-3		
	6294197*PED	Dec	18, 2017				
<u>VALSARTAN - DIOVAN</u>							
N021283 002	5399578	Mar	21, 2012				
	5399578*PED	Sep	21, 2012				
	5972990	Oct	26, 2016		U-692		
	5972990*PED	Apr	26, 2017				
	6294197	Jun	18, 2017		U-3		
	6294197*PED	Dec	18, 2017				
<u>VALSARTAN - DIOVAN</u>							
N021283 003	5399578	Mar	21, 2012				
	5399578*PED	Sep	21, 2012				
	5972990	Oct	26, 2016		U-692		
	5972990*PED	Apr	26, 2017				
	6294197	Jun	18, 2017		U-3		
	6294197*PED	Dec	18, 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
<u>VALSARTAN - DIOVAN</u>							
N021283 004	5399578	Mar	21, 2012				
	5399578*PED	Sep	21, 2012				
	5972990	Oct	26, 2016		U-692		
	5972990*PED	Apr	26, 2017				
	6294197	Jun	18, 2017		U-3		
	6294197*PED	Dec	18, 2017				
<u>VANDETANIB - VANDETANIB</u>							
N022405 001	7173038	Aug	14, 2021	DS	DP	NCE	Apr 06, 2016
	8067427	Aug	08, 2028		DP	ODE	Apr 06, 2018
	RE42353	Sep	23, 2017	DS	DP		
<u>VANDETANIB - VANDETANIB</u>							
N022405 002	7173038	Aug	14, 2021	DS	DP	NCE	Apr 06, 2016
	8067427	Aug	08, 2028		DP	ODE	Apr 06, 2018
	RE42353	Sep	23, 2017	DS	DP		
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>							
N021400 001	6362178	Oct	31, 2018	DS	DP	U-533	
	7696206	Oct	31, 2018	DS	DP	U-533	
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>							
N021400 002	6362178	Oct	31, 2018	DS	DP	U-533	
	7696206	Oct	31, 2018	DS	DP	U-533	
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>							
N021400 003	6362178	Oct	31, 2018	DS	DP	U-533	
	7696206	Oct	31, 2018	DS	DP	U-533	
<u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u>							
N021400 004	6362178	Oct	31, 2018	DS	DP	U-533	
	7696206	Oct	31, 2018	DS	DP	U-533	
<u>VARDENAFIL HYDROCHLORIDE - STAXYN</u>							
N200179 001	6362178	Oct	31, 2018			U-155	
	7696206	Oct	31, 2018			U-155	
<u>VARENICLINE TARTRATE - CHANTIX</u>							
N021928 001	6410550	May	10, 2020	DS	DP	U-56	
	6890927	May	06, 2022	DS	DP	U-56	M-105
	7265119	Aug	03, 2022	DS	DP	U-56	Jul 22, 2014
<u>VARENICLINE TARTRATE - CHANTIX</u>							
N021928 002	6410550	May	10, 2020	DS	DP	U-56	
	6890927	May	06, 2022	DS	DP	U-56	M-105
	7265119	Aug	03, 2022	DS	DP	U-56	Jul 22, 2014
<u>VELAGLUCERASE ALFA - VPRIV</u>							
N022575 001							NCE Feb 26, 2015
<u>VELAGLUCERASE ALFA - VPRIV</u>							
N022575 002							NCE Feb 26, 2015
<u>VEMURAFENIB - ZELBORAF</u>							
N202429 001	7504509	Oct	22, 2026	DS	DP	NCE	Aug 17, 2016
	7863288	Jun	20, 2029	DS	DP	ODE	Aug 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
VENLAFAXINE HYDROCHLORIDE - EFFEXOR XR						
N020699 001	5916923	Jun 28, 2013		U-398		
	5916923*PED	Dec 28, 2013		U-398		
	6274171	Mar 20, 2017				
	6274171*PED	Sep 20, 2017				
	6310101	Jun 28, 2013		U-46		
	6403120	Mar 20, 2017		U-451		
	6403120	Mar 20, 2017		U-535		
	6403120*PED	Sep 20, 2017		U-535		
	6403120*PED	Sep 20, 2017		U-451		
	6419958	Mar 20, 2017		U-459		
	6419958	Mar 20, 2017		U-535		
	6419958*PED	Sep 20, 2017		U-535		
	6419958*PED	Sep 20, 2017		U-459		
	6444708	Jun 28, 2013		U-398		
	6444708*PED	Dec 28, 2013		U-398		
VENLAFAXINE HYDROCHLORIDE - EFFEXOR XR						
N020699 002	5916923	Jun 28, 2013		U-398		
	5916923*PED	Dec 28, 2013		U-398		
	6274171	Mar 20, 2017				
	6274171*PED	Sep 20, 2017				
	6310101	Jun 28, 2013		U-46		
	6403120	Mar 20, 2017		U-451		
	6403120	Mar 20, 2017		U-535		
	6403120*PED	Sep 20, 2017		U-451		
	6403120*PED	Sep 20, 2017		U-535		
	6419958	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-459		
	6419958*PED	Sep 20, 2017		U-535		
	6419958*PED	Sep 20, 2017		U-459		
	6444708	Jun 28, 2013		U-398		
	6444708*PED	Dec 28, 2013		U-398		
VENLAFAXINE HYDROCHLORIDE - EFFEXOR XR						
N020699 003	5916923	Jun 28, 2013		U-398		
	5916923*PED	Dec 28, 2013		U-398		
	6274171	Mar 20, 2017				
	6274171*PED	Sep 20, 2017				
	6403120	Mar 20, 2017		U-535		
	6403120	Mar 20, 2017		U-451		
	6403120*PED	Sep 20, 2017		U-535		
	6403120*PED	Sep 20, 2017		U-451		
	6419958	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-459		
	6419958*PED	Sep 20, 2017		U-459		
	6419958*PED	Sep 20, 2017		U-535		
	6444708	Jun 28, 2013		U-398		
	6444708*PED	Dec 28, 2013		U-398		
VENLAFAXINE HYDROCHLORIDE - EFFEXOR XR						
N020699 004	5916923	Jun 28, 2013		U-398		
	5916923*PED	Dec 28, 2013		U-398		
	6274171	Mar 20, 2017				
	6274171*PED	Sep 20, 2017				
	6310101	Jun 28, 2013		U-46		
	6403120	Mar 20, 2017		U-535		
	6403120	Mar 20, 2017		U-451		
	6403120*PED	Sep 20, 2017		U-535		
	6403120*PED	Sep 20, 2017		U-451		
	6419958	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-459		
	6419958*PED	Sep 20, 2017		U-535		
	6419958*PED	Sep 20, 2017		U-459		
	6444708	Jun 28, 2013		U-398		
	6444708*PED	Dec 28, 2013		U-398		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote 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>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
N022104 001	6403120	Mar 20, 2017		U-839		
	6403120	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-839		
	6419958	Mar 20, 2017		U-535		
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
N022104 002	6403120	Mar 20, 2017		U-839		
	6403120	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-839		
	6419958	Mar 20, 2017		U-535		
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
N022104 003	6403120	Mar 20, 2017		U-839		
	6403120	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-839		
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
N022104 004	6403120	Mar 20, 2017		U-839		
	6403120	Mar 20, 2017		U-535		
	6419958	Mar 20, 2017		U-839		
	6419958	Mar 20, 2017		U-535		
<u>VERAPAMIL HYDROCHLORIDE - COVERA-HS</u>						
N020552 001	6096339	Apr 04, 2017		U-365		
<u>VERAPAMIL HYDROCHLORIDE - COVERA-HS</u>						
N020552 002	6096339	Apr 04, 2017		U-365		
<u>VERTEPORFIN - VISUDYNE</u>						
N021119 001	5095030	Sep 09, 2011	DS			
	5707608	Aug 02, 2015				
	5756541	Mar 11, 2016		U-357		
	5770619	Jan 06, 2015		U-357		
	5798349	Aug 25, 2015		U-357		
	6074666	Feb 05, 2012				
<u>VIGABATRIN - SABRIL</u>						
N020427 001					NCE	Aug 21, 2014
<u>VIGABATRIN - SABRIL</u>						
N022006 001					NCE	Aug 21, 2014
					ODE	Aug 21, 2016
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N022567 001	5532241	Sep 29, 2014	DS DP		NCE	Jan 21, 2016
	7834020	Jun 05, 2022	DS DP	U-839		
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N022567 002	5532241	Sep 29, 2014	DS DP		NCE	Jan 21, 2016
	7834020	Jun 05, 2022	DS DP	U-839		
<u>VILAZODONE HYDROCHLORIDE - VIIBRYD</u>						
N022567 003	5532241	Sep 29, 2014	DS DP		NCE	Jan 21, 2016
	7834020	Jun 05, 2022	DS DP	U-839		
<u>VORICONAZOLE - VFEND</u>						
N021266 001	5364938	Nov 15, 2011	DS			
	5567817	May 24, 2016	DS DP	U-540		
<u>VORICONAZOLE - VFEND</u>						
N021266 002	5364938	Nov 15, 2011	DS			
	5567817	May 24, 2016	DS DP	U-540		
<u>VORICONAZOLE - VFEND</u>						
N021267 001	5364938	Nov 15, 2011	DS			
	5567817	May 24, 2016	DS DP	U-540		
	6632803	Jun 02, 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
VORICONAZOLE - VFEND							
N021630 001	5364938	Nov	15, 2011	DS			
	5567817	May	24, 2016	DS DP	U-540		
VORINOSTAT - ZOLINZA							
N021991 001	6087367	Oct	04, 2011		U-776		
	7399787	Feb	09, 2025		U-892		
	7456219	Nov	14, 2026	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		
	RE38506	Nov	29, 2012	DS DP			
ZAFIRLUKAST - ACCOLATE							
N020547 001	5294636	Dec	11, 2011				
	5319097	Dec	11, 2011				
	5482963	Jan	09, 2013				
	5612367	Mar	18, 2014		U-189		
	6143775	Dec	11, 2011				
ZAFIRLUKAST - ACCOLATE							
N020547 003	5294636	Dec	11, 2011				
	5319097	Dec	11, 2011				
	5482963	Jan	09, 2013				
	5612367	Mar	18, 2014		U-189		
	6143775	Dec	11, 2011				
ZANAMIVIR - RELENZA							
N021036 001	5360817	Jul	26, 2013	DS DP			
	5648379	Jul	15, 2014		U-721		
	5648379	Jul	15, 2014		U-722		
	5648379	Jul	15, 2014		U-274		
	6294572	Dec	15, 2014	DS DP			
ZICONOTIDE - PRIALT							
N021060 003	5364842	Dec	30, 2016		U-48		
	5364842	Dec	30, 2016		U-55		
	5795864	Jun	27, 2015	DP			
	5859186	Dec	30, 2011		U-55		
	5859186	Dec	30, 2011		U-48		
ZICONOTIDE ACETATE - PRIALT							
N021060 001	5364842	Dec	30, 2016		U-48		
	5364842	Dec	30, 2016		U-55		
	5795864	Jun	27, 2015	DP			
	5859186	Dec	30, 2011		U-55		
	5859186	Dec	30, 2011		U-48		
ZICONOTIDE ACETATE - PRIALT							
N021060 002	5364842	Dec	30, 2016		U-48		
	5364842	Dec	30, 2016		U-55		
	5795864	Jun	27, 2015	DP			
	5859186	Dec	30, 2011		U-48		
	5859186	Dec	30, 2011		U-55		
ZICONOTIDE ACETATE - PRIALT							
N021060 004	5364842	Dec	30, 2016		U-55		
	5364842	Dec	30, 2016		U-48		
	5795864	Jun	27, 2015	DP			
	5859186	Dec	30, 2011		U-55		
	5859186	Dec	30, 2011		U-48		
ZILEUTON - ZYFLO CR							
N022052 001	5422123	Jun	06, 2012	DP			
	6183778	Sep	21, 2013	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 DELIST REQUESTED		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>								
N020825 001	4831031	Mar 02, 2012	DS	DP	U-720		I-615	Nov 20, 2012
	5312925	Sep 01, 2012	DS	DP				
	6150366	May 27, 2019		DP				
	6245766	Dec 18, 2018			U-601			
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>								
N020825 002	4831031	Mar 02, 2012	DS	DP	U-720		I-615	Nov 20, 2012
	5312925	Sep 01, 2012	DS	DP				
	6150366	May 27, 2019		DP				
	6245766	Dec 18, 2018			U-601			
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>								
N020825 003	4831031	Mar 02, 2012	DS	DP	U-720		I-615	Nov 20, 2012
	5312925	Sep 01, 2012	DS	DP				
	6150366	May 27, 2019		DP				
	6245766	Dec 18, 2018			U-601			
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>								
N020825 004	4831031	Mar 02, 2012	DS	DP	U-720		I-615	Nov 20, 2012
	5312925	Sep 01, 2012	DS	DP				
	6150366	May 27, 2019		DP				
	6245766	Dec 18, 2018			U-601			
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>								
N021483 001	4831031	Mar 02, 2012	DS	DP	U-720			
	5312925	Sep 01, 2012	DS	DP	U-720			
	6150366	May 27, 2019		DP	U-719			
	6245766	Dec 18, 2018			U-601			
	7175855	May 18, 2020		DP				
<u>ZIPRASIDONE MESYLATE - GEODON</u>								
N020919 001	4831031	Mar 02, 2012	DS	DP	U-720			
	6110918	Mar 26, 2017						
	6232304	Apr 01, 2017						
	6399777	Apr 01, 2017						
<u>ZOLEDRONIC ACID - RECLAST</u>								
N021817 001	4939130	Sep 02, 2012	DS	DP	U-662		I-595	May 29, 2012
	4939130*PED	Mar 02, 2013					I-584	Mar 15, 2012
	7932241	Feb 05, 2028		DP			I-581	Dec 19, 2011
	7932241*PED	Aug 05, 2028						
	8052987	Mar 19, 2024			U-1199			
<u>ZOLEDRONIC ACID - ZOMETA</u>								
N021223 001	4939130	Sep 02, 2012	DS	DP	U-53			
	4939130*PED	Mar 02, 2013						
<u>ZOLEDRONIC ACID - ZOMETA</u>								
N021223 002	4939130	Sep 02, 2012	DS	DP	U-53		PED	Sep 20, 2011
	4939130*PED	Mar 02, 2013						
<u>ZOLEDRONIC ACID - ZOMETA</u>								
N021223 003	4939130	Sep 02, 2012	DS	DP	U-53			
	4939130*PED	Mar 02, 2013						
	7932241	Feb 05, 2028		DP				
<u>ZOLMITRIPTAN - ZOMIG</u>								
N020768 001	5466699	Nov 14, 2012						
	5466699*PED	May 14, 2013						
	5863935	Nov 14, 2012						
	5863935*PED	May 14, 2013						
<u>ZOLMITRIPTAN - ZOMIG</u>								
N020768 002	5466699	Nov 14, 2012						
	5466699*PED	May 14, 2013						
	5863935	Nov 14, 2012						
	5863935*PED	May 14, 2013						

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ZOLMITRIPTAN - ZOMIG							
N021450 004	5466699	Nov	14, 2012			U-436	
	5466699*PED	May	14, 2013				
	6750237	Nov	28, 2020		DP		
	6750237*PED	May	28, 2021				
	7220767	Nov	28, 2020		DP		
	7220767*PED	May	28, 2021				
ZOLMITRIPTAN - ZOMIG-ZMT							
N021231 001	5466699	Nov	14, 2012				
	5466699*PED	May	14, 2013				
ZOLMITRIPTAN - ZOMIG-ZMT							
N021231 002	5466699	Nov	14, 2012				
	5466699*PED	May	14, 2013				
ZOLPIDEM TARTRATE - AMBIEN CR							
N021774 001	6514531	Dec	01, 2019		DP		
	6514531*PED	Jun	01, 2020				
ZOLPIDEM TARTRATE - AMBIEN CR							
N021774 002	6514531	Dec	01, 2019		DP		
	6514531*PED	Jun	01, 2020				
ZOLPIDEM TARTRATE - EDLUAR							
N021997 001	6761910	Sep	24, 2019		DP	U-674	
ZOLPIDEM TARTRATE - EDLUAR							
N021997 002	6761910	Sep	24, 2019		DP	U-674	
ZOLPIDEM TARTRATE - INTERMEZZO							
N022328 001	7658945	Apr	15, 2027		DP	U-1194	
	7682628	Feb	16, 2025			U-1194	
ZOLPIDEM TARTRATE - INTERMEZZO							
N022328 002	7658945	Apr	15, 2027		DP	U-1194	
	7682628	Feb	16, 2025			U-1194	
ZOLPIDEM TARTRATE - ZOLPIMIST							
N022196 001	7632517	Oct	01, 2017			U-70	

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 51

PATENT & EXCLUSIVITY ABBREVIATIONS

D	NEW DOSING SCHEDULE (SEE INDIVIDUAL REFERENCES)
I	NEW INDICATION (SEE INDIVIDUAL REFERENCES)
M	MISCELLANEOUS EXCLUSIVITY CODES (SEE INDIVIDUAL REFERENCES)
NC	NEW COMBINATION
NCE	NEW CHEMICAL ENTITY
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
PC	PATENT CHALLENGE
PED	PEDIATRIC EXCLUSIVITY
RTO	RX TO OTC SWITCH OR OTC USE
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
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 PHOPHYLAXIX 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

PATENT AND EXCLUSIVITY TERMS

ADB 2 of 51

EXCLUSIVITY DOSING SCHEDULE

- 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 ADIMISTRATION OF CISATRICURIUM 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 CONVERT 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/M² FOLLOWED BY CISPLATIN AT A DOSE OF 75MG/M² 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 3 of 51

EXCLUSIVITY DOSING SCHEDULE

- 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
- 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 DEFECIT 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 BEFPRE 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

PATENT AND EXCLUSIVITY TERMS

ADB 4 of 51

EXCLUSIVITY DOSING SCHEDULE

- D-117 50 MG TABLET FOR INITIATION OF DOSE TITRATION FOR BIPOLEAR 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 REGIMEN ADJUSTMENTS
- D-121 CHANGE TO REMOVE 20 MG MAXIMUM DOSAGE RESTRICTION
- 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 DARUNIVIR 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

EXCLUSIVITY INDICATION

- I-1 DYSMENORRHEA
- I-2 CHOLANGIOPANCREATOGRAPHY
- I-3 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
- I-4 PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
- I-5 HYSTEROSALPINGOGRAPHY
- I-6 TREATMENT OF JUVENILE ARTHRITIS
- I-7 BIOPSY PROVEN MINIMAL CHANGE NEPHROTIC SYNDROME IN CHILDREN
- I-8 ADULT INTRAVENOUS CONTRAST-EHANCED 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

PATENT AND EXCLUSIVITY TERMS

ADB 5 of 51

EXCLUSIVITY INDICATION

- 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 OMNIPACQUE 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
 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 ANGIOCARDIOPHY
 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

PATENT AND EXCLUSIVITY TERMS

ADB 6 of 51

EXCLUSIVITY INDICATION

- 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
- 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 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

PATENT AND EXCLUSIVITY TERMS

ADB 7 of 51

EXCLUSIVITY INDICATION

- 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 EROSIONAL 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 (AaDO₂) 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 8 of 51

EXCLUSIVITY INDICATION

- 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 PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE)
- I-187 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 REGIMENT, 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 SULFONYL UREAS 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

PATENT AND EXCLUSIVITY TERMS

ADB 9 of 51

EXCLUSIVITY INDICATION

- 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)
- I-219 USE OF LIPITOR BY PATIENTS WITH PRIMARY DYSBETA LIPOPROTEINEMIA (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 PRIOSEC (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
- 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 SODIUM AND OUTPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM

PATENT AND EXCLUSIVITY TERMS

ADB 10 of 51

EXCLUSIVITY INDICATION

- 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 SYMPTOMATIC 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 INFLAMMATION
- 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 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

PATENT AND EXCLUSIVITY TERMS

ADB 11 of 51

EXCLUSIVITY INDICATION

- 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 12 of 51

EXCLUSIVITY INDICATION

- 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 SYSTOMS 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
- 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-CONSTRICTION 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 ANTI COAGULANT 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 HYPERSECRETRY 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 THE 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

PATENT AND EXCLUSIVITY TERMS

ADB 13 of 51

EXCLUSIVITY INDICATION

- 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
- 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 INTRAOCCULAR 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>=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 RECUR AFTER STEM CELL TRNSPLT OR RESIST TO INTERFERON ALPHA THERAPY. NO CONTROLLED TRIALS DEMONSTRATING A CLINICAL BENEFIT SUCH AS IMPROVE 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
- 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, HYPOMANIA, MIXED EPISODES) IN PATIENTS TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- I-405 TREATMENT OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) USING AN INTERMITTENT DOSING REGIMENT
- I-406 PREVENTION OF CYTOMEGALOVIRUS DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPositive/RECIPIENT CMV SERONEGATIVE)

PATENT AND EXCLUSIVITY TERMS

ADB 14 of 51

EXCLUSIVITY INDICATION

- 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 FLOXATIN 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 PACLITAXEL FOR THE FIRST-LINE TREATMENT OF PATIENTS 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

PATENT AND EXCLUSIVITY TERMS

ADB 15 of 51

EXCLUSIVITY INDICATION

- 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 EROSiVE 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/HYPOPEmA 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 BOTH AGENTS DO NOT RESULT iN ADEQUATE GLYCEmiC CONTROL (TRIPLE THERAPY)
- I-454 MAINTENANCE OF CLiNiCAL REMiSSION OF MiLD TO MODERATE CHRONiS 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 COMPLiATIONS
- 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 KiDiNEY 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 iNFECTION
- 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 iN PATIENTS WITH LOCALLy ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER
- I-474 TREATMENT OF IRON DEFiCiENCY ANEMiA iN PERITONEAL DIALYSIS DEPENDANT CHRONiC KiDiNEY DISEASE iN PATIENTS RECEIVING AN ERYTHROPOiETiN
- I-475 PREVENTION OF NAUSEA AND VOMiTing ASSOCIATED WITH iNITAL AND REPEAT COURSES OF MODERATELY EMETOGEniC CANCER CHEMOTHERAPY
- I-476 TREATMENT OF DiABETiC FOOT iNFECTIONS WITHOUT OSTeOmyELiTIS

PATENT AND EXCLUSIVITY TERMS

ADB 16 of 51

EXCLUSIVITY INDICATION

- 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. THETAIOTAOOMICRON 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
- 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 EPIPHYES
- 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 HYPERSECRETOORY 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
- I-510 ADULT DERMAFIBROSARCOMA PROTUBERANS (DFSP)
- I-511 ADULT MYELODYSPLASTIC SYNDROME/MYELOPROLIFERATIVE DISEASES (MDS/MDP)
- I-512 ADULT PH+ ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) MONOTHERAPY

PATENT AND EXCLUSIVITY TERMS

ADB 17 of 51

EXCLUSIVITY INDICATION

- 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 HOMEBOX 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 RECCURENCE 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 BIPOLAR 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
- 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 PERITONIS AND ABSCESSSES

PATENT AND EXCLUSIVITY TERMS

ADB 18 of 51

EXCLUSIVITY INDICATION

- 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 EXACERATIONS 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 DISORDER (ADHD) IN CHILDREN AND ADOLESCENTS
- I-563 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES 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
- 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 (FREDICKSON 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 GASTROINTESTINAL 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

PATENT AND EXCLUSIVITY TERMS

ADB 19 of 51

EXCLUSIVITY INDICATION

- 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 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 POSTMENARCHAL 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 POSTMENARCHAL 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 EROSIVE 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 20 of 51

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)

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
- 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 & PK INFORMATION IN CHILDREN 6 MONTHS TO LESS THAN 6 YEARS OF AGE ADDED TO PKG 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

PATENT AND EXCLUSIVITY TERMS

ADB 21 of 51

EXCLUSIVITY MISCELLANEOUS

- 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 TRUOPT HAVE BEEN DEMONSTRATED IN PEDIATRIC PATIENTS IN A 3 MONTH, MULTI-CENTER DOUBLE MASKED ACTIVE-TREATMENT-CONTROLLED 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 DENISTY 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 PRMTRS
- M-59 RESULTS OF THE T20-310 STUDY WHICH EVALUATED THE PHARMACOKINETICS, SAFETY, AND ANTIVIRAL ACTIVITY OF FUZEON IN TREATMENT EXPERIENCED PEDIATRIC SUBJECTS AND ADOLSCENTS 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 ROSUVASTATIN 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

PATENT AND EXCLUSIVITY TERMS

ADB 22 of 51

EXCLUSIVITY MISCELLANEOUS

- 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
- 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 ARGAGATROBAN IN CERTAIN PEDIATRIC PATIENTS WITH HEPARIN-INDUCED THROMBOCYTOPENIA (HIT) OR HEPARIN-INDUCED THROMBOCYTOPENIA WITH THROMBOSIS (HITT)
- 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 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

PATENT AND EXCLUSIVITY TERMS

ADB 23 of 51

EXCLUSIVITY MISCELLANEOUS

- 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

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 HYPOTHALMIC 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 ENZYMO PATHIES
- U-15 METHOD OF LOWERING INTRAOCULAR 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 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

PATENT AND EXCLUSIVITY TERMS

ADB 24 of 51

PATENT USE

- U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCERAL 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
- U-62 CORONARY ARTERIOGRAPHY, LEFT VENTRICULOGRAPHY, CT IMAGING OF THE BODY, INTRAVENOUS EXCRETORY UROGRAPHY, INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY AND VENOGRAPHY
- U-63 ISOPRENALENE 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 HEARTBURNDUE 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

PATENT AND EXCLUSIVITY TERMS

ADB 25 of 51

PATENT USE

- 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-CONTRAINDICATED
 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
 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 EROSIONAL ESOPHAGITIS, POORLY RESPONSIVE SYMPTOMATIC GERD AND PATHOLOGICAL HYPERSECRETORY CONDITIONS AND MAINTENANCE HEALING OF EROSIONAL 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 PERSSARY
 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 H₂-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

PATENT AND EXCLUSIVITY TERMS

ADB 26 of 51

PATENT USE

- 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
- 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 HYPERCHOLESTROLEMIA
- 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 27 of 51

PATENT USE

- U-184 TREATING ALLERGIC EYE DISEASES IN HUMANS
 U-185 METHOD OF TREATING HYPERTENSION
 U-186 METHOD FOR TREATING GI DISORDERS CAUSED BY H. RYLORI WHICH COMPRISSES 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 ISOTOAC 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 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 MEHTOD 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

PATENT AND EXCLUSIVITY TERMS

ADB 28 of 51

PATENT USE

- 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 HYPERSECRETORY 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
- 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 INTRAOCULAR 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 DYSENTERIA; ACUTE TREATMENT OF MIGRAINE ATTACKS IN ADULTS
- U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMINISTERING 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

PATENT AND EXCLUSIVITY TERMS

ADB 29 of 51

PATENT USE

- 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 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 VASOCONSTRICKTION 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 POLYPS
 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 COMPRISES 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
 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

PATENT AND EXCLUSIVITY TERMS

ADB 30 of 51

PATENT USE

- 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 CONDITIONS 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 PHARMACOKINETICS OF A DRUG THAT IS METABOLIZED BY CYTOCHROME P450 MONOOXYGENASE BY ADMINISTERING 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 ADMINISTRATION WITH A REVERSE TRANSCRIPTASE INHIBITOR
- U-351 INHIBITING PROTEASE WITH LOPINAVIR AND INHIBITING AN HIV INFECTION WITH LOPINAVIR
- U-352 INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH A REVERSE TRANSCRIPTASE INHIBITOR
- U-353 PREVENTION AND TREATMENT OF OSTEOPOROSIS
- 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 ADMINISTER 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

PATENT AND EXCLUSIVITY TERMS

ADB 31 of 51

PATENT USE

- 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 RECURRENCE
- 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 32 of 51

PATENT USE

- U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASING 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 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 LEVOCARITINE 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 REGIMENT 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

PATENT AND EXCLUSIVITY TERMS

ADB 33 of 51

PATENT USE

- U-447 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- 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 RECURRENT
- 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 HYPERPLASIA
- 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
- 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)

PATENT AND EXCLUSIVITY TERMS

ADB 34 of 51

PATENT USE

- 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 CHAMBER 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 35 of 51

PATENT USE

- 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 RESONACE 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.
- 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 PERIDONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIDONTAL 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 THYROGLOBULIN (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)

PATENT AND EXCLUSIVITY TERMS

ADB 36 of 51

PATENT USE

- 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 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 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 SERTONERGIC AND NORADRENERGIC ACTIVITY IN THE CNS
- U-606 USE OF IRINOTECAN IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLRECTAL CANCER

PATENT AND EXCLUSIVITY TERMS

ADB 37 of 51

PATENT USE

- 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 IIA, 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 POLYPHS
- 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
- 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 ANTI NEOPLASTIC 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 EROSIVE 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

PATENT AND EXCLUSIVITY TERMS

ADB 38 of 51

PATENT USE

- 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 EXEDIN 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 39 of 51

PATENT USE

- 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 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 (AVP) 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 EXERCISE-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 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)

PATENT AND EXCLUSIVITY TERMS

ADB 40 of 51

PATENT USE

- 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
- 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 REVLIIMID (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 INTRAOCCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOME 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

PATENT AND EXCLUSIVITY TERMS

ADB 41 of 51

PATENT USE

- 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 42 of 51

PATENT USE

- 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
- 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 INTRAOCCULAR 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 43 of 51

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 HYPERVOLMIC 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 CLEVIDIPREX 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 44 of 51

PATENT USE

- 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 ALLOGENIC 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 ABY SUSCEPTIBLE STRAINS
- U-925 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTILES) 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)
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 45 of 51

PATENT USE

- 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 EROSION ESOPHAGITIS
- U-948 TREATMENT OF DIABETES MELLITUS
- U-949 HEALING OF ALL GRADES OF EROSION ESOPHAGITIS (EE) FOR UP TO 8 WEEKS
- U-950 MAINTAIN HEALING OF EROSION 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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 46 of 51

PATENT USE

- 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
- 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 REDUCT ELEVATED TOTAL CHOLESTEROL (TC), LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES, AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIX DYSLIPIDEMIA
- U-997 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY DOSING AT INTERVALS OF 24 HOURS
- U-998 ADJUNCITVE 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
- 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)

PATENT AND EXCLUSIVITY TERMS

ADB 47 of 51

PATENT USE

- 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 MUSCULSKELETAL 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
- 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 MEDICATED DISEASE IN A MAMMAL BY ADMININSTERING 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 DSYFUNCTION 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

PATENT AND EXCLUSIVITY TERMS

ADB 48 of 51

PATENT USE

- 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 REVLIMID (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
- 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

PATENT AND EXCLUSIVITY TERMS

ADB 49 of 51

PATENT USE

- 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 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 REDUCTINO 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 TRETMENT 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 HYPERTRIGLYCDERIDEMIA 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
- 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 PATINETS WITH HYPERPHENYLALANINEMIA (HPA)

PATENT AND EXCLUSIVITY TERMS

ADB 50 of 51

PATENT USE

- 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 BROCHODILATOR 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 WIHT 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
- 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 SULFYONYLUREA(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 MELITUS 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

PATENT AND EXCLUSIVITY TERMS

ADB 51 of 51

PATENT USE

- 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