

APPROVED DRUG PRODUCTS

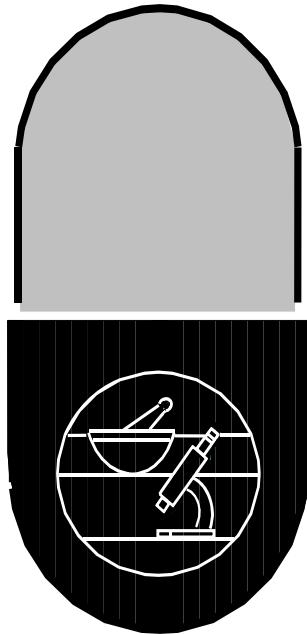
With Therapeutic Equivalence Evaluations



The "Orange Book"

FDA data supplied by [DrugPatentWatch.com](https://www.drugpatentwatch.com)

**CUMULATIVE
SUPPLEMENT 07
July 2009**



**APPROVED
DRUG PRODUCTS**

**WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS**

29th EDITION

**Department of Health and Human Services
Food and Drug Administration
Center for Drug Evaluation and Research
Office of Generic Drugs**

2009

Prepared By
Office of Generic Drugs
Center for Drug Evaluation and Research
Food and Drug Administration

**APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS**

29th EDITION

Cumulative Supplement 7

July 2009

CONTENTS

	<i>PAGE</i>
1.0 INTRODUCTION	iii
1.1 How to use the Cumulative Supplement	iii
1.2 Cumulative Supplement Content.....	iv
1.3 Applicant Name Changes.....	v
1.4 Availability of the Edition	vi
1.5 Report of Counts for the Prescription Drug Product List	vi
1.6 Cumulative Supplement Legend	vii
 DRUG PRODUCT LISTS	
Prescription Drug Product List	1-1
OTC Drug Product List	2-1
Drug Products with Approval under Section 505 of the Act	
Administered by the Center for Biologics Evaluation and Research List.....	3-1
Orphan Product Designations and Approvals List	4-1
Drug Products Which Must Demonstrate in vivo Bioavailability	
Only if Product Fails to Achieve Adequate Dissolution	5-1
 PATENT AND EXCLUSIVITY INFORMATION ADDENDUM	
A. Patent and Exclusivity Lists	A-1
B. Patent and Exclusivity Terms	B-1

**APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS**

29th EDITION

**CUMULATIVE SUPPLEMENT 7
July 2009**

1.0 INTRODUCTION

1.1 HOW TO USE THE CUMULATIVE SUPPLEMENT

This Cumulative Supplement is one of a series of monthly updates to the Approved Drug Products with Therapeutic Equivalence Evaluations, 29th Edition (the List). The List is composed of four parts: approved prescription drug products with therapeutic equivalence evaluations; over-the-counter (OTC) drug products that require approved applications as a condition of marketing; drug products with approval under Section 505 of the Act administered by the Center for Biologics Evaluation and Research; and products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.

The Cumulative Supplement provides, among other things, information on newly approved drugs and, if necessary, revised therapeutic equivalence evaluations and updated patent and exclusivity data. The Addendum contains appropriate drug patent and exclusivity information required of the Agency by the "Drug Price Competition and Patent Term Restoration Act of 1984" for the Prescription, OTC, and Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research Lists.

Because all parts of the publication are subject to changes, additions, or deletions, the List must be used in conjunction with the most current Cumulative Supplement. Users may wish to mark to the left of the ingredient(s) in the List to indicate that changes to that entry appear in the Cumulative Supplement. Drug product information is provided in each Cumulative Supplement for completeness to assist in locating the proper place in the List for the revision.

The presence of any therapeutic equivalence code indicates that the drug product is multisource; the deletion of a therapeutic equivalence code indicates that the drug product has become single source. (An infrequent exception exists when a therapeutic equivalence code is revised. In that case, the deletion of the therapeutic equivalence code is followed immediately by the addition of the revised one.)

Products that have never been marketed, are for exportation, are for military use, or have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons, will be flagged in this Cumulative Supplement with the "@" symbol to designate their non-marketed status. All products having a "@" symbol in the 12th Cumulative Supplement of the 29th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 30th Edition. The current Edition Section 2., How To Use The Drug Product Lists, describes the layout and usage of the List.

New additions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >A>. The Patent and Exclusivity List new additions are indicated by the symbol >A> to the left of Patent Number or Exclusivity Code. The >A> symbol is then dropped in subsequent Cumulative Supplements for that item.

New deletions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >D> (DELETE) to the left of the line. The information line with the >D> symbol is dropped in subsequent Cumulative Supplements for that item.

The Patent and Exclusivity List is arranged in alphabetical order by active ingredient name(s) and trade name. The trade name will follow the active ingredient name separated by a dash symbol. Also shown is the application number and product number (FDA's internal file number) for reference purposes. All patents with their expiration dates are displayed for each application number. Drug substance and drug product patents are indicated as such with DS or DP in the Patent codes column. Use patents are indicated with the symbol "U" followed by a number representing a specific use. Exclusivity information for a specific drug is indicated by an abbreviation followed by the date upon which the exclusivity expires. Refer to the Exclusivity Terms, Section B, in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations. Refer to Section 1.3 for internet access to the most current list of Patent and Exclusivity terms.

1.2 CUMULATIVE SUPPLEMENT CONTENT

Since February 2005, we have been providing daily Electronic Orange Book (EOB) product information for new generic drug approvals. Daily generic updates provide the consumer with the current list of approved generic products which is important for substitution purposes. Previously, a first-time-generic product approved early in the month would not be published in the Cumulative Supplement (CS) for several weeks.

The CS monthly update publish goal is by the end of the following month's second work week (e.g., November's supplement will be updated by the end of the second full work week in December).

Currently, the monthly PDF CS includes:

- Generic product ANDA (Abbreviated New Drug Approval) approvals as of the date of publication.
- All product changes received and processed as of the date of publication.
 - o Refer to CS Section 1.8 Cumulative Supplement Legend for types of changes
 - o Discontinued products will be processed as of the date of publication. There will be circumstances where a product is discontinued in one month, however, it will be reported in a different month's CS. For example, the Orange Book Staff received a letter November 7 that the product has been discontinued from manufacturing and marketing. The Orange Book subsequently publishes the October CS on November 14. The product will show in the October CS that it is discontinued even though the date of discontinuance is the day that the Orange Book Staff receives notification (November 7).
- New Drug Application (NDA) approvals (20,000 and 50,000 series) appear in the CS month they were approved.

- Patent information, also updated daily in the EOB, is current to the date of publication.
- Exclusivity information is updated monthly and current to the date of publication.

Every effort is made to ensure the Cumulative Supplement is current and accurate. Applicant holders are requested to inform the FDA Orange Book Staff (OBS) of any changes or corrections. The OBS can be contacted by email at drugproducts@cder.fda.gov. Send Changes by FAX: 240-276-8974; mail to:

FDA/CDER Orange Book Staff
Office of Generic Drugs, HFD-610
7500 Standish Place
Rockville, MD 20855-2773

1.3 APPLICANT NAME CHANGES

It is not practical to identify in the Cumulative Supplement each and every product involved when an applicant transfers its entire line of approved drug products to another applicant, or when an applicant changes its name. Therefore, the cumulation of these transfers and name changes will be identified in this section only. Where only partial lines of approved products are transferred between applicants, each approved product involved will appear as an applicant name change entry in the Cumulative Supplement.

It is also not practical to identify each and every product involved when an applicant name is changed to meet internal publication standards (e.g., MSD or Zenith [Former Abbreviated Names] are changed, respectively, to Merck Sharp Dohme or Zenith Labs [New Abbreviated Names]). When this occurs, each product involved (either currently in the Cumulative Supplement or in the following year's edition) will reflect the new abbreviated name. Consequently, it will not appear as an applicant name change entry in the Cumulative Supplement nor will the cumulation of these name changes appear in this section. The Electronic Orange Book Query, updated monthly, will contain the most current applicant holder name.

<u>FORMER APPLICANT NAME</u> <u>(FORMER ABBREVIATED NAME)</u>	<u>NEW APPLICANT NAME</u> <u>(NEW ABBREVIATED NAME)</u>
ABLE LABORATORIES INC (ABLE)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARM)
AXIOM PHARMACEUTICAL CORP (AXIOM PHARM)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARM)
DABUR ONCOLOGY PLC (DABUR ONCOLOGY PLC)	FRESENIUS KABI ONCOLOGY PLC (FRESENIUS KABI ONCOL)
HALSEY DRUG CO INC (HALSEY)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARM)
IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS (IVAX PAHRMS)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARM)

NORTON WATERFORD LTD
(NORTON WATERFORD)

IVAX PHARMACEUTICALS INC SUB TEVA
PHARMACEUTICALS USA
(IVAX SUB TEVA PHARM)

TORPHARM INC
(TORPHARM)

APOTEX INC
(APOTEX)

ZENITH GOLDLINE
(ZENTH GOLDLINE)

IVAX PHARMACEUTICALS INC SUB TEVA
PHARMACEUTICALS USA
(IVAX SUB TEVA PHARM)

1.4 AVAILABILITY OF THE EDITION

Since 1997, the Electronic Orange Book Query (EOBQ) <http://www.accessdata.fda.gov/scripts/cder/ob/default.cfm>, has been available on the internet and has become the updated-every-month Orange Book. The Query provides searching of the approved drug list by active ingredient, proprietary name, applicant holder, applicant number or patent number. Product search categories are: prescription, over-the-counter, discontinued drugs. There are links to patent and exclusivity information that may be applicable to each product.

Commencing with the 25th edition, the Annual Edition and monthly Cumulative Supplements have been provided in downloadable Portable Document Format (PDF) at the EOB home page by clicking on Publications. The PDF annual and cumulative supplements duplicate previous paper versions. Over time, there will be an archive for the annuals and each year's December Cumulative Supplement.

The downloaded Annual Edition and Cumulative Supplements are also available in a paper version (Approved Drug Products with Therapeutic Equivalence Evaluations, ADP) from the U.S. Government Printing Office: <http://bookstore.gpo.gov>; toll free 866-512-1800.

There are historical lists of Orange Book cumulative supplement product monthly changes at <http://www.fda.gov/Drugs/InformationOnDrugs/ucm086229.htm>. There are ASCII text files of the Orange Book drug product, patent, and exclusivity data at <http://www.fda.gov/Drugs/InformationOnDrugs/ucm129689.htm>. The drug product text files are provided in eobzip.zip format. The files are updated concurrently with the monthly cumulative supplements. The annual Orange Book Edition Appendices A, B, and C in PDF format are updated quarterly.

Effective August 18, 2003, patent submissions for publication in the Orange Book and Docket *95S-0117 need to be submitted on form FDA-3542 which may be downloaded from the FDA Forms List, <http://www.fda.gov/opacom/morechoices/fdaforms/default.html>.

The current listing of the Orphan Product Designations and Approvals is available at <http://www.fda.gov/orphan/designat/list.htm>.

1.5 REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST

DESCRIPTION OF REPORT

This report provides summary counts derived from the product information in the Prescription Drug Product List and the current Cumulative Supplement. Products included in the counts are domestically marketed drug products approved for both safety and effectiveness under section 505 of the Federal Food, Drug, and Cosmetic Act. Excluded are approved drug products marketed by distributors; those marketed solely abroad; and

those now regarded as medical devices, biologics or foods.

The baseline column (Dec 2008) refers to the products in the Prescription Drug Product List. For each three-month period, a column of quarterly data is added which incorporates counts of product activity from the previous quarter(s) with those in the baseline count.

DEFINITIONS

Drug Product

For this report, a drug product is the representation in the Prescription Drug Product List of an active moiety (molecular entity and its salts, esters and derivatives) either as a single ingredient or as a combination product provided in a specific dosage form and strength for a given route of administration with approval for marketing by a firm under a particular generic or trade name.

New Molecular Entity

A new molecular entity is considered an active moiety that has not previously been approved (either as the parent compound or as a salt, ester or derivative of the parent compound) in the United States for use in a drug product either as a single ingredient or as part of a combination.

REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST COUNTS CUMULATIVE BY QUARTER

<u>CATEGORIES COUNTED</u>	<u>DEC 2008</u>	<u>MAR 2009</u>	<u>JUN 2009</u>	<u>SEPT 2009</u>
DRUG PRODUCTS LISTED	12751	12910	13027	
SINGLE SOURCE	2433 (19.1%)	2449 (19.0%)	2451 (18.8%)	
MULTISOURCE	10229 (80.2%)	10372 (80.3%)	10487 (80.5%)	
THERAPEUTICALLY EQUIVALENT	10072 (79.0%)	10216 (79.1%)	10333 (79.3%)	
NOT THERAPEUTICALLY EQUIVALENT	157 (1.2%)	156 (1.2%)	154 (1.2%)	
EXCEPTIONS ¹	89 (0.7%)	89 (0.7%)	89 (0.7%)	
NEW MOLECULAR ENTITIES APPROVED	15	5	6	
NUMBER OF APPLICANTS	719	724	732	

¹Amino acid-containing products of varying composition (see Introduction, page xx of the List).

1.6 CUMULATIVE SUPPLEMENT LEGEND

The List is sorted by Ingredient(s) and, within each grouping, by the Dosage Form; Route and then by trade name.

The individual product record contains the Therapeutic Equivalence Code, Reference Listed Drug symbol, applicant holder, strength(s), New Drug Application number, product number, and approval date. The last two columns describe the action. The Action Month is the CS month the action occurred. The OB Action is the type of change that has occurred.

New ingredient(s), new dosage form; route(s), new trade names, and new product additions are preceded by >A> during the action month. The change month is the current CS month; the change code for new approvals is NEWA.

Following months will display the same information without the >A>.

Changes to currently listed products will list two records. The deleted product record will be preceded by >D>. The product record change addition being made will be preceded by >A>. Following months will display only the >A> record without the >A>. All changes that occur to the product through the Annual year will be listed. The change month and change code will document the change.

The change code and description:

NEWA	New drug product approval usually in the supplement month.
CAHN	Applicant holder firm name has changed.
CAIN	Change. There has been a change in the Ingredient(s) name. All products will be deleted under the old name and all products will be added under the changed ingredient(s) name.
CDFR	Change. Dosage Form; Route of Administration.
CFTG	Change. A first time generic for the innovator product. A TE Code is added.
CMFD	Change. The product is moved from the Discontinued Section due to a change in marketing status.
CMS1	Change. Miscellaneous addition to list.
CMS2	Change. Miscellaneous deletion from list.
CPOT	Change. Potency amount/unit.
CRLD	Change. Reference Listed Drug.
CTEC	Change. Therapeutic Equivalence Code.
CTNA	Change. Trade Name.
DISC	Discontinued. The Rx or OTC listed product is not being marketed and will be moved to the discontinued section in the next edition.

PRESCRIPTION DRUG PRODUCT LIST - 28TH EDITION

RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

1-1

ACARBOSE

TABLET; ORAL

ACARBOSE

AB	IMPAX LABS	25MG	N78441 001	May 14, 2009	May	NEWA
AB		50MG	N78441 002	May 14, 2009	May	NEWA
AB		100MG	N78441 003	May 14, 2009	May	NEWA

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

>D>	AB	MUTUAL PHARM	325MG;50MG;40MG	N40601 001	Jul 29, 2005	Jul	DISC
>A>	@		325MG;50MG;40MG	N40601 001	Jul 29, 2005	Jul	DISC

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

@ SANDOZ 300MG;30MG

N81250 001 Jul 16, 1992 Mar DISC

@ 300MG;60MG

N81249 001 Jul 16, 1992 Mar DISC

TYLENOL W/ CODEINE NO. 4

AA	ORTHO MCNEIL JANSSEN	300MG;60MG	N85055 004		Mar	CMFD
----	----------------------	------------	------------	--	-----	------

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

@ MALLINCKRODT 500MG;5MG

N89006 001 Aug 09, 1985 Feb CTNA

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

PERCOCET

AA	+	ENDO PHARMS	325MG;2.5MG	N40330 001	Jun 25, 1999	May	CTEC
----	---	-------------	-------------	------------	--------------	-----	------

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL

PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN

@ SANDOZ 650MG;65MG

N89959 001 Jul 18, 1989 Mar DISC

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

@ SANDOZ 650MG;100MG

N70443 001 Jan 23, 1986 Mar DISC

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

AB	MYLAN	325MG;37.5MG	N77858 001	Sep 26, 2008	Apr	CAHN
----	-------	--------------	------------	--------------	-----	------

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

@ HOSPIRA EQ 500MG BASE/VIAL

N40108 001 Oct 30, 1995 Apr DISC

ACETIC ACID, GLACIAL

SOLUTION/DROPS; OTIC

VOSOL

>D>	@ HI TECH PHARMA	2%	N12179 001		Jul	CMFD
>A>	AT	2%	N12179 001		Jul	CMFD

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

@ BELCHER PHARMS	200MG	N74889 001	Oct 31, 1997	Jun	CAHN
------------------	-------	------------	--------------	-----	------

TABLET; ORAL

ACYCLOVIR

@ BELCHER PHARMS	400MG	N74891 001	Oct 31, 1997	Jun	CAHN
------------------	-------	------------	--------------	-----	------

@	800MG	N74891 002	Oct 31, 1997	Jun	CAHN
---	-------	------------	--------------	-----	------

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

@ HOSPIRA	EQ 500MG BASE/VIAL	N74758 001	Apr 22, 1997	Apr	DISC
-----------	--------------------	------------	--------------	-----	------

@	EQ 1GM BASE/VIAL	N74758 002	Apr 22, 1997	Apr	DISC
---	------------------	------------	--------------	-----	------

>A> ACYCLOVIR; HYDROCORTISONE

>A> CREAM; TOPICAL

>A> ACYCLOVIR AND HYDROCORTISONE

>A>	+ MEDIVIR	5%;1%	N22436 001	Jul 31, 2009	Jul	NEWA
-----	-----------	-------	------------	--------------	-----	------

ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

AP	LUITPOLD	3MG/ML	N90010 001	Apr 28, 2009	Apr	NEWA
----	----------	--------	------------	--------------	-----	------

AP	STRIDES ARCOLAB LTD	3MG/ML	N78686 001	May 13, 2009	May	NEWA
----	---------------------	--------	------------	--------------	-----	------

AP	WOCKHARDT	3MG/ML	N90220 001	Jul 20, 2009	Jun	NEWA
----	-----------	--------	------------	--------------	-----	------

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

@ ARMSTRONG PHARMS	0.09MG/INH	N72273 001	Aug 14, 1996	Jan	DISC
--------------------	------------	------------	--------------	-----	------

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

@ BAUSCH AND LOMB	EQ 0.083% BASE	N75358 001	Mar 29, 2000	Apr	DISC
-------------------	----------------	------------	--------------	-----	------

AN	HOLOPACK INTL	EQ 0.083% BASE	N77839 001	Dec 16, 2008	Jan	CAHN
----	---------------	----------------	------------	--------------	-----	------

AN	TEVA PARENTERAL	EQ 0.083% BASE	N75343 001	Nov 09, 1999	Apr	CAHN
----	-----------------	----------------	------------	--------------	-----	------

TABLET; ORAL

ALBUTEROL SULFATE

@ SANDOZ	EQ 2MG BASE	N72151 001	Dec 05, 1989	Mar	DISC
----------	-------------	------------	--------------	-----	------

@	EQ 4MG BASE	N72152 001	Dec 05, 1989	Mar	DISC
---	-------------	------------	--------------	-----	------

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE

AN	TEVA PARENTERAL	EQ 0.083% BASE;0.017%	N76724 001	Dec 31, 2007	Apr	CAHN
----	-----------------	-----------------------	------------	--------------	-----	------

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ALCLOMETASONE DIPROPIONATE

AB	GLENMARK GENERICS	0.05%	N79061 001	Jun 23, 2009	Jun	NEWA
----	-------------------	-------	------------	--------------	-----	------

OINTMENT; TOPICAL

ALCLOMETASONE DIPROPIONATE

>A>	AB	GLENMARK GENERICS	0.05%	N79227 001	Jul 30, 2009	Jul	NEWA
-----	----	-------------------	-------	------------	--------------	-----	------

ALENDRONATE SODIUM

TABLET; ORAL

ALENDRONATE SODIUM

AB	SANDOZ	EQ 5MG BASE	N75871 001	Apr 22, 2009	Apr	NEWA
AB		EQ 10MG BASE	N75871 002	Apr 22, 2009	Apr	NEWA
AB		EQ 35MG BASE	N75871 004	Apr 22, 2009	Apr	NEWA
AB		EQ 40MG BASE	N75871 003	Apr 22, 2009	Apr	NEWA
AB		EQ 70MG BASE	N75871 005	Apr 22, 2009	Apr	NEWA

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

@	SANDOZ	100MG	N70268 001	Dec 31, 1985	Mar	DISC
@		300MG	N70269 001	Dec 31, 1985	Mar	DISC

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

AB	+	PAR PHARM	5MG	N70346 001	Jan 22, 1986	Jan	CTEC
AB		SIGMAPHARM LABS LLC	5MG	N79133 001	Jan 30, 2009	Jan	NEWA
AB		MIDAMOR					
AB		PADDOCK LABS	5MG	N18200 001		May	CMFD

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

@	SANDOZ	EQ 5MG ANHYDROUS;50MG	N73357 001	Nov 27, 1991	Mar	DISC
---	--------	-----------------------	------------	--------------	-----	------

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMINOCAPROIC ACID

@	HOSPIRA	250MG/ML	N70888 001	Jun 16, 1988	Apr	DISC
---	---------	----------	------------	--------------	-----	------

TABLET; ORAL

AMICAR

	XANODYNE PHARM	1GM	N15197 002	Jun 24, 2004	Jan	NEWA
--	----------------	-----	------------	--------------	-----	------

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

AP	+	HOSPIRA	25MG/ML	N87242 001	Oct 26, 1983	Jun	CRLD
		AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%					
	@	HOSPIRA	100MG/100ML	N88147 002	May 03, 1983	Apr	DISC
	@		200MG/100ML	N88147 003	May 03, 1983	Apr	DISC

>D> SUPPOSITORY; RECTAL

>D> TRUPHYLLINE

>D>		G AND W LABS	250MG	N85498 001	Mar 23, 1983	Jul	DISC
-----	--	--------------	-------	------------	--------------	-----	------

>A>	@		250MG	N85498 001	Mar 23, 1983	Jul	DISC
-----	---	--	-------	------------	--------------	-----	------

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

@ INTL MEDICATION SYS 50MG/ML

N21594 001 Feb 04, 2004 Jun DISC

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

AB MYLAN 200MG

N75188 001 Feb 24, 1999 Apr CAHN

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

AB ALKEM EQ 2.5MG BASE

N78925 001 May 04, 2009 Apr NEWA

AB EQ 5MG BASE

N78925 002 May 04, 2009 Apr NEWA

AB EQ 10MG BASE

N78925 003 May 04, 2009 Apr NEWA

AB GLENMARK GENERICS EQ 2.5MG BASE

N78552 001 Apr 08, 2009 Mar NEWA

AB EQ 5MG BASE

N78552 002 Apr 08, 2009 Mar NEWA

AB EQ 10MG BASE

N78552 003 Apr 08, 2009 Mar NEWA

AB ORCHID HLTHCARE EQ 2.5MG BASE

N78453 001 Jul 02, 2009 Jun NEWA

AB EQ 5MG BASE

N78453 002 Jul 02, 2009 Jun NEWA

AB EQ 10MG BASE

N78453 003 Jul 02, 2009 Jun NEWA

AB SYNTHON PHARMS EQ 2.5MG BASE

N77080 001 Jun 27, 2007 Jan CAHN

AB EQ 5MG BASE

N77080 002 Jun 27, 2007 Jan CAHN

AB EQ 10MG BASE

N77080 003 Jun 27, 2007 Jan CAHN

AMLODIPINE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

EXFORGE HCT

NOVARTIS

5MG;12.5MG;160MG

N22314 001 Apr 30, 2009 Apr NEWA

5MG;25MG;160MG

N22314 002 Apr 30, 2009 Apr NEWA

10MG;12.5MG;160MG

N22314 003 Apr 30, 2009 Apr NEWA

+ 10MG;25MG;320MG

N22314 005 Apr 30, 2009 Apr NEWA

10MG;25MG;160MG

N22314 004 Apr 30, 2009 Apr NEWA

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

AB APOTEX 250MG;EQ 125MG BASE

N65333 001 Feb 24, 2009 Feb NEWA

AB 500MG;EQ 125MG BASE

N65333 002 Feb 24, 2009 Feb NEWA

APRACLOINIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

APRACLOINIDINE HYDROCHLORIDE

AT AKORN INC EQ 0.5% BASE

N77764 001 Mar 12, 2009 Feb NEWA

IOPIDINE

AT + ALCON EQ 0.5% BASE

N20258 001 Jul 30, 1993 Feb CFTG

ARMODAFINIL

TABLET; ORAL

NUVIGIL

CEPHALON

100MG

N21875 002 Mar 26, 2009 Mar CMFD

200MG

N21875 005 Mar 26, 2009 Mar NEWA

ARTEMETHER; LUMEFANTRINE

TABLET; ORAL

COARTEM

NOVARTIS

20MG;120MG

N22268 001 Apr 07, 2009 Apr NEWA

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

@ SANDOZ

325MG;50MG;40MG

N86398 002 Apr 06, 1984 Mar DISC

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENGESIC

@ SOLCO HLTHCARE

385MG;30MG;25MG

N75141 001 May 29, 1998 Jan CAHN

ORPHENGESIC FORTE

@ SOLCO HLTHCARE

770MG;60MG;50MG

N75141 002 May 29, 1998 Jan CAHN

ATENOLOL

TABLET; ORAL

ATENOLOL

AB IPCA LABS LTD

25MG

N77877 001 Dec 27, 2006 Jun CAHN

AB

50MG

N77877 002 Dec 27, 2006 Jun CAHN

AB

100MG

N77877 003 Dec 27, 2006 Jun CAHN

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

@ HOSPIRA

10MG/ML

N74740 001 Mar 28, 1997 Jan DISC

ATRACURIUM BESYLATE PRESERVATIVE FREE

@ HOSPIRA

10MG/ML

N74741 001 Mar 28, 1997 Jan DISC

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

+ VALEANT

0.025MG;1MG

N17744 002 Jun CMFD

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON-PLUS

>D> @ BIONICHE PHARMA

0.14MG/ML;10MG/ML

N19677 001 Nov 06, 1991 Jul CMFD

>A>

+

0.14MG/ML;10MG/ML

N19677 001 Nov 06, 1991 Jul CMFD

AZACITIDINE

INJECTABLE; IV-SC

VIDAZA

+ CELGENE

100MG/VIAL

N50794 001 May 19, 2004 Mar CAHN

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AZELASTINE HYDROCHLORIDE

>A>

APOTEX INC

0.05%

N78621 001 Aug 03, 2009 Jul NEWA

>A>

OPTIVAR

>D>

+

MEDA PHARMS

0.05%

N21127 001 May 22, 2000 Jul CFTG

>A>

AT

+

0.05%

N21127 001 May 22, 2000 Jul CFTG

SPRAY, METERED; NASAL
 AZELASTINE HYDROCHLORIDE
 @ APOTEX INC

EQ 0.125MG BASE/SPRAY

N77954 001 Apr 30, 2009 Apr DISC

AZITHROMYCIN

INJECTABLE; INJECTION

AZITHROMYCIN

AP HOSPIRA EQ 500MG BASE/VIAL

N65500 001 Jun 26, 2009 Jun NEWA

AP EQ 500MG BASE/VIAL

N65511 001 Jun 26, 2009 Jun NEWA

AP SAGENT STRIDES EQ 500MG BASE/VIAL

N65506 001 Mar 24, 2009 Mar NEWA

BACLOFEN

TABLET; ORAL

BACLOFEN

AB MYLAN 10MG

N77181 001 Jul 29, 2005 Apr CAHN

AB 20MG

N77121 002 Jul 29, 2005 Apr CAHN

BENDAMUSTINE HYDROCHLORIDE

POWDER; IV (INFUSION)

TREANDA

+ CEPHALON 25MG/VIAL

N22249 002 May 01, 2009 May NEWA

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

>D> BT + SANOFI AVENTIS US 5%;EQ 1% BASE

N50756 001 Dec 21, 2000 Jul CFTG

>A> AB + 5%;EQ 1% BASE

N50756 001 Dec 21, 2000 Jul CFTG

>A> CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE

>A> AB DOW PHARM SCIENCES 5%;EQ 1% BASE

N65443 001 Aug 11, 2009 Jul NEWA

BENZTROPINE MESYLATE

INJECTABLE; INJECTION

BENZTROPINE MESYLATE

>A> AP NEXUS PHARMS 1MG/ML

N90233 001 Jul 28, 2009 Jul NEWA

COGENTIN

>D> + LUNDBECK INC 1MG/ML

N12015 001 Jul CFTG

>A> AP + 1MG/ML

N12015 001 Jul CFTG

+ 1MG/ML

N12015 001 Apr CAHN

BENZYL ALCOHOL

LOTION; TOPICAL

BENZYL ALCOHOL

+ SCIELE PHARMA INC 5%

N22129 001 Apr 09, 2009 Apr NEWA

ULESFIA

+ SCIELE PHARMA INC 5%

N22129 001 Apr 09, 2009 May CTNA

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BESIVANCE

+ BAUSCH AND LOMB EQ 0.6% BASE

N22308 001 May 28, 2009 May NEWA

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE

>A> AB PHARMAFORCE 3MG/ML;EQ 3MG BASE/ML

N90747 001 Jul 31, 2009 Jul NEWA

INJECTABLE; INJECTION

CELESTONE SOLUSPAN

>D>	+	SCHERING	3MG/ML;EQ 3MG BASE/ML	N14602 001	Jul	CFTG
>A>	AB	+	3MG/ML;EQ 3MG BASE/ML	N14602 001	Jul	CFTG

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE

OINTMENT; TOPICAL

TACLONEX

+	LEO PHARM	0.064%;0.005%	N21852 001	Jan 09, 2006	Mar	CAHN
---	-----------	---------------	------------	--------------	-----	------

BETAMETHASONE VALERATE

AEROSOL, FOAM; TOPICAL

LUXIQ

+	CONNECTICS	EQ 0.12% BASE	N20934 001	Feb 28, 1999	May	CAHN
---	------------	---------------	------------	--------------	-----	------

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

AA	SUN PHARM INDS INC	5MG	N40897 001	Apr 22, 2009	Apr	NEWA
AA		10MG	N40897 002	Apr 22, 2009	Apr	NEWA
AA		25MG	N40897 003	Apr 22, 2009	Apr	NEWA
AA		50MG	N40897 004	Apr 22, 2009	Apr	NEWA

BICALUTAMIDE

TABLET; ORAL

BICALUTAMIDE

AB	ACCORD HLTHCARE INC	50MG	N78917 001	Jul 06, 2009	Jun	NEWA	
AB	KUDCO IRELAND	50MG	N77995 001	Jul 06, 2009	Jun	NEWA	
AB	MYLAN	50MG	N79185 001	Jul 06, 2009	Jun	NEWA	
AB	SANDOZ	50MG	N78575 001	Jul 06, 2009	Jun	NEWA	
AB	SUN PHARMA GLOBAL	50MG	N79110 001	Jul 06, 2009	Jun	NEWA	
AB	SYNTHON PHARMS	50MG	N77973 001	Jul 06, 2009	Jun	NEWA	
AB	TEVA	50MG	N76932 001	Jul 06, 2009	Jun	NEWA	
AB	ZYDUS PHARMS USA INC	50MG	N79089 001	Jul 06, 2009	Jun	NEWA	
	CASODEX						
AB	+	ASTRAZENECA	50MG	N20498 001	Oct 04, 1995	Jun	CFTG

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

BRIMONIDINE TARTRATE

AT	TEVA PARENTERAL	0.2%	N76372 001	Sep 10, 2004	Apr	CAHN
----	-----------------	------	------------	--------------	-----	------

BROMOCRIPTINE MESYLATE

TABLET; ORAL

CYCLOSET

+	VEROSCIENCE	EQ 0.8MG BASE	N20866 001	May 05, 2009	May	NEWA
---	-------------	---------------	------------	--------------	-----	------

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMFED-DM

+	MORTON GROVE	2MG/5ML;10MG/5ML;30MG/5ML	N88811 001	Jun 07, 1985	Jun	CTNA
	@ WOCKHARDT EU	2MG/5ML;10MG/5ML;30MG/5ML	N89681 001	Dec 22, 1988	Jun	DISC
AA		2MG/5ML;10MG/5ML;30MG/5ML	N89681 001	Dec 22, 1988	Apr	CAHN

BUDESONIDE

SUSPENSION; INHALATION

BUDESONIDE

AN	APOTEX	0.25MG/2ML	N78202 001	Mar 30, 2009	Mar	NEWA
AN		0.5MG/2ML	N78202 002	Mar 30, 2009	Mar	NEWA
AN	TEVA PARENTERAL	0.25MG/2ML	N77519 001	Nov 18, 2008	Apr	CAHN
AN		0.5MG/2ML	N77519 002	Nov 18, 2008	Apr	CAHN

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

SENSORCAINE

AP	APP PHARMS	0.25%	N70552 001	May 21, 1986	Jun	CAHN
AP		0.5%	N70553 001	May 21, 1986	Jun	CAHN
AP		0.75%	N70554 001	May 21, 1986	Jun	CAHN

INJECTABLE; SPINAL

SENSORCAINE

AP	APP PHARMS	0.75%	N71202 001	Apr 15, 1987	Jun	CAHN
----	------------	-------	------------	--------------	-----	------

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

SENSORCAINE

AP	APP PHARMS	0.25%;0.0091MG/ML	N70966 001	Oct 13, 1987	Jun	CAHN
AP		0.25%;0.0091MG/ML	N70967 001	Oct 13, 1987	Jun	CAHN
AP		0.5%;0.0091MG/ML	N70968 001	Oct 13, 1987	Jun	CAHN

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

AB1	WATSON LABS	100MG	N79095 001	Mar 24, 2009	Mar	NEWA
AB2		150MG	N79094 001	Mar 24, 2009	Mar	NEWA
AB1		150MG	N79095 002	Mar 24, 2009	Mar	NEWA
AB1		200MG	N79095 003	Mar 24, 2009	Mar	NEWA

WELLBUTRIN XL

>A>	AB3 +	BIOVAIL LABS INTL	150MG	N21515 001	Aug 28, 2003	Jul	CAHN
>A>	AB3		300MG	N21515 002	Aug 28, 2003	Jul	CAHN
>D>	AB3 +	SMITHKLINE BEECHAM	150MG	N21515 001	Aug 28, 2003	Jul	CAHN
>D>	AB3		300MG	N21515 002	Aug 28, 2003	Jul	CAHN

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPAR

AB	+	BRISTOL MYERS SQUIBB	15MG	N18731 003	Apr 22, 1996	Feb	CRLD
		@	30MG	N18731 004	Apr 22, 1996	Feb	DISC

BUSPIRONE HYDROCHLORIDE

AB	DR REDDYS LABS LTD	5MG	N78246 001	Feb 27, 2009	Feb	NEWA
AB		10MG	N78246 002	Feb 27, 2009	Feb	NEWA
AB		15MG	N78246 003	Feb 27, 2009	Feb	NEWA
AB		30MG	N78246 004	Feb 27, 2009	Feb	NEWA
AB	MYLAN	5MG	N75467 001	Feb 28, 2002	Apr	CAHN
AB		7.5MG	N75467 002	Mar 28, 2001	Apr	CAHN
AB		10MG	N75467 003	Feb 28, 2002	Apr	CAHN
AB		15MG	N75467 004	Feb 28, 2002	Apr	CAHN
	@ SANDOZ	5MG	N75413 001	Mar 19, 2002	Mar	DISC
	@	10MG	N75413 002	Mar 19, 2002	Mar	DISC
	@	15MG	N75413 003	Mar 19, 2002	Mar	DISC

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

AP	HIKMA FARMACEUTICA	1MG/ML	N78400 001	May 01, 2009	Apr	NEWA
AP		2MG/ML	N78247 001	Apr 29, 2009	Apr	NEWA
AP		2MG/ML	N78400 002	May 01, 2009	Apr	NEWA

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFFEINE CITRATE

AP	PADDOCK LABS	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N77233 001	Sep 21, 2006	Apr	CAHN
----	--------------	------------------------------------	------------	--------------	-----	------

CALCITONIN, SALMON

SPRAY, METERED; NASAL

CALCITONIN-SALMON

AB	MDRNA	200 IU/SPRAY	N76979 001	Jun 08, 2009	May	NEWA
----	-------	--------------	------------	--------------	-----	------

CALCITRIOL

CAPSULE; ORAL

ROCALTROL

AB	VALIDUS PHARMS	0.25UGM	N18044 001		Mar	CAHN
AB	+	0.5UGM	N18044 002		Mar	CAHN

OINTMENT; TOPICAL

VECTICAL

	+	GALDERMA LABS LP	3UGM/GM	N22087 001	Jan 23, 2009	Jan	NEWA
--	---	------------------	---------	------------	--------------	-----	------

SOLUTION; ORAL

ROCALTROL

AA	+	VALIDUS PHARMS	1UGM/ML	N21068 001	Nov 20, 1998	Mar	CAHN
----	---	----------------	---------	------------	--------------	-----	------

CAPECITABINE

TABLET; ORAL

XELODA

>D>	HLR	150MG	N20896 001	Apr 30, 1998	Jul	CAHN
>D>	+	500MG	N20896 002	Apr 30, 1998	Jul	CAHN
>A>	HOFFMANN LA ROCHE	150MG	N20896 001	Apr 30, 1998	Jul	CAHN
>A>	+	500MG	N20896 002	Apr 30, 1998	Jul	CAHN

CARBAMAZEPINE

TABLET; ORAL

CARBAMAZEPINE

	TORRENT PHARMS	100MG	N77272 001	Dec 07, 2005	Apr	CAHN
AB		200MG	N77272 002	Dec 07, 2005	Apr	CAHN
		300MG	N77272 003	Dec 07, 2005	Apr	CAHN
		400MG	N77272 004	Dec 07, 2005	Apr	CAHN

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

AB	TORRENT PHARMS	100MG	N75712 001	Jul 05, 2001	Jan	CAHN
----	----------------	-------	------------	--------------	-----	------

TABLET, EXTENDED RELEASE; ORAL

CARBAMAZEPINE

AB	TARO	100MG	N78115 001	Mar 31, 2009	Mar	NEWA
AB		200MG	N78115 002	Mar 31, 2009	Mar	NEWA
AB		400MG	N78115 003	Mar 31, 2009	Mar	NEWA

TEGRETOL -XR

AB	NOVARTIS	100MG	N20234 001	Mar 25, 1996	Mar	CFTG
----	----------	-------	------------	--------------	-----	------

TABLET, EXTENDED RELEASE; ORAL

TEGRETOL-XR

AB	NOVARTIS	200MG	N20234 002	Mar 25, 1996	Mar	CFTG
AB	+	400MG	N20234 003	Mar 25, 1996	Mar	CFTG

CARBENICILLIN INDANYL SODIUM

TABLET; ORAL

GEOCILLIN

@ PFIZER

EQ 382MG BASE

N50435 001

Mar DISC

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

@ SANDOZ

10MG;100MG

N73586 001 Jun 29, 1995 Mar DISC

@

25MG;100MG

N73587 001 Jun 29, 1995 Mar DISC

@

25MG;250MG

N73620 001 Jun 29, 1995 Mar DISC

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

@ KV PHARM

50MG;200MG

N76663 001 Jun 24, 2004 Apr DISC

>D>

TABLET, FOR SUSPENSION; ORAL

>D>

CARBILEV

>D>

RANBAXY

10MG;100MG

N76643 001 Jun 10, 2005 Jul DISC

>A>

@

10MG;100MG

N76643 001 Jun 10, 2005 Jul DISC

>D>

25MG;100MG

N76643 002 Jun 10, 2005 Jul DISC

>A>

@

25MG;100MG

N76643 002 Jun 10, 2005 Jul DISC

>D>

+

25MG;250MG

N76643 003 Jun 10, 2005 Jul DISC

>A>

@

25MG;250MG

N76643 003 Jun 10, 2005 Jul DISC

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

>A> AB

SUN PHARM INDS

10MG;100MG

N78690 001 Jul 31, 2009 Jul NEWA

>A> AB

25MG;100MG

N78690 002 Jul 31, 2009 Jul NEWA

>A> AB

25MG;250MG

N78690 003 Jul 31, 2009 Jul NEWA

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

AP + WATSON LABS

50MG/VIAL

N76162 001 Oct 14, 2004 Apr CAHN

AP +

150MG/VIAL

N76162 002 Oct 14, 2004 Apr CAHN

AP +

450MG/VIAL

N76162 003 Oct 14, 2004 Apr CAHN

INJECTABLE; IV (INFUSION)

CARBOPLATIN

>D>

@ ABRAXIS PHARM

EQ 50MG/5ML (10MG/ML)

N77247 001 Oct 21, 2004 Jul CPOT

>A>

@

50MG/5ML (10MG/ML)

N77247 001 Oct 21, 2004 Jul CPOT

>D> AP

EQ 50MG/5ML (10MG/ML)

N77266 001 Feb 15, 2006 Jul CPOT

>A> AP

50MG/5ML (10MG/ML)

N77266 001 Feb 15, 2006 Jul CPOT

>D>

@

EQ 150MG/15ML (10MG/ML)

N77247 002 Oct 21, 2004 Jul CPOT

>A>

@

150MG/15ML (10MG/ML)

N77247 002 Oct 21, 2004 Jul CPOT

>D> AP

EQ 150MG/15ML (10MG/ML)

N77266 002 Feb 15, 2006 Jul CPOT

>A> AP

150MG/15ML (10MG/ML)

N77266 002 Feb 15, 2006 Jul CPOT

>D> AP

EQ 450MG/45ML (10MG/ML)

N77247 003 Oct 21, 2004 Jul CPOT

>A> AP

450MG/45ML (10MG/ML)

N77247 003 Oct 21, 2004 Jul CPOT

>D> AP

EQ 450MG/45ML (10MG/ML)

N77266 003 Feb 15, 2006 Jul CPOT

>A> AP

450MG/45ML (10MG/ML)

N77266 003 Feb 15, 2006 Jul CPOT

>D> AP

EQ 600MG/60ML (10MG/ML)

N77266 004 Feb 15, 2006 Jul CPOT

>A> AP

600MG/60ML (10MG/ML)

N77266 004 Feb 15, 2006 Jul CPOT

>A> AP

AKORN

50MG/5ML (10MG/ML)

N90475 001 Jul 29, 2009 Jul NEWA

INJECTABLE; IV (INFUSION)

CARBOPLATIN

>A>	AP	AKORN	150MG/15ML (10MG/ML)	N90475 002	Jul 29, 2009	Jul	NEWA
>A>	AP		450MG/45ML (10MG/ML)	N90475 003	Jul 29, 2009	Jul	NEWA
>D>	AP	BEDFORD LABS	EQ 50MG/5ML (10MG/ML)	N77244 001	Oct 15, 2004	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77244 001	Oct 15, 2004	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77244 002	Oct 15, 2004	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77244 002	Oct 15, 2004	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77244 003	Oct 15, 2004	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77244 003	Oct 15, 2004	Jul	CPOT
>D>	AP		EQ 600MG/60ML (10MG/ML)	N77244 004	Jan 20, 2006	Jul	CPOT
>A>	AP		600MG/60ML (10MG/ML)	N77244 004	Jan 20, 2006	Jul	CPOT
>D>	AP	EBEWE PHARMA	EQ 50MG/5ML (10MG/ML)	N78280 001	May 08, 2008	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N78280 001	May 08, 2008	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N78280 002	May 08, 2008	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N78280 002	May 08, 2008	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N78280 003	May 08, 2008	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N78280 003	May 08, 2008	Jul	CPOT
>D>	AP	FRESENIUS KABI ONCOL	EQ 150MG/15ML (10MG/ML)	N77432 002	Sep 29, 2006	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77432 003	Sep 29, 2006	Jul	CPOT
>D>	AP		EQ 50MG/5ML (10MG/ML)	N77432 001	Sep 29, 2006	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77432 002	Sep 29, 2006	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77432 003	Sep 29, 2006	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77432 001	Sep 29, 2006	Jul	CPOT
>D>	AP	GENERAMEDIX	EQ 50MG/5ML (10MG/ML)	N77998 001	Apr 24, 2007	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77998 001	Apr 24, 2007	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77998 002	Apr 24, 2007	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77998 002	Apr 24, 2007	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77998 003	Apr 24, 2007	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77998 003	Apr 24, 2007	Jul	CPOT
>D>	AP	HOSPIRA	EQ 50MG/5ML (10MG/ML)	N76517 001	Oct 14, 2004	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N76517 001	Oct 14, 2004	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N76517 002	Oct 14, 2004	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N76517 002	Oct 14, 2004	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N76517 003	Oct 14, 2004	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N76517 003	Oct 14, 2004	Jul	CPOT
>D>	AP		EQ 600MG/60ML (10MG/ML)	N77059 001	Nov 23, 2004	Jul	CPOT
>A>	AP		600MG/60ML (10MG/ML)	N77059 001	Nov 23, 2004	Jul	CPOT
>D>	AP	PHARMACHEMIE	EQ 50MG/5ML (10MG/ML)	N77269 001	Oct 14, 2004	Jul	CPOT
>A>	AP	+	50MG/5ML (10MG/ML)	N77269 001	Oct 14, 2004	Jul	CPOT
>D>	AP	+	EQ 150MG/15ML (10MG/ML)	N77269 002	Oct 14, 2004	Jul	CPOT
>A>	AP	+	150MG/15ML (10MG/ML)	N77269 002	Oct 14, 2004	Jul	CPOT
>D>	AP	+	EQ 450MG/45ML (10MG/ML)	N77269 003	Oct 14, 2004	Jul	CPOT
>A>	AP	+	450MG/45ML (10MG/ML)	N77269 003	Oct 14, 2004	Jul	CPOT
>D>	AP	+	EQ 600MG/60ML (10MG/ML)	N77269 004	Dec 28, 2007	Jul	CPOT
>A>	AP	+	600MG/60ML (10MG/ML)	N77269 004	Dec 28, 2007	Jul	CPOT
>D>	AP	PHARMACHEMIE BV	EQ 50MG/5ML (10MG/ML)	N77679 001	Feb 25, 2009	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77679 001	Feb 25, 2009	Jul	CPOT
	AP		EQ 50MG/5ML (10MG/ML)	N77679 001	Feb 25, 2009	Feb	NEWA
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77679 002	Feb 25, 2009	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77679 002	Feb 25, 2009	Jul	CPOT
	AP		EQ 150MG/15ML (10MG/ML)	N77679 002	Feb 25, 2009	Feb	NEWA
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77679 003	Feb 25, 2009	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77679 003	Feb 25, 2009	Jul	CPOT
	AP		EQ 450MG/45ML (10MG/ML)	N77679 003	Feb 25, 2009	Feb	NEWA
>D>	AP	PLIVA LACHEMA	EQ 50MG/5ML (10MG/ML)	N78631 001	Dec 02, 2008	Jul	CPOT

INJECTABLE; IV (INFUSION)

CARBOPLATIN

>A>	AP	PLIVA LACHEMA	50MG/5ML (10MG/ML)	N78631 001	Dec 02, 2008	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N78631 002	Dec 02, 2008	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N78631 002	Dec 02, 2008	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N78631 003	Dec 02, 2008	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N78631 003	Dec 02, 2008	Jul	CPOT
>D>	AP		EQ 600MG/60ML (10MG/ML)	N78631 004	Dec 02, 2008	Jul	CPOT
>A>	AP		600MG/60ML (10MG/ML)	N78631 004	Dec 02, 2008	Jul	CPOT
>D>	AP	SPECTRUM PHARMS	EQ 50MG/5ML (10MG/ML)	N77096 001	Jun 14, 2005	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77096 001	Jun 14, 2005	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77096 002	Jun 14, 2005	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77096 002	Jun 14, 2005	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77096 003	Jun 14, 2005	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77096 003	Jun 14, 2005	Jul	CPOT
>D>	AP	SUN PHARM INDS	EQ 50MG/5ML (10MG/ML)	N77926 001	Sep 19, 2008	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77926 001	Sep 19, 2008	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77926 002	Sep 19, 2008	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77926 002	Sep 19, 2008	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77926 003	Sep 19, 2008	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77926 003	Sep 19, 2008	Jul	CPOT
>D>	AP	+ TEVA PARENTERAL	EQ 50MG/5ML (10MG/ML)	N77139 001	Sep 21, 2005	Jul	CPOT
>A>	AP	+	50MG/5ML (10MG/ML)	N77139 001	Sep 21, 2005	Jul	CPOT
>D>	AP		EQ 50MG/5ML (10MG/ML)	N77389 001	Mar 30, 2007	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77389 001	Mar 30, 2007	Jul	CPOT
>D>	AP	+	EQ 150MG/15ML (10MG/ML)	N77139 002	Sep 21, 2005	Jul	CPOT
>A>	AP	+	150MG/15ML (10MG/ML)	N77139 002	Sep 21, 2005	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77389 002	Mar 30, 2007	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77389 002	Mar 30, 2007	Jul	CPOT
>D>	AP	+	EQ 450MG/45ML (10MG/ML)	N77139 003	Sep 21, 2005	Jul	CPOT
>A>	AP	+	450MG/45ML (10MG/ML)	N77139 003	Sep 21, 2005	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77389 003	Mar 30, 2007	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77389 003	Mar 30, 2007	Jul	CPOT
>D>	AP	+	EQ 600MG/60ML (10MG/ML)	N77139 004	Sep 21, 2005	Jul	CPOT
>A>	AP	+	600MG/60ML (10MG/ML)	N77139 004	Sep 21, 2005	Jul	CPOT
>D>	AP	WATSON LABS	EQ 50MG/5ML (10MG/ML)	N77861 001	Jan 18, 2007	Jul	CPOT
>A>	AP		50MG/5ML (10MG/ML)	N77861 001	Jan 18, 2007	Jul	CPOT
>D>	AP		EQ 150MG/15ML (10MG/ML)	N77861 002	Jan 18, 2007	Jul	CPOT
>A>	AP		150MG/15ML (10MG/ML)	N77861 002	Jan 18, 2007	Jul	CPOT
>D>	AP		EQ 450MG/45ML (10MG/ML)	N77861 003	Jan 18, 2007	Jul	CPOT
>A>	AP		450MG/45ML (10MG/ML)	N77861 003	Jan 18, 2007	Jul	CPOT
>D>	AP		EQ 600MG/60ML (10MG/ML)	N77861 004	Jan 18, 2007	Jul	CPOT
>A>	AP		600MG/60ML (10MG/ML)	N77861 004	Jan 18, 2007	Jul	CPOT

PARAPLATIN

>D>	AP	+ BRISTOL MYERS SQUIBB	EQ 50MG/5ML (10MG/ML)	N20452 001	Jul 14, 2003	Jul	CPOT
>A>	AP	+	50MG/5ML (10MG/ML)	N20452 001	Jul 14, 2003	Jul	CPOT
>D>	AP	+	EQ 150MG/15ML (10MG/ML)	N20452 002	Jul 14, 2003	Jul	CPOT
>A>	AP	+	150MG/15ML (10MG/ML)	N20452 002	Jul 14, 2003	Jul	CPOT
>D>	AP	+	EQ 450MG/45ML (10MG/ML)	N20452 003	Jul 14, 2003	Jul	CPOT
>A>	AP	+	450MG/45ML (10MG/ML)	N20452 003	Jul 14, 2003	Jul	CPOT
>D>	AP	+	EQ 600MG/60ML (10MG/ML)	N20452 004	Jan 15, 2004	Jul	CPOT
>A>	AP	+	600MG/60ML (10MG/ML)	N20452 004	Jan 15, 2004	Jul	CPOT

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

>A>	AA	AUROBINDO PHARMA	350MG	N40792 001	Aug 06, 2009	Jul	NEWA
		@ SANDOZ	350MG	N81025 001	Apr 13, 1989	Mar	DISC

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF

@ GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

N50461 003

Jun DISC

@

EQ 10GM BASE/VIAL

N50461 005

Jun DISC

CEFAZOLIN SODIUM

AP CEPHAZONE PHARMA

EQ 500MG BASE/VIAL

N65280 001 Mar 18, 2009 Mar NEWA

AP

EQ 1GM BASE/VIAL

N65280 002 Mar 18, 2009 Mar NEWA

AP

EQ 10GM BASE/VIAL

N65295 001 Mar 18, 2009 Mar NEWA

AP

EQ 20GM BASE/VIAL

N65296 001 Mar 18, 2009 Mar NEWA

@ GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

N64033 001 Oct 31, 1993 Mar DISC

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

AP ACS DOBFAR

EQ 1GM BASE/VIAL

N65414 001 Jun 12, 2009 Jun NEWA

AP

EQ 2GM BASE/VIAL

N65414 002 Jun 12, 2009 Jun NEWA

CEFTAZIDIME

INJECTABLE; INJECTION

TAZICEF

>D> @ HOSPIRA

1GM/VIAL

N64032 001 Oct 31, 1993 Jul CMFD

>A> AP

1GM/VIAL

N64032 001 Oct 31, 1993 Jul CMFD

>D> @

2GM/VIAL

N64032 002 Oct 31, 1993 Jul CMFD

>A> AP

2GM/VIAL

N64032 002 Oct 31, 1993 Jul CMFD

CEFTRIAXONE SODIUM

INJECTABLE; IM-IV

CEFTRIAXONE

>D> AP SANDOZ

EQ 250MG BASE/VIAL

N65169 001 May 09, 2005 Jul CRLD

>A> AP +

EQ 250MG BASE/VIAL

N65169 001 May 09, 2005 Jul CRLD

>D> AP

EQ 500MG BASE/VIAL

N65169 002 May 09, 2005 Jul CRLD

>A> AP +

EQ 500MG BASE/VIAL

N65169 002 May 09, 2005 Jul CRLD

>D> AP

EQ 1GM BASE/VIAL

N65169 003 May 09, 2005 Jul CRLD

>A> AP +

EQ 1GM BASE/VIAL

N65169 003 May 09, 2005 Jul CRLD

AP +

EQ 2GM BASE/VIAL

N65169 004 May 09, 2005 Mar CRLD

ROCEPHIN

>D> @ HLR

EQ 250MG BASE/VIAL

N50585 001 Dec 21, 1984 Jul CAHN

>D> @

EQ 500MG BASE/VIAL

N50585 002 Dec 21, 1984 Jul CAHN

>D> @

EQ 1GM BASE/VIAL

N50585 003 Dec 21, 1984 Jul CAHN

>D> @

EQ 2GM BASE/VIAL

N50585 004 Dec 21, 1984 Jul CAHN

>A> @ HOFFMANN LA ROCHE

EQ 250MG BASE/VIAL

N50585 001 Dec 21, 1984 Jul CAHN

>A> @

EQ 500MG BASE/VIAL

N50585 002 Dec 21, 1984 Jul CAHN

>A> @

EQ 1GM BASE/VIAL

N50585 003 Dec 21, 1984 Jul CAHN

>A> @

EQ 2GM BASE/VIAL

N50585 004 Dec 21, 1984 Jul CAHN

INJECTABLE; INJECTION

ROCEPHIN

>D> @ HLR

EQ 10GM BASE/VIAL

N50585 005 Dec 21, 1984 Jul CAHN

>A> @ HOFFMANN LA ROCHE

EQ 10GM BASE/VIAL

N50585 005 Dec 21, 1984 Jul CAHN

INJECTABLE; INJECTION

	ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER								
>D>	@ HLR	EQ 10MG BASE/ML	N50624 001	Feb 11, 1987	Jul	CAHN			
>D>	@	EQ 20MG BASE/ML	N50624 002	Feb 11, 1987	Jul	CAHN			
>D>	@	EQ 40MG BASE/ML	N50624 003	Feb 11, 1987	Jul	CAHN			
>A>	@ HOFFMANN LA ROCHE	EQ 10MG BASE/ML	N50624 001	Feb 11, 1987	Jul	CAHN			
>A>	@	EQ 20MG BASE/ML	N50624 002	Feb 11, 1987	Jul	CAHN			
>A>	@	EQ 40MG BASE/ML	N50624 003	Feb 11, 1987	Jul	CAHN			

CEFTRIAZONE SODIUM; LIDOCAINEINJECTABLE; INJECTION

	ROCEPHIN KIT								
>D>	@ HLR	EQ 500MG BASE/VIAL,N/A;N/A,1%	N50585 007	May 08, 1996	Jul	CAHN			
>D>	@	EQ 1GM BASE/VIAL,N/A;N/A,1%	N50585 006	May 08, 1996	Jul	CAHN			
>A>	@ HOFFMANN LA ROCHE	EQ 500MG BASE/VIAL,N/A;N/A,1%	N50585 007	May 08, 1996	Jul	CAHN			
>A>	@	EQ 1GM BASE/VIAL,N/A;N/A,1%	N50585 006	May 08, 1996	Jul	CAHN			

CETIRIZINE HYDROCHLORIDESYRUP; ORAL

	CETIRIZINE HYDROCHLORIDE								
AA	DR REDDYS LABS LTD	5MG/5ML	N78870 001	Apr 27, 2009	Apr	NEWA			

CHLORHEXIDINE GLUCONATESOLUTION; DENTAL

	CHLORHEXIDINE GLUCONATE								
AT	XTTRIUM	0.12%	N77789 001	Jun 18, 2009	Jun	NEWA			

CHLOROTHIAZIDETABLET; ORALDIURIL

	@ LUNDBECK INC	250MG	N11145 004		Apr	CAHN			
	@	500MG	N11145 002		Apr	CAHN			

CHLOROTHIAZIDE SODIUMINJECTABLE; INJECTIONDIURIL

	+ LUNDBECK INC	EQ 500MG BASE/VIAL	N11145 005		Apr	CAHN			
--	----------------	--------------------	------------	--	-----	------	--	--	--

CHLORPROPAMIDETABLET; ORALCHLORPROPAMIDE

	@ SANDOZ	100MG	N88725 001	Aug 31, 1984	Mar	DISC			
	@	250MG	N88726 001	Aug 31, 1984	Mar	DISC			

CHLORZOXAZONETABLET; ORALCHLORZOXAZONE

	@ SANDOZ	250MG	N89852 001	May 04, 1988	Mar	DISC			
	@	500MG	N89853 001	May 04, 1988	Mar	DISC			
AA	WATSON LABS	500MG	N89859 001	May 04, 1988	Apr	CAHN			

CHOLINE FENOFIBRATECAPSULE, DELAYED RELEASE; ORALTRILIPIX

	ABBOTT LABS	EQ 45MG FENOFIBRIC ACID	N22224 001	Dec 15, 2008	Jan	CTNA			
	+	EQ 135MG FENOFIBRIC ACID	N22224 002	Dec 15, 2008	Jan	CTNA			

CHYMOPAPAIN

INJECTABLE; INJECTION

CHYMODIACTIN

@ CHART MEDCL

4,000 UNITS/VIAL

N18663 002 Aug 21, 1984 Mar CAHN

@

10,000 UNITS/VIAL

N18663 001 Nov 10, 1982 Mar CAHN

CIMETIDINE

TABLET; ORAL

CIMETIDINE

@ SANDOZ

200MG

N74100 001 Jan 31, 1995 Mar DISC

@

300MG

N74100 002 Jan 31, 1995 Mar DISC

@

400MG

N74100 003 Jan 31, 1995 Mar DISC

@

800MG

N74100 004 Jan 31, 1995 Mar DISC

CIPROFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS; OTIC

CETRAXAL

>D> + LABORATORIOS SALVAT EQ 0.2% BASE

N21918 001 May 01, 2009 Jul CAHN

+ EQ 0.2% BASE

N21918 001 May 01, 2009 May NEWA

>A> + WRASER PHARMS EQ 0.2% BASE

N21918 001 May 01, 2009 Jul CAHN

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

AB MYLAN EQ 250MG BASE

N75685 002 Jun 09, 2004 Mar CMFD

AB EQ 500MG BASE

N75685 003 Jun 09, 2004 Mar CMFD

AB EQ 750MG BASE

N75685 001 Jun 09, 2004 Mar CMFD

@ TEVA EQ 250MG BASE

N76136 001 Jun 09, 2004 Jan DISC

@ EQ 500MG BASE

N76136 002 Jun 09, 2004 Jan DISC

@ EQ 750MG BASE

N76136 003 Jun 09, 2004 Jan DISC

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

AB AMNEAL PHARMS EQ 10MG BASE

N77045 003 Apr 29, 2005 Feb CAHN

AB EQ 20MG BASE

N77045 002 Apr 29, 2005 Feb CAHN

AB EQ 40MG BASE

N77045 001 Apr 29, 2005 Feb CAHN

AB GLENMARK GENERICS EQ 10MG BASE

N77654 001 Feb 27, 2009 Feb NEWA

AB EQ 20MG BASE

N77654 002 Feb 27, 2009 Feb NEWA

AB EQ 40MG BASE

N77654 003 Feb 27, 2009 Feb NEWA

CLARITHROMYCIN

TABLET, EXTENDED RELEASE; ORAL

CLARITHROMYCIN

>D> + RANBAXY 1GM

N65210 001 Jan 26, 2005 Jul DISC

>A> @ 1GM

N65210 001 Jan 26, 2005 Jul DISC

CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLODERM

+ DOW PHARM SCIENCES 0.1%

N17765 001 Jun CAHN

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

>D> AB KALI LABS 0.5MG

N77147 001 May 02, 2005 Jul DISC

>A> @ 0.5MG

N77147 001 May 02, 2005 Jul DISC

TABLET; ORAL

CLONAZEPAM

>D>	AB	KALI LABS	1MG	N77147 002	May 02, 2005	Jul	DISC
>A>		@	1MG	N77147 002	May 02, 2005	Jul	DISC
>D>	AB		2MG	N77147 003	May 02, 2005	Jul	DISC
>A>		@	2MG	N77147 003	May 02, 2005	Jul	DISC

CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE

@ SANDOZ	0.1MG	N70887 001	Aug 31, 1988	Mar	DISC
@	0.2MG	N70886 001	Aug 31, 1988	Mar	DISC
@	0.3MG	N71294 001	Aug 31, 1988	Mar	DISC

CLOPIDOGREL BISULFATE

TABLET; ORAL

CLOPIDOGREL BISULFATE

@ DR REDDYS LABS INC	EQ 75MG BASE	N76273 001	Jan 14, 2008	Jun	DISC
----------------------	--------------	------------	--------------	-----	------

PLAVIX

SANOFI AVENTIS US	EQ 75MG BASE	N20839 001	Nov 17, 1997	Jun	CTEC
-------------------	--------------	------------	--------------	-----	------

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM

@ SANDOZ	3.75MG	N72219 001	Aug 26, 1988	Mar	DISC
----------	--------	------------	--------------	-----	------

TRANXENE

@ LUNDBECK INC	3.75MG	N17105 001		Apr	CAHN
----------------	--------	------------	--	-----	------

@	7.5MG	N17105 002		Apr	CAHN
---	-------	------------	--	-----	------

@	15MG	N17105 003		Apr	CAHN
---	------	------------	--	-----	------

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

@ SANDOZ	7.5MG	N72513 001	May 11, 1990	Mar	DISC
----------	-------	------------	--------------	-----	------

@	15MG	N72514 001	May 11, 1990	Mar	DISC
---	------	------------	--------------	-----	------

TRANXENE

AB	LUNDBECK INC	3.75MG	N17105 006		Apr	CAHN
----	--------------	--------	------------	--	-----	------

AB		7.5MG	N17105 007		Apr	CAHN
----	--	-------	------------	--	-----	------

AB	+	15MG	N17105 008		Apr	CAHN
----	---	------	------------	--	-----	------

TRANXENE SD

@ LUNDBECK INC	11.25MG	N17105 005		May	DISC
----------------	---------	------------	--	-----	------

	11.25MG	N17105 005		Apr	CAHN
--	---------	------------	--	-----	------

@	22.5MG	N17105 004		May	DISC
---	--------	------------	--	-----	------

+	22.5MG	N17105 004		Apr	CAHN
---	--------	------------	--	-----	------

CLOXACILLIN SODIUM

CAPSULE; ORAL

CLOXAPEN

@ GLAXOSMITHKLINE	EQ 250MG BASE	N61806 001		Jun	DISC
-------------------	---------------	------------	--	-----	------

@	EQ 500MG BASE	N61806 002		Jun	DISC
---	---------------	------------	--	-----	------

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

TRIACIN-C

+	STI PHARMA LLC	10MG/5ML;30MG/5ML;1.25MG/5ML	N88704 001	Mar 22, 1985	Apr	CAHN
---	----------------	------------------------------	------------	--------------	-----	------

CODEINE SULFATE

>A>	TABLET; ORAL					
>A>	CODEINE SULFATE					
>A>	ROXANE	15MG	N22402 001	Jul 16, 2009	Jul	NEWA
>A>		30MG	N22402 002	Jul 16, 2009	Jul	NEWA
>A>	+	60MG	N22402 003	Jul 16, 2009	Jul	NEWA

COLCHICINE

	TABLET; ORAL					
	COLCRYS					
>A>	+ AR HOLDING CO INC	0.6MG	N22352 001	Jul 29, 2009	Jul	CAHN

COLISTIMETHATE SODIUM

	INJECTABLE; INJECTION					
	COLISTIMETHATE SODIUM					
AP	PADDOCK	EQ 150MG BASE/VIAL	N65177 001	Mar 19, 2004	Apr	CTNA
AP	X GEN PHARMS	EQ 150MG BASE/VIAL	N64216 001	Feb 26, 1999	Apr	CTNA

CROMOLYN SODIUM

	SOLUTION; INHALATION					
	CROMOLYN SODIUM					
AN	+ TEVA PARENTERAL	10MG/ML	N75271 001	Jan 18, 2000	Apr	CAHN

CYANOCOBALAMIN

	GEL, METERED; NASAL					
	NASCOBAL					
	@ PAR PHARM	0.5MG/INH	N19722 001	Nov 05, 1996	Mar	CAHN
	SPRAY, METERED; NASAL					
	NASCOBAL					
	+ PAR PHARM CO	0.5MG/SPRAY	N21642 001	Jan 31, 2005	May	CAHN

CYCLOPHOSPHAMIDE

	INJECTABLE; INJECTION					
	CYTOXAN					
	@ BAXTER HLTHCARE	100MG/VIAL	N12142 001		Feb	CAHN
	@	200MG/VIAL	N12142 002		Feb	CAHN
	@	500MG/VIAL	N12142 003		Feb	CAHN
	@	1GM/VIAL	N12142 004	Aug 30, 1982	Feb	CAHN
	@	2GM/VIAL	N12142 005	Aug 30, 1982	Feb	CAHN
	LYOPHILIZED CYTOXAN					
	+ BAXTER HLTHCARE	100MG/VIAL	N12142 006	Dec 05, 1985	Feb	CAHN
	+	200MG/VIAL	N12142 007	Dec 10, 1985	Feb	CAHN
AP	+	500MG/VIAL	N12142 008	Jan 04, 1984	Feb	CAHN
AP	+	1GM/VIAL	N12142 010	Sep 24, 1985	Feb	CAHN
AP	+	2GM/VIAL	N12142 009	Dec 10, 1984	Feb	CAHN
	TABLET; ORAL					
	CYTOXAN					
	@ BAXTER HLTHCARE	25MG	N12141 002		Feb	CAHN
	@	50MG	N12141 001		Feb	CAHN

DACTINOMYCIN

	INJECTABLE; INJECTION					
	COSMEGEN					
	+ LUNDBECK INC	0.5MG/VIAL	N50682 001		Apr	CAHN

DARUNAVIR ETHANOLATE

TABLET; ORAL

PREZISTA

>A>	CENTOCOR ORTHO	EQ 75MG BASE	N21976 004	Dec 18, 2008	Jul	CAHN
>A>		EQ 150MG BASE	N21976 005	Dec 18, 2008	Jul	CAHN
>A>		EQ 300MG BASE	N21976 001	Jun 23, 2006	Jul	CAHN
>A>		EQ 400MG BASE	N21976 003	Oct 21, 2008	Jul	CAHN
>A>	+	EQ 600MG BASE	N21976 002	Feb 25, 2008	Jul	CAHN
>D>	TIBOTEC	EQ 75MG BASE	N21976 004	Dec 18, 2008	Jul	CAHN
>D>		EQ 150MG BASE	N21976 005	Dec 18, 2008	Jul	CAHN
		EQ 150MG BASE	N21976 005	Dec 18, 2008	Jun	CMFD
>D>		EQ 300MG BASE	N21976 001	Jun 23, 2006	Jul	CAHN
>D>		EQ 400MG BASE	N21976 003	Oct 21, 2008	Jul	CAHN
>D>	+	EQ 600MG BASE	N21976 002	Feb 25, 2008	Jul	CAHN

DAUNORUBICIN CITRATE

>D> INJECTABLE, LIPOSOMAL; INJECTION

>D> DAUNOXOME

>D>	+	DIATOS	EQ 2MG BASE/ML	N50704 002	Apr 08, 1996	Jul	DISC
>A>	@		EQ 2MG BASE/ML	N50704 002	Apr 08, 1996	Jul	DISC

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

AP	WATSON LABS	500MG/VIAL	N76806 001	Mar 31, 2006	Apr	CAHN
AP		2GM/VIAL	N76806 002	Mar 31, 2006	Apr	CAHN

DEGARELIX ACETATE

POWDER; SUBCUTANEOUS

FIRMAGON

	FERRING	EQ 80MG BASE/VIAL	N22201 001	Dec 24, 2008	Jun	CTNA
+		EQ 120MG BASE/VIAL	N22201 002	Dec 24, 2008	Jun	CTNA

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-21

DESOGESTREL AND ETHINYL ESTRADIOL

@	DURAMED PHARMS BARR	0.15MG;0.03MG	N75256 001	Aug 12, 1999	Feb	DISC
---	---------------------	---------------	------------	--------------	-----	------

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

PRISTIQ

>D>	WYETH PHARMS INC	EQ 50MG BASE	N21992 001	Feb 29, 2008	Jul	CRLD
>A>	+	EQ 50MG BASE	N21992 001	Feb 29, 2008	Jul	CRLD

DEXAMETHASONE

ELIXIR; ORAL

DEXAMETHASONE

AA	+	STI PHARMA LLC	0.5MG/5ML	N84754 001		Apr	CAHN
----	---	----------------	-----------	------------	--	-----	------

IMPLANT; INTRAVITREAL

OZURDEX

+	ALLERGAN	0.7MG	N22315 001	Jun 17, 2009	Jun	NEWA
---	----------	-------	------------	--------------	-----	------

DEXAMETHASONE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC

TOBRADEX ST

+	ALCON	0.05%;0.3%	N50818 001	Feb 13, 2009	Feb	NEWA
---	-------	------------	------------	--------------	-----	------

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE; ORAL

KAPIDEX

	TAKEDA PHARMS	30MG	N22287 001	Jan 30, 2009	Jan	NEWA
+		60MG	N22287 002	Jan 30, 2009	Jan	NEWA

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

AA	+	BARR	10MG	N40361 002	Jan 31, 2001	Mar	CRLD
		DEXTROSTAT					
		@ SHIRE	5MG	N84051 001		Mar	DISC
		@	10MG	N84051 002		Mar	DISC

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 60% IN PLASTIC CONTAINER

@	HOSPIRA	60GM/100ML	N19346 001	Jan 25, 1985	Apr	DISC
---	---------	------------	------------	--------------	-----	------

DIATRIZOATE MEGLUMINE

INJECTABLE; INJECTION

RENO-60

@	BRACCO	60%	N10040 016		Jun	DISC
---	--------	-----	------------	--	-----	------

RENO-DIP

@	BRACCO	30%	N10040 012		Jun	DISC
---	--------	-----	------------	--	-----	------

SOLUTION; URETERAL

RENO-30

@	BRACCO	30%	N10040 021		Jun	DISC
---	--------	-----	------------	--	-----	------

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

RENOGRAFIN-60

@	BRACCO	52%;8%	N10040 006		Jun	DISC
---	--------	--------	------------	--	-----	------

DIAZEPAM

TABLET; ORAL

DIAZEPAM

@	SANDOZ	2MG	N70302 001	Dec 20, 1985	Mar	DISC
---	--------	-----	------------	--------------	-----	------

@		5MG	N70303 001	Dec 20, 1985	Mar	DISC
---	--	-----	------------	--------------	-----	------

@		10MG	N70304 001	Dec 20, 1985	Mar	DISC
---	--	------	------------	--------------	-----	------

DICLOFENAC POTASSIUM

CAPSULE; ORAL

ZIPSOR

+	XANODYNE PHARM	25MG	N22202 001	Jun 18, 2009	Jun	NEWA
---	----------------	------	------------	--------------	-----	------

FOR SOLUTION; ORAL

CAMBIA

+	KOWA PHARMS	50MG	N22165 001	Jun 17, 2009	Jun	NEWA
---	-------------	------	------------	--------------	-----	------

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDE

AA	COREPHARMA	25MG	N40828 001	Nov 05, 2008	Feb	CTEC
----	------------	------	------------	--------------	-----	------

TENUATE

AA	+ WATSON PHARMS	25MG	N11722 002		Feb	CTEC
----	-----------------	------	------------	--	-----	------

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

@ SANDOZ 500MG

N74604 001 Jun 10, 1996 Mar DISC

+ TEVA 500MG

N73673 001 Jul 31, 1992 Mar CTEC

DIGOXIN

INJECTABLE; INJECTION

DIGOXIN

@ HOSPIRA 0.25MG/ML

N40206 001 Aug 28, 1998 Apr DISC

TABLET; ORAL

DIGOXIN

AB IMPAX LABS 0.125MG

N78556 001 Jul 20, 2009 Jun NEWA

AB 0.25MG

N78556 002 Jul 20, 2009 Jun NEWA

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILT-CD

AB3 APOTEX 300MG

N76151 004 May 20, 2004 Apr CAHN

DILTIAZEM HYDROCHLORIDE

>D> + MYLAN 120MG

N74910 003 May 02, 1997 Jul CTEC

>A> BC + 120MG

N74910 003 May 02, 1997 Jul CTEC

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

+ BARR 50MG

N80738 001 Mar CRLD

@ LNK 25MG

N87977 001 Jan 27, 1983 Mar DISC

@ 50MG

N87978 001 Jan 27, 1983 Mar DISC

@ SANDOZ 25MG

N80832 001 Mar DISC

@ 50MG

N80832 002 Mar DISC

@ VALEANT PHARM INTL 50MG

N80592 001 Mar DISC

@ WATSON LABS 25MG

N80728 001 Mar DISC

@ 50MG

N80727 001 Mar DISC

DIPYRIDAMOLE

TABLET; ORAL

DIPYRIDAMOLE

@ SANDOZ 25MG

N86944 002 Apr 16, 1991 Mar DISC

@ 50MG

N87562 001 Feb 25, 1992 Mar DISC

@ 75MG

N87561 001 Feb 25, 1992 Mar DISC

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS; ORAL

DEPAKOTE

AB + ABBOTT EQ 125MG VALPROIC ACID

N19680 001 Sep 12, 1989 Jan CFTG

DIVALPROEX SODIUM

AB DR REDDYS LABS LTD EQ 125MG VALPROIC ACID

N78979 001 Jan 23, 2009 Jan NEWA

CAPSULE, DELAYED REL PELLETS; ORAL

DIVALPROEX SODIUM

AB	ZYDUS PHARMS USA INC	EQ 125MG VALPROIC ACID	N78919 001	Jan 27, 2009	Jan	NEWA
----	----------------------	------------------------	------------	--------------	-----	------

TABLET, DELAYED RELEASE; ORAL

DIVALPROEX SODIUM

AB	MYLAN	EQ 125MG VALPROIC ACID	N90062 001	Mar 17, 2009	Mar	NEWA
AB		EQ 250MG VALPROIC ACID	N90062 002	Mar 17, 2009	Mar	NEWA
AB		EQ 500MG VALPROIC ACID	N90062 003	Mar 17, 2009	Mar	NEWA
AB	ZYDUS PHARMS USA INC	EQ 125MG VALPROIC ACID	N77100 001	Mar 05, 2009	Feb	NEWA
AB		EQ 250MG VALPROIC ACID	N77100 002	Mar 05, 2009	Feb	NEWA
AB		EQ 500MG VALPROIC ACID	N77100 003	Mar 05, 2009	Feb	NEWA

TABLET, EXTENDED RELEASE; ORAL

DEPAKOTE ER

AB	ABBOTT	EQ 250MG VALPROIC ACID	N21168 002	May 31, 2002	Jan	CFTG
AB	+	EQ 500MG VALPROIC ACID	N21168 001	Aug 04, 2000	Jan	CFTG

DIVALPROEX SODIUM

AB	ANCHEN PHARMS	EQ 250MG VALPROIC ACID	N78445 001	Feb 26, 2009	Feb	NEWA	
>A>	AB	EQ 500MG VALPROIC ACID	N78445 002	Aug 04, 2009	Jul	NEWA	
AB	IMPAX LABS	EQ 250MG VALPROIC ACID	N78791 001	May 06, 2009	Apr	NEWA	
>A>	AB	EQ 500MG VALPROIC ACID	N78791 002	Aug 04, 2009	Jul	NEWA	
AB	MYLAN	EQ 250MG VALPROIC ACID	N77567 001	Jan 29, 2009	Jan	NEWA	
AB		EQ 500MG VALPROIC ACID	N77567 002	Jan 29, 2009	Jan	NEWA	
>A>	AB	TEVA PHARMS	EQ 500MG VALPROIC ACID	N78700 001	Aug 03, 2009	Jul	NEWA
AB	WOCKHARDT	EQ 250MG VALPROIC ACID	N78705 002	Feb 10, 2009	Jan	NEWA	
>A>	AB	EQ 500MG VALPROIC ACID	N78705 001	Aug 04, 2009	Jul	NEWA	
AB	ZYDUS PHARMS USA INC	EQ 250MG VALPROIC ACID	N78239 001	Feb 27, 2009	Feb	NEWA	
>A>	AB	EQ 500MG VALPROIC ACID	N78239 002	Aug 04, 2009	Jul	NEWA	

DORIPENEM

INJECTABLE; IV (INFUSION)

DORIBAX

+	ORTHO MCNEIL JANSSEN	500MG/VIAL	N22106 001	Oct 12, 2007	Jun	CAHN
---	----------------------	------------	------------	--------------	-----	------

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

AT	ALCON	EQ 2% BASE	N78981 001	Apr 13, 2009	Mar	NEWA
AT	BAUSCH AND LOMB	EQ 2% BASE	N90143 001	Jun 25, 2009	Jun	NEWA

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE

AT	BAUSCH AND LOMB	EQ 2% BASE;EQ 0.5% BASE	N90037 001	Jul 14, 2009	Jun	NEWA
----	-----------------	-------------------------	------------	--------------	-----	------

DOXERCALCIFEROL

CAPSULE; ORAL

HECTOROL

>A>	GENZYME	1UGM	N20862 003	Jul 13, 2009	Jul	NEWA
-----	---------	------	------------	--------------	-----	------

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

+	PAR PHARM	EQ 150MG BASE	N65055 003	Jul 15, 2005	Jan	CRLD
---	-----------	---------------	------------	--------------	-----	------

TABLET; ORAL

DOXYCYCLINE

AB	MUTUAL PHARM	EQ 50MG BASE	N65471 001	Apr 17, 2009	Mar	NEWA
----	--------------	--------------	------------	--------------	-----	------

TABLET; ORAL

DOXYCYCLINE

AB	MUTUAL PHARM	EQ 75MG BASE	N65471 002	Apr 17, 2009	Mar	NEWA
AB		EQ 100MG BASE	N65471 003	Apr 17, 2009	Mar	NEWA

DRONABINOL

CAPSULE; ORAL

DRONABINOL

AB	SVC PHARMA	2.5MG	N78292 001	Jun 27, 2008	Mar	CAHN
AB		5MG	N78292 002	Jun 27, 2008	Mar	CAHN
AB		10MG	N78292 003	Jun 27, 2008	Mar	CAHN

>A> DRONEDARONE HYDROCHLORIDE

>A> TABLET; ORAL

>A> MULTAQ

>A>	+	SANOFI AVENTIS US	EQ 400MG BASE	N22425 001	Jul 01, 2009	Jul	NEWA
-----	---	-------------------	---------------	------------	--------------	-----	------

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL

DROSPIRENONE AND ETHINYL ESTRADIOL

AB	BARR	3MG;0.02MG	N78515 001	Mar 30, 2009	Mar	NEWA
----	------	------------	------------	--------------	-----	------

YAZ

AB	+	BAYER HLTHCARE	3MG;0.02MG	N21676 001	Mar 16, 2006	Mar	CFTG
----	---	----------------	------------	------------	--------------	-----	------

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON

AP	BIONICHE PHARMA	10MG/ML	N88873 001	Aug 06, 1985	Jun	CMFD
----	-----------------	---------	------------	--------------	-----	------

EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

@	BRISTOL MYERS SQUIBB	100MG	N20972 002	Sep 17, 1998	Jun	DISC
---	----------------------	-------	------------	--------------	-----	------

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

@	SANDOZ	2.5MG	N75048 001	Aug 22, 2000	Mar	DISC
---	--------	-------	------------	--------------	-----	------

@		5MG	N75048 002	Aug 22, 2000	Mar	DISC
---	--	-----	------------	--------------	-----	------

@		10MG	N75048 003	Aug 22, 2000	Mar	DISC
---	--	------	------------	--------------	-----	------

@		20MG	N75048 004	Aug 22, 2000	Mar	DISC
---	--	------	------------	--------------	-----	------

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

LEXXEL

@	ASTRAZENECA	5MG;2.5MG	N20668 002	Oct 28, 1998	May	DISC
---	-------------	-----------	------------	--------------	-----	------

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

AP	+	HOSPIRA	1.25MG/ML	N75458 001	Aug 22, 2000	Feb	CRLD
----	---	---------	-----------	------------	--------------	-----	------

VASOTEC

@	BIOVAIL LABS INTL	1.25MG/ML	N19309 001	Feb 09, 1988	Feb	DISC
---	-------------------	-----------	------------	--------------	-----	------

EPINEPHRINE

INJECTABLE; IM-SC

TWINJECT 0.15

+ SCIELE PHARMA INC EQ 0.15MG /DELIVERY N20800 002 May 28, 2004 Jun CAHN

TWINJECT 0.3

+ SCIELE PHARMA INC EQ 0.3MG /DELIVERY N20800 001 May 30, 2003 Jun CAHN

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE DENTAL

+ DENTSPLY PHARM 0.005MG/ML;4% N21383 001 Apr CTNA

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE DENTAL WITH EPINEPHRINE

AP + DENTSPLY PHARM 0.01MG/ML;2% N21381 001 Apr CTNA

AP + 0.02MG/ML;2% N21381 002 Apr CTNA

XYLOCAINE W/ EPINEPHRINE

AP + APP PHARMS 0.02MG/ML;2% N06488 005 Apr CMFD

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

EPIRUBICIN HYDROCHLORIDE

AP AKORN INC 50MG/25ML (2MG/ML) N90163 001 Jun 24, 2009 Jun NEWA

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

EPOPROSTENOL SODIUM

+ ACTELION EQ 1.5MG BASE/VIAL N22260 001 Jun 27, 2008 Apr CAHN

ERGOCALCIFEROL

CAPSULE; ORAL

ERGOCALCIFEROL

AA ORIT LABS LLC 50,000 IU N40833 001 May 20, 2009 May NEWA

ERYTHROMYCIN

OINTMENT; OPHTHALMIC

ERYTHROMYCIN

>D> AT AKORN 0.5% N64030 001 Jul 18, 1996 Jul DISC

>A> @ 0.5% N64030 001 Jul 18, 1996 Jul DISC

OINTMENT; TOPICAL

AKNE-MYCIN

+ DOW PHARM SCIENCES 2% N50584 001 Jan 10, 1985 Jun CAHN

SOLUTION; TOPICAL

SANSAC

>D> @ CORIA 2% N62522 001 Jan 24, 1985 Jul CAHN

>A> @ DOW PHARM SCIENCES 2% N62522 001 Jan 24, 1985 Jul CAHN

ERYTHROMYCIN ESTOLATE

SUSPENSION; ORAL

ERYTHROMYCIN ESTOLATE

@ ALPHARMA US PHARMS EQ 125MG BASE/5ML

N62353 001 Nov 18, 1982 Apr DISC

@ EQ 250MG BASE/5ML

N62409 001 Dec 16, 1982 Apr DISC

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

@ ALPHARMA US PHARMS EQ 200MG BASE/5ML

N62200 001

Apr DISC

@ EQ 400MG BASE/5ML

N62200 002

Apr DISC

ERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYL

GRANULE; ORAL

ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL

+ BARR EQ 200MG BASE/5ML;EQ 600MG
BASE/5ML

N62759 001 May 20, 1988 Jun CMFD

ERYZOLE

@ ALRA EQ 200MG BASE/5ML;EQ 600MG
BASE/5ML

N62758 001 Jun 15, 1988 Jan DISC

PEDIAZOLE

@ ROSS LABS EQ 200MG BASE/5ML;EQ 600MG
BASE/5ML

N50529 001

Jan DISC

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

@ HOSPIRA EQ 1GM BASE/VIAL

N50182 003

Feb DISC

AP + EQ 1GM BASE/VIAL

N62638 002 Oct 31, 1986 Feb CRDL

ESCITALOPRAM OXALATE

TABLET; ORAL

LEXAPRO

>D> AB FOREST LABS EQ 5MG BASE

N21323 001 Aug 14, 2002 Jul CTEC

>A> EQ 5MG BASE

N21323 001 Aug 14, 2002 Jul CTEC

>D> AB EQ 10MG BASE

N21323 002 Aug 14, 2002 Jul CTEC

>A> EQ 10MG BASE

N21323 002 Aug 14, 2002 Jul CTEC

>D> AB + EQ 20MG BASE

N21323 003 Aug 14, 2002 Jul CTEC

>A> + EQ 20MG BASE

N21323 003 Aug 14, 2002 Jul CTEC

ESTRADIOL

TABLET; ORAL

INNOFEM

@ NOVO NORDISK INC 0.5MG

N40312 001 Nov 19, 1999 Apr DISC

@ 1MG

N40312 002 Nov 19, 1999 Apr DISC

@ 2MG

N40312 003 Nov 19, 1999 Apr DISC

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDE

AB LUPIN 100MG

N78939 001 Jun 17, 2009 Jun NEWA

AB 400MG

N78939 002 Jun 17, 2009 Jun NEWA

MYAMBUTOL

AB STI PHARMA LLC 100MG

N16320 001

Mar CAHN

@ 200MG

N16320 002

Mar CAHN

AB 400MG

N16320 003

Mar CAHN

@ 500MG

N16320 004

Mar CAHN

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-21

DEMULEN 1/35-21

@ GD SEARLE LLC 0.035MG;1MG

N18168 001

Apr DISC

TABLET; ORAL-21

DEMULEN 1/50-21

@ GD SEARLE LLC

0.05MG;1MG

N16927 001

Apr DISC

ZOVIA 1/35E-21

@ WATSON LABS

0.035MG;1MG

N72720 001

Dec 30, 1991 Apr DISC

ZOVIA 1/50E-21

@ WATSON LABS

0.05MG;1MG

N72722 001

Dec 30, 1991 Apr DISC

TABLET; ORAL-28

DEMULEN 1/35-28

@ GD SEARLE LLC

0.035MG;1MG

N18160 001

Apr DISC

DEMULEN 1/50-28

@ GD SEARLE LLC

0.05MG;1MG

N16936 001

Apr DISC

ZOVIA 1/50E-28

+ WATSON LABS

0.05MG;1MG

N72723 001

Dec 30, 1991 Apr CRLD

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-21

TRIPHASIL-21

@ AKRIMAX PHARMS

0.03MG,0.04MG,0.03MG;0.05MG,0.075
MG,0.125MG

N19192 001

Nov 01, 1984 Apr CAHN

TABLET; ORAL-28

TRIPHASIL-28

@ WYETH PHARMS INC

0.03MG,0.04MG,0.03MG;0.05MG,0.075
MG,0.125MG

N19190 001

Nov 01, 1984 Apr DISC

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)

WATSON LABS

0.035MG,0.035MG;0.5MG,1MG

N71041 001

Sep 24, 1991 Apr CRLD

TABLET; ORAL-28

NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

WATSON LABS

0.035MG,0.035MG;0.5MG,1MG

N71044 001

Apr 01, 1988 Apr CTEC

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-21

LOESTRIN 21 1.5/30

AB WARNER CHILCOTT

0.03MG;1.5MG

N17875 001

Apr CRLD

LOESTRIN 21 1/20

AB WARNER CHILCOTT

0.02MG;1MG

N17876 001

Apr CRLD

TABLET; ORAL-28

LOESTRIN FE 1/20

AB WARNER CHILCOTT

0.02MG;1MG

N17354 001

Apr CRLD

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

ORTHO TRI-CYCLEN LO

AB + ORTHO MCNEIL JANSSEN

0.025MG,0.025MG,0.025MG;0.18MG,0.
215MG,0.25MG

N21241 001

Aug 22, 2002 Jun CFTG

TRI LO SPRINTC

AB BARR

0.025MG,0.025MG,0.025MG;0.18MG,0.
215MG,0.25MG

N76784 001

Jun 29, 2009 Jun NEWA

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

OGESTREL 0.5/50-21

@ WATSON LABS

0.05MG;0.5MG

N75406 001

Dec 15, 1999 Apr DISC

TABLET; ORAL - 21

OVRAL

@ AKRIMAX PHARMS

0.05MG;0.5MG

N16672 001

Mar CAHN

ETHOTOIN

TABLET; ORAL

PEGANONE

+ LUNDBECK INC

250MG

N10841 001

Apr CAHN

@

500MG

N10841 003

Apr CAHN

ETOPOSIDE

CAPSULE; ORAL

ETOPOSIDE

+ GENPHARM

50MG

N75635 001 Sep 19, 2001 Feb CRLD

VEPESID

@ BRISTOL MYERS SQUIBB

50MG

N19557 001 Dec 30, 1986 Feb DISC

EVEROLIMUS

TABLET; ORAL

AFINITOR

NOVARTIS

5MG

N22334 001 Mar 30, 2009 Mar NEWA

+

10MG

N22334 002 Mar 30, 2009 Mar NEWA

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

AB ALEMBIC LTD

20MG

N78916 001 May 22, 2009 May NEWA

AB

40MG

N78916 002 May 22, 2009 May NEWA

@ SANDOZ

20MG

N75302 001 Apr 16, 2001 Mar DISC

@

40MG

N75302 002 Apr 16, 2001 Mar DISC

FEBUXOSTAT

TABLET; ORAL

ULORIC

TAKEDA PHARMS

40MG

N21856 001 Feb 13, 2009 Feb NEWA

+

80MG

N21856 002 Feb 13, 2009 Feb NEWA

FENOPROFEN CALCIUM

CAPSULE; ORAL

NALFON

>A>

+ PEDINOL

EQ 200MG BASE

N17604 003

Jul CTNA

@

EQ 300MG BASE

N17604 002

Feb DISC

>A>

EQ 400MG BASE

N17604 004 Jul 21, 2009 Jul NEWA

>D>

NALFON 200

>D>

+ PEDINOL

EQ 200MG BASE

N17604 003

Jul CTNA

+

EQ 200MG BASE

N17604 003

Feb CRLD

TABLET; ORAL

FENOPROFEN CALCIUM

@ SANDOZ

EQ 600MG BASE

N72396 001 Oct 17, 1988 Mar DISC

FENTANYL CITRATE

>A>

FILM; BUCCAL

>A>

ONSOLIS

>A>

BIODELIVERY SCI

EQ 0.2MG BASE

N22266 001 Jul 16, 2009 Jul NEWA

>A>

EQ 0.4MG BASE

N22266 002 Jul 16, 2009 Jul NEWA

>A>

EQ 0.6MG BASE

N22266 003 Jul 16, 2009 Jul NEWA

>A>	FILM; BUCCAL								
>A>	ONSOLIS								
>A>	BIODELIVERY SCI	EQ 0.8MG BASE	N22266 004	Jul 16, 2009	Jul	NEWA			
>A>		EQ 1.2MG BASE	N22266 005	Jul 16, 2009	Jul	NEWA			
	INJECTABLE; INJECTION								
	SUBLIMAZE PRESERVATIVE FREE								
AP	+ AKORN	EQ 0.05MG BASE/ML	N16619 001		May	CAHN			
	<u>FERUMOXYTOL</u>								
	SOLUTION; INTRAVENOUS								
	FERAHEME								
	+ AMAG PHARMS INC	EQ 510MG IRON/17ML (EQ 30MG IRON/ML)	N22180 001	Jun 30, 2009	Jun	NEWA			
	<u>FLECAINIDE ACETATE</u>								
	TABLET; ORAL								
	FLECAINIDE ACETATE								
AB	APOTEX INC	50MG	N79164 001	Jul 09, 2009	Jun	NEWA			
AB		100MG	N79164 002	Jul 09, 2009	Jun	NEWA			
AB		150MG	N79164 003	Jul 09, 2009	Jun	NEWA			
	@ SANDOZ	50MG	N76030 001	Oct 28, 2002	Mar	DISC			
	@	100MG	N76030 002	Oct 28, 2002	Mar	DISC			
	@	150MG	N76030 003	Oct 28, 2002	Mar	DISC			
	<u>FLUCONAZOLE</u>								
	INJECTABLE; INJECTION								
	FLUCANAZOLE								
>A>	AP ACS DOBFAR INFO SA	200MG/100ML (2MG/ML)	N79104 001	Jul 30, 2009	Jul	NEWA			
>A>	AP	400MG/200ML (2MG/ML)	N79104 002	Jul 30, 2009	Jul	NEWA			
	<u>FLUDARABINE PHOSPHATE</u>								
	INJECTABLE; INJECTION								
	FLUDARA								
AP	+ GENZYME	50MG/VIAL	N20038 001	Apr 18, 1991	Jun	CAHN			
	FLUDARABINE PHOSPHATE								
AP	ACTAVIS TOTOWA	50MG/VIAL	N78610 001	Feb 11, 2009	Feb	NEWA			
	TABLET; ORAL								
>D>	FLUDARABINE PHOSPHATE								
>D>	+ SANOFI AVENTIS US	10MG	N22273 001	Dec 18, 2008	Jul	CTNA			
	+	10MG	N22273 001	Dec 18, 2008	Jun	CAHN			
>A>	OFORTA								
>A>	+ SANOFI AVENTIS US	10MG	N22273 001	Dec 18, 2008	Jul	CTNA			
	<u>FLUMAZENIL</u>								
	INJECTABLE; INJECTION								
	FLUMAZENIL								
AP	HIKMA FARMACEUTICA	0.5MG/5ML (0.1MG/ML)	N78527 001	Mar 23, 2009	Mar	NEWA			
AP		1MG/10ML (0.1MG/ML)	N78527 002	Mar 23, 2009	Mar	NEWA			
	<u>FLUOCINOLONE ACETONIDE</u>								
	OIL; TOPICAL								
	DERMA-SMOOTH/FS								
	+ HILL DERMAC	0.01%	N19452 002	Nov 09, 2005	Feb	NEWA			
	OIL/DROPS; OTIC								
>A>	DERMOTIC								
>A>	+ HILL DERMAC	0.01%	N19452 003	Nov 09, 2005	Jul	CTNA			

OIL/DROPS; OTIC									
>D>	FLUOCINOLONE ACETONIDE								
>D>	+	HILL DERMAC	0.01%		N19452	003	Nov 09, 2005	Jul	CTNA
SHAMPOO; TOPICAL									
CAPEX									
	+	GALDERMA LABS LP	0.01%		N20001	001	Aug 27, 1990	Jun	CTNA
<u>FLUOROMETHOLONE ACETATE; TOBRAMYCIN</u>									
SUSPENSION/DROPS; OPHTHALMIC									
TOBRASONE									
	@	ALCON	0.1%;0.3%		N50628	001	Jul 21, 1989	Jun	DISC
<u>FLUOROURACIL</u>									
SOLUTION; TOPICAL									
FLUOROPLEX									
	@	ELORAC	1%		N16765	001		Feb	CAHN
<u>FLUOXETINE HYDROCHLORIDE</u>									
CAPSULE; ORAL									
FLUOXETINE HYDROCHLORIDE									
AB1		ALEMBIC LTD	EQ 10MG BASE		N90223	001	Mar 19, 2009	Mar	NEWA
AB1			EQ 20MG BASE		N90223	002	Mar 19, 2009	Mar	NEWA
AB			EQ 40MG BASE		N90223	003	Mar 19, 2009	Mar	NEWA
AB1		BEIJING DOUBLE CRANE	EQ 10MG BASE		N76165	001	Feb 01, 2002	Mar	CAHN
AB1			EQ 20MG BASE		N76165	002	Feb 01, 2002	Mar	CAHN
>A>	AB1	LANDELA PHARM	EQ 10MG BASE		N75464	001	Jan 30, 2002	Jul	CAHN
>A>	AB1		EQ 20MG BASE		N75464	002	Jan 30, 2002	Jul	CAHN
>D>	AB1	RXELITE	EQ 10MG BASE		N75464	001	Jan 30, 2002	Jul	CAHN
>D>	AB1		EQ 20MG BASE		N75464	002	Jan 30, 2002	Jul	CAHN
SOLUTION; ORAL									
FLUOXETINE HYDROCHLORIDE									
AA		AUROBINDO PHARM	EQ 20MG BASE/5ML		N79209	001	Mar 20, 2009	Mar	NEWA
PROZAC									
	@	LILLY	EQ 20MG BASE/5ML		N20101	001	Apr 24, 1991	Jun	DISC
TABLET; ORAL									
FLUOXETINE HYDROCHLORIDE									
AB		MYLAN	EQ 10MG BASE		N75755	001	Aug 02, 2001	Apr	CAHN
	+		EQ 20MG BASE		N75755	002	Aug 02, 2001	Apr	CAHN
<u>FLURAZEPAM HYDROCHLORIDE</u>									
CAPSULE; ORAL									
FLURAZEPAM HYDROCHLORIDE									
	@	SANDOZ	15MG		N71716	001	Jul 31, 1991	Mar	DISC
	@		30MG		N71717	001	Jul 31, 1991	Mar	DISC
<u>FLURBIPROFEN</u>									
TABLET; ORAL									
FLURBIPROFEN									
	@	SANDOZ	50MG		N74448	001	Jul 28, 1995	Mar	DISC
	@		100MG		N74448	002	Jul 28, 1995	Mar	DISC
<u>FLUVOXAMINE MALEATE</u>									
TABLET; ORAL									
FLUVOXAMINE MALEATE									
	@	IVAX PHARMS	25MG		N75898	001	Mar 12, 2001	Mar	DISC
	@		50MG		N75898	002	Mar 12, 2001	Mar	DISC

TABLET; ORAL

FLUVOXAMINE MALEATE

@ IVAX PHARMS

100MG

N75898 003 Mar 12, 2001 Mar DISC

FOLIC ACID

TABLET; ORAL

FOLIC ACID

AA INVAGEN PHARMS

1MG

N90035 001 Jun 09, 2009 Jun NEWA

FOMEPIZOLE

INJECTABLE; INJECTION

FOMEPIZOLE

AP GENERAMEDIX 1.5GM/1.5ML (1GM/ML)

N79033 001 Apr 07, 2009 Mar NEWA

FOSFOMYCIN TROMETHAMINE

FOR SUSPENSION; ORAL

MONUROL

+ ZAMBON SPA

EQ 3GM BASE/PACKET

N50717 001 Dec 19, 1996 Jun CAHN

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

@ SANDOZ

10MG

N76188 001 Oct 08, 2004 Mar DISC

@

20MG

N76188 002 Oct 08, 2004 Mar DISC

@

40MG

N76188 003 Oct 08, 2004 Mar DISC

AB + TEVA

40MG

N76139 003 Nov 25, 2003 Jun CRLD

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

AB AUROBINDO PHARMA

10MG;12.5MG

N79245 001 Jul 09, 2009 Jun NEWA

AB

20MG;12.5MG

N79245 002 Jul 09, 2009 Jun NEWA

AB INVAGEN PHARMS

10MG;12.5MG

N90228 001 Jul 09, 2009 Jun NEWA

AB

20MG;12.5MG

N90228 002 Jul 09, 2009 Jun NEWA

AB + TEVA

20MG;12.5MG

N76945 002 Jul 05, 2006 Jun CRLD

MONOPRIL-HCT

@ BRISTOL MYERS SQUIBB

10MG;12.5MG

N20286 002 Nov 30, 1994 Feb DISC

@

20MG;12.5MG

N20286 001 Nov 30, 1994 Feb DISC

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

>D> AP BAXTER HLTHCARE

10MG/ML

N71439 001 Sep 14, 1990 Jul DISC

>A>

@

10MG/ML

N71439 001 Sep 14, 1990 Jul DISC

TABLET; ORAL

FUROSEMIDE

AB IPCA LABS LTD

20MG

N78010 001 Sep 18, 2006 Jun CAHN

AB

40MG

N78010 002 Sep 18, 2006 Jun CAHN

AB

80MG

N78010 003 Sep 18, 2006 Jun CAHN

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

>D> AB SANDOZ

100MG

N75428 001 Jan 24, 2006 Jul DISC

>A>

@

100MG

N75428 001 Jan 24, 2006 Jul DISC

>D> AB

300MG

N75428 002 Jan 24, 2006 Jul DISC

CAPSULE; ORAL

GABAPENTIN

>A>		@ SANDOZ	300MG	N75428 002	Jan 24, 2006	Jul	DISC
>D>	AB		400MG	N75428 003	Jan 24, 2006	Jul	DISC
>A>		@	400MG	N75428 003	Jan 24, 2006	Jul	DISC

TABLET; ORAL

GABAPENTIN

>D>	AB	RANBAXY	600MG	N76605 001	Sep 14, 2005	Jul	DISC
>A>		@	600MG	N76605 001	Sep 14, 2005	Jul	DISC
>D>	AB		800MG	N76605 002	Sep 14, 2005	Jul	DISC
>A>		@	800MG	N76605 002	Sep 14, 2005	Jul	DISC
>D>	AB	SANDOZ	600MG	N76120 001	Jan 27, 2006	Jul	DISC
>A>		@	600MG	N76120 001	Jan 27, 2006	Jul	DISC
>D>	AB		800MG	N76120 002	Jan 27, 2006	Jul	DISC
>A>		@	800MG	N76120 002	Jan 27, 2006	Jul	DISC

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

AB	IMPAX LABS	EQ 8MG BASE	N78484 001	May 27, 2009	May	NEWA
AB		EQ 16MG BASE	N78484 002	May 27, 2009	May	NEWA
AB		EQ 24MG BASE	N78484 003	May 27, 2009	May	NEWA

SOLUTION; ORAL

GALANTAMINE HYDROBROMIDE

AA	ROXANE	4MG/ML	N78185 001	Jan 30, 2009	Jan	NEWA
AA	+ ORTHO MCNEIL JANSSEN	4MG/ML	N21224 001	Jun 22, 2001	Jan	CFTG

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

AB	BEIJING YABAO	EQ 4MG BASE	N77604 001	Feb 06, 2009	Apr	CAHN
AB		EQ 8MG BASE	N77604 002	Feb 06, 2009	Apr	CAHN
AB		EQ 12MG BASE	N77604 003	Feb 06, 2009	Apr	CAHN
AB	MYLAN	EQ 4MG BASE	N77590 001	May 29, 2009	May	NEWA
AB		EQ 4MG BASE	N77603 001	Aug 28, 2008	Apr	CAHN
AB		EQ 8MG BASE	N77590 002	May 29, 2009	May	NEWA
AB		EQ 8MG BASE	N77603 002	Aug 28, 2008	Apr	CAHN
AB		EQ 12MG BASE	N77590 003	May 29, 2009	May	NEWA
AB		EQ 12MG BASE	N77603 003	Aug 28, 2008	Apr	CAHN
AB	PAR PHARM	EQ 4MG BASE	N77604 001	Feb 06, 2009	Jan	NEWA
AB		EQ 8MG BASE	N77604 002	Feb 06, 2009	Jan	NEWA
AB		EQ 12MG BASE	N77604 003	Feb 06, 2009	Jan	NEWA
AB	ROXANE	EQ 4MG BASE	N77608 001	Feb 11, 2009	Jan	NEWA
AB		EQ 8MG BASE	N77608 002	Feb 11, 2009	Jan	NEWA
AB		EQ 12MG BASE	N77608 003	Feb 11, 2009	Jan	NEWA
AB	SANDOZ	EQ 4MG BASE	N77589 001	Jun 22, 2009	Jun	NEWA
AB		EQ 8MG BASE	N77589 002	Jun 22, 2009	Jun	NEWA
AB		EQ 12MG BASE	N77589 003	Jun 22, 2009	Jun	NEWA
AB	TEVA PHARMS	EQ 4MG BASE	N77587 001	Jul 09, 2009	Jun	NEWA
AB		EQ 8MG BASE	N77587 002	Jul 09, 2009	Jun	NEWA
AB		EQ 12MG BASE	N77587 003	Jul 09, 2009	Jun	NEWA

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE

	@ ROCHE PALO	EQ 500MG BASE/VIAL	N19661 001	Jun 23, 1989	Jun	DISC
--	--------------	--------------------	------------	--------------	-----	------

INJECTABLE; INJECTION

GANCICLOVIR SODIUM

	+ BEDFORD	EQ 500MG BASE/VIAL	N76222 001	Jul 16, 2003	Jun	CRLD
--	-----------	--------------------	------------	--------------	-----	------

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

>D>	RANBAXY	3MG	N77366 001	Oct 06, 2005	Jul	DISC
>A>	@	3MG	N77366 001	Oct 06, 2005	Jul	DISC
>D>		6MG	N77366 002	Oct 06, 2005	Jul	DISC
>A>	@	6MG	N77366 002	Oct 06, 2005	Jul	DISC

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

DUETACT

	+ TAKEDA GLOBAL	2MG;30MG	N21925 001	Jul 28, 2006	May	CRLD
		4MG;30MG	N21925 002	Jul 28, 2006	May	CRLD

GLYBURIDE

TABLET; ORAL

GLYBURIDE

AB	+ TEVA	5MG	N74388 003	Aug 29, 1995	Mar	CRLD
	MICRONASE					
	@ PHARMACIA AND UPJOHN	1.25MG	N17498 001	May 01, 1984	Mar	DISC
	@	2.5MG	N17498 002	May 01, 1984	Mar	DISC
	@	5MG	N17498 003	May 01, 1984	Mar	DISC

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

AB	+ BRISTOL MYERS SQUIBB	2.5MG;500MG	N21178 002	Jul 31, 2000	May	CRLD
AB		5MG;500MG	N21178 003	Jul 31, 2000	May	CRLD
	GLYBURIDE AND METFORMIN	HYDROCHLORIDE				
AB	DR REDDYS LABS INC	1.25MG;250MG	N79009 001	Jun 03, 2009	May	NEWA
AB		2.5MG;500MG	N79009 002	Jun 03, 2009	May	NEWA
AB		5MG;500MG	N79009 003	Jun 03, 2009	May	NEWA

GLYCOPYRROLATE

TABLET; ORAL

GLYCOPYRROLATE

AA	WEST WARD	1MG	N40836 001	Mar 05, 2009	Feb	NEWA
AA		2MG	N40836 002	Mar 05, 2009	Feb	NEWA

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

AP	SANDOZ	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	N78534 001	Apr 30, 2009	Apr	NEWA
AP		EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N78531 001	Apr 30, 2009	Apr	NEWA
AP		EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	N78531 002	Apr 30, 2009	Apr	NEWA

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

AB	DR REDDYS LABS LTD	EQ 1MG BASE	N78846 001	Feb 27, 2009	Feb	NEWA
AB	NATCO PHARMA	EQ 1MG BASE	N78969 001	Jun 22, 2009	Jun	NEWA

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

AB	MYLAN	10MG	N70278 002	Jul 16, 2009	Jun	NEWA
AB		20MG	N70278 003	Jul 16, 2009	Jun	NEWA

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALOPERIDOL DECANOATE

@ SANDOZ

EQ 50MG BASE/ML

N76463 001 Jun 24, 2005 Apr DISC

@

EQ 100MG BASE/ML

N76463 002 Jun 24, 2005 Apr DISC

HALOPERIDOL LACTATE

INJECTABLE; INJECTION

HALOPERIDOL

@ SANDOZ

EQ 5MG BASE/ML

N76464 001 Sep 29, 2004 Apr DISC

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

SUPPRELIN LA

+ ENDO PHARM

50MG

N22058 001 May 03, 2007 Apr CAHN

VANTAS

+ ENDO PHARM

50MG

N21732 001 Oct 12, 2004 Apr CAHN

+ ENDO PHARMS

50MG

N21732 001 Oct 12, 2004 Mar CAHN

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN

@ ENDO PHARMS

1.5MG/5ML;5MG/5ML

N05213 002 Jul 26, 1988 Feb DISC

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

AA + HI TECH PHARMA

1.5MG/5ML;5MG/5ML

N40613 001 Feb 08, 2008 Feb CRLD

TABLET; ORAL

HYCODAN

@ ENDO PHARMS

1.5MG;5MG

N05213 001 Jul 26, 1988 Feb DISC

TUSSIGON

AA + KING PHARMS

1.5MG;5MG

N88508 001 Jul 30, 1985 Feb CRLD

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

AP AKORN

20MG/ML

N40730 001 Apr 21, 2009 Mar NEWA

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

AA GLENMARK PHARMS LTD

10MG

N90527 001 May 27, 2009 May NEWA

AA

25MG

N90527 002 May 27, 2009 May NEWA

AA

50MG

N90527 003 May 27, 2009 May NEWA

AA

100MG

N90527 004 May 27, 2009 May NEWA

@ SANDOZ

10MG

N83241 001 Mar DISC

@

25MG

N83560 001 Mar DISC

@

50MG

N83561 001 Mar DISC

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

AB IPCA LABS LTD

12.5MG

N79237 001 Apr 02, 2009 Mar NEWA

SOLUTION; ORAL									
HYDROCHLOROTHIAZIDE									
	@ ROXANE	50MG/5ML		N88587 001	Jul 02, 1984	Mar	DISC		
TABLET; ORAL									
HYDROCHLOROTHIAZIDE									
	@ SANDOZ	25MG		N87565 001	Mar 09, 1982	Mar	DISC		
	@	50MG		N84912 001		Mar	DISC		
<u>HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE</u>									
TABLET; ORAL									
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE									
	@ SANDOZ	25MG;40MG		N71060 001	Aug 26, 1987	Mar	DISC		
	@	25MG;80MG		N71061 001	Aug 26, 1987	Mar	DISC		
<u>HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE</u>									
TABLET; ORAL									
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE									
AB	RANBAXY	12.5MG;EQ 10MG BASE		N78211 001	Mar 04, 2009	Feb	NEWA		
AB		12.5MG;EQ 20MG BASE		N78211 002	Mar 04, 2009	Feb	NEWA		
AB		25MG;EQ 20MG BASE		N78211 003	Mar 04, 2009	Feb	NEWA		
<u>HYDROCHLOROTHIAZIDE; SPIRONOLACTONE</u>									
TABLET; ORAL									
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE									
	@ SANDOZ	25MG;25MG		N86881 001		Mar	DISC		
<u>HYDROCORTISONE</u>									
LOTION; TOPICAL									
CETACORT									
	@ CORIA	0.5%		N80426 002		Jun	DISC		
	@	1%		N80426 001		Jun	DISC		
<u>HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE</u>									
SUSPENSION; OPHTHALMIC									
TERRA-CORTRIL									
	@ PFIZER	1.5%;EQ 5MG BASE/ML		N61016 001		Feb	DISC		
<u>HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE</u>									
AEROSOL, METERED; TOPICAL									
PROCTOFOAM HC									
BX	UCB INC	1%;1%		N86195 001		Apr	CAHN		
<u>HYDROCORTISONE ACETATE; UREA</u>									
CREAM; TOPICAL									
CARMOL HC									
AT	+ NYCOMED US	1%;10%		N80505 001		Apr	CAHN		
<u>HYDROMORPHONE HYDROCHLORIDE</u>									
INJECTABLE; INJECTION									
DILAUDID									
+	PURDUE PHARM PRODS	1MG/ML		N19034 003	Apr 30, 2009	May	CMFD		
		1MG/ML		N19034 003	Apr 30, 2009	Apr	NEWA		
+		2MG/ML		N19034 004	Apr 30, 2009	May	CRLD		
		2MG/ML		N19034 004	Apr 30, 2009	Apr	NEWA		
+		4MG/ML		N19034 005	Apr 30, 2009	May	CRLD		
		4MG/ML		N19034 005	Apr 30, 2009	Apr	NEWA		

TABLET; ORAL									
HYDROMORPHONE HYDROCHLORIDE									
AB	ROXANE	4MG	N74597 003	May 29, 2009	May	NEWA			
<u>HYDROXYUREA</u>									
CAPSULE; ORAL									
HYDROXYUREA									
AB	BARR	500MG	N75143 001	Oct 16, 1998	Feb	CMFD			
<u>HYDROXYZINE HYDROCHLORIDE</u>									
INJECTABLE; INJECTION									
HYDROXYZINE HYDROCHLORIDE									
AP	+ ABRAXIS PHARM	25MG/ML	N87329 001		Jun	CRLD			
AP	+	50MG/ML	N87329 002		Jun	CRLD			
VISTARIL									
	@ PFIZER	25MG/ML	N11111 001		Jun	DISC			
	@	50MG/ML	N11111 002		Jun	DISC			
TABLET; ORAL									
HYDROXYZINE HYDROCHLORIDE									
	@ SANDOZ	10MG	N87869 001	Dec 20, 1982	Mar	DISC			
	@	25MG	N87870 001	Dec 20, 1982	Mar	DISC			
	@	50MG	N87871 001	Dec 20, 1982	Mar	DISC			
<u>HYDROXYZINE PAMOATE</u>									
CAPSULE; ORAL									
HYDROXYZINE PAMOATE									
	@ SANDOZ	EQ 25MG HCL	N81127 001	Jun 28, 1991	Mar	DISC			
<u>IBANDRONATE SODIUM</u>									
TABLET; ORAL									
BONIVA									
>D>	+ ROCHE	EQ 2.5MG BASE	N21455 001	May 16, 2003	Jul	DISC			
>A>	@	EQ 2.5MG BASE	N21455 001	May 16, 2003	Jul	DISC			
<u>IBUPROFEN</u>									
SOLUTION; INTRAVENOUS									
CALDOLOR									
	CUMBERLAND PHARMS	400MG/4ML (100MG/ML)	N22348 001	Jun 11, 2009	Jun	NEWA			
	+	800MG/8ML (100MG/ML)	N22348 002	Jun 11, 2009	Jun	NEWA			
TABLET; ORAL									
IBUPROFEN									
AB	+ DR REDDYS LA	800MG	N75682 003	Nov 14, 2001	Feb	CRLD			
	@ SANDOZ	300MG	N70734 001	Jun 12, 1986	Mar	DISC			
	@	400MG	N70735 001	Jun 12, 1986	Mar	DISC			
	@	600MG	N70736 001	Jun 12, 1986	Mar	DISC			
	@	800MG	N72169 001	Dec 11, 1987	Mar	DISC			
AB	SHASUN USA	400MG	N78329 001	Feb 05, 2009	Jan	NEWA			
AB		600MG	N78329 002	Feb 05, 2009	Jan	NEWA			
AB		800MG	N78329 003	Feb 05, 2009	Jan	NEWA			
MOTRIN									
	@ MCNEIL CONSUMER	300MG	N17463 003		Apr	DISC			
	@	400MG	N17463 002		Feb	DISC			
	@	600MG	N17463 004		Feb	DISC			
	@	800MG	N17463 005	May 22, 1985	Feb	DISC			

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

NEOPROFEN

+	LUNDBECK INC	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N21903 001	Apr 13, 2006	Apr	CAHN
---	--------------	------------------------------------	------------	--------------	-----	------

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDARUBICIN HYDROCHLORIDE

>A>	AP	APP PHARMS	1MG/ML	N65440 001	Aug 04, 2009	Jul	NEWA
-----	----	------------	--------	------------	--------------	-----	------

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

@	BAXTER HLTHCARE	1GM/VIAL	N19763 001	Dec 30, 1988	Feb	CAHN
---	-----------------	----------	------------	--------------	-----	------

@		3GM/VIAL	N19763 002	Dec 30, 1988	Feb	CAHN
---	--	----------	------------	--------------	-----	------

IFOSFAMIDE; MESNA

INJECTABLE; INJECTION

IFEX/MESNEX KIT

+	BAXTER HLTHCARE	1GM/VIAL;100MG/ML	N19763 003	Oct 10, 1992	Feb	CAHN
---	-----------------	-------------------	------------	--------------	-----	------

+		3GM/VIAL;100MG/ML	N19763 004	Oct 10, 1992	Feb	CAHN
---	--	-------------------	------------	--------------	-----	------

ILOPERIDONE

TABLET; ORAL

FANAPT

	VANDA PHARMS INC	1MG	N22192 001	May 06, 2009	May	NEWA
--	------------------	-----	------------	--------------	-----	------

		2MG	N22192 002	May 06, 2009	May	NEWA
--	--	-----	------------	--------------	-----	------

		4MG	N22192 003	May 06, 2009	May	NEWA
--	--	-----	------------	--------------	-----	------

		6MG	N22192 004	May 06, 2009	May	NEWA
--	--	-----	------------	--------------	-----	------

		8MG	N22192 005	May 06, 2009	May	NEWA
--	--	-----	------------	--------------	-----	------

		10MG	N22192 006	May 06, 2009	May	NEWA
--	--	------	------------	--------------	-----	------

+		12MG	N22192 007	May 06, 2009	May	NEWA
---	--	------	------------	--------------	-----	------

INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

@	MERCK	EQ 333MG BASE	N20685 005	Dec 17, 1998	Jun	DISC
---	-------	---------------	------------	--------------	-----	------

INDOMETHACIN

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

AB	+	SANDOZ	75MG	N74464 001	May 28, 1998	Feb	CTEC
----	---	--------	------	------------	--------------	-----	------

INDOMETHACIN

AB		AVANTHI INC	75MG	N79175 001	Mar 06, 2009	Feb	NEWA
----	--	-------------	------	------------	--------------	-----	------

SUSPENSION; ORAL

INDOCIN

+	IROKO PHARMS	25MG/5ML	N18332 001	Oct 10, 1985	Mar	CAHN
---	--------------	----------	------------	--------------	-----	------

INDOMETHACIN SODIUM

INJECTABLE; INJECTION

INDOCIN

AP	+	LUNDBECK INC	EQ 1MG BASE/VIAL	N18878 001	Jan 30, 1985	Apr	CAHN
----	---	--------------	------------------	------------	--------------	-----	------

INSULIN GLULISINE RECOMBINANT

INJECTABLE; SUBCUTANEOUS

APIDRA SOLOSTAR

SANOFI AVENTIS US 300 UNITS/3ML

N21629 003 Feb 24, 2009 Feb NEWA

INSULIN RECOMBINANT HUMAN

POWDER; INHALATION

EXUBERA

@ PFIZER 1MG/INH

N21868 001 Jan 27, 2006 Jun DISC

@ 3MG/INH

N21868 002 Jan 27, 2006 Jun DISC

IPRATROPIUM BROMIDE

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

AN TEVA PARENTERAL 0.02%

N75313 001 Feb 07, 2000 Apr CAHN

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

AP PHARMAFORCE 40MG/2ML (20MG/ML)

N90016 001 Jan 28, 2009 Mar NEWA

AP 40MG/2ML (20MG/ML)

N90016 001 Jan 28, 2009 Jan NEWA

AP 100MG/5ML (20MG/ML)

N90016 002 Jan 28, 2009 Mar NEWA

AP 100MG/5ML (20MG/ML)

N90016 002 Jan 28, 2009 Jan NEWA

ISOSORBIDE DINITRATE

TABLET; SUBLINGUAL

ISOSORBIDE DINITRATE

@ SANDOZ 2.5MG

N86225 001 Feb 19, 1988 Mar DISC

@ 5MG

N86222 001 Feb 19, 1988 Mar DISC

ISOTRETINOIN

CAPSULE; ORAL

AC CUTANE

>D> AB HLR 10MG

N18662 002 May 07, 1982 Jul DISC

>D> AB + 20MG

N18662 004 Mar 28, 1983 Jul DISC

>D> AB + 40MG

N18662 003 May 07, 1982 Jul DISC

>A> HOFFMANN LA ROCHE 10MG

N18662 002 May 07, 1982 Jul DISC

>A> @ 20MG

N18662 004 Mar 28, 1983 Jul DISC

>A> @ 40MG

N18662 003 May 07, 1982 Jul DISC

KETOCONAZOLE

GEL; TOPICAL

XOLEGEL

+ STIEFEL LABS INC 2%

N21946 001 Jul 28, 2006 Jan CAHN

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

@ SANDOZ 50MG

N74024 001 Dec 29, 1995 Mar DISC

@ 75MG

N74024 002 Dec 29, 1995 Mar DISC

KETOROLAC TROMETHAMINE

>A> SOLUTION/DROPS; OPHTHALMIC

>A> ACUVAIL

>A> + ALLERGAN 0.45% N22427 001 Jul 22, 2009 Jul NEWA

LAMOTRIGINE

TABLET; ORAL

LAMOTRIGINE

AB	APOTEX INC	25MG	N78625 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N78625 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N78625 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N78625 004	Jan 27, 2009	Jan	NEWA
AB	AUROBINDO PHARMA	25MG	N78956 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N78956 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N78956 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N78956 004	Jan 27, 2009	Jan	NEWA
AB	CADISTA PHARMS	25MG	N79132 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N79132 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N79132 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N79132 004	Jan 27, 2009	Jan	NEWA
AB	DR REDDYS LABS LTD	25MG	N76708 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N76708 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N76708 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N76708 004	Jan 27, 2009	Jan	NEWA
AB	GENPHARM ULC	25MG	N77428 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N77428 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N77428 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N77428 004	Jan 27, 2009	Jan	NEWA
AB	MATRIX LABS LTD	25MG	N78443 001	Feb 11, 2009	Jan	NEWA
AB		100MG	N78443 002	Feb 11, 2009	Jan	NEWA
AB		150MG	N78443 003	Feb 11, 2009	Jan	NEWA
AB		200MG	N78443 004	Feb 11, 2009	Jan	NEWA
AB	MYLAN	25MG	N77420 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N77420 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N77420 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N77420 004	Jan 27, 2009	Jan	NEWA
AB	ROXANE	25MG	N77392 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N77392 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N77392 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N77392 004	Jan 27, 2009	Jan	NEWA
AB	SANDOZ	25MG	N78645 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N78645 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N78645 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N78645 004	Jan 27, 2009	Jan	NEWA
AB	TARO PHARM INDS	25MG	N78525 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N78525 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N78525 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N78525 004	Jan 27, 2009	Jan	NEWA
AB	TORRENT PHARMS	25MG	N78947 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N78947 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N78947 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N78947 004	Jan 27, 2009	Jan	NEWA
AB	UPSHER SMITH	25MG	N78310 001	Feb 04, 2009	Jan	NEWA
AB		100MG	N78310 002	Feb 04, 2009	Jan	NEWA
AB		150MG	N78310 003	Feb 04, 2009	Jan	NEWA

TABLET; ORAL

LAMOTRIGINE

AB	UPSHER SMITH	200MG	N78310 004	Feb 04, 2009	Jan	NEWA
AB	WOCKHARDT	25MG	N78982 001	Jan 27, 2009	Jan	NEWA
AB		100MG	N78982 002	Jan 27, 2009	Jan	NEWA
AB		150MG	N78982 003	Jan 27, 2009	Jan	NEWA
AB		200MG	N78982 004	Jan 27, 2009	Jan	NEWA
AB	ZYDUS PHARMS USA	25MG	N77633 001	Jan 27, 2009	Jan	NEWA
		50MG	N77633 002	Jan 27, 2009	Jan	NEWA
AB		100MG	N77633 003	Jan 27, 2009	Jan	NEWA
AB		150MG	N77633 004	Jan 27, 2009	Jan	NEWA
AB		200MG	N77633 005	Jan 27, 2009	Jan	NEWA
		250MG	N77633 006	Jan 27, 2009	Jan	NEWA

TABLET, CHEWABLE; ORAL

LAMOTRIGINE

AB	GLENMARK GENERICS	5MG	N79099 001	Feb 19, 2009	Feb	NEWA
AB		25MG	N79099 002	Feb 19, 2009	Feb	NEWA
AB	TARO	5MG	N79204 001	Feb 04, 2009	Jan	NEWA
AB		25MG	N79204 002	Feb 04, 2009	Jan	NEWA

TABLET, EXTENDED RELEASE; ORAL

LAMICTAL XR

	SMITHKLINE BEECHAM	25MG	N22115 001	May 29, 2009	May	NEWA
>D>		50MG	N22115 002	May 29, 2009	Jul	CRLD
>A>	+	50MG	N22115 002	May 29, 2009	Jul	CRLD
		50MG	N22115 002	May 29, 2009	May	NEWA
		100MG	N22115 003	May 29, 2009	May	NEWA
>D>	+	200MG	N22115 004	May 29, 2009	Jul	CRLD
>A>		200MG	N22115 004	May 29, 2009	Jul	CRLD
	+	200MG	N22115 004	May 29, 2009	May	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

LAMICTAL ODT

	SMITHKLINE BEECHAM	25MG	N22251 001	May 08, 2009	May	NEWA
>D>		50MG	N22251 002	May 08, 2009	Jul	CRLD
>A>	+	50MG	N22251 002	May 08, 2009	Jul	CRLD
		50MG	N22251 002	May 08, 2009	May	NEWA
		100MG	N22251 003	May 08, 2009	May	NEWA
>D>	+	200MG	N22251 004	May 08, 2009	Jul	CRLD
>A>		200MG	N22251 004	May 08, 2009	Jul	CRLD
	+	200MG	N22251 004	May 08, 2009	May	NEWA

LANSOPRAZOLE

FOR SUSPENSION, DELAYED RELEASE; ORAL

PREVACID

@	TAKEDA PHARMS NA	15MG/PACKET	N21281 001	May 03, 2001	May	DISC
@		30MG/PACKET	N21281 002	May 03, 2001	May	DISC

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

@	COREPHARMA	EQ 5MG BASE	N74544 001	Aug 28, 1997	Jun	CAHN
@		EQ 25MG BASE	N74544 002	Aug 28, 1997	Jun	CAHN
	ROXANE	EQ 10MG BASE	N72734 001	Feb 22, 1993	Jun	CTEC
@	XANODYNE PHARM	EQ 5MG BASE	N18459 001	Jan 30, 1986	May	DISC
@		EQ 10MG BASE	N71962 001	Nov 19, 1987	Jun	DISC

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION

VIADUR

@ ORTHO MCNEIL JANSSEN EQ 65MG BASE

N21088 001 Mar 03, 2000 Jun DISC

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

AP SUN PHARMA GLOBAL 1MG/0.2ML

N78885 001 Mar 09, 2009 Feb NEWA

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

AN DEY EQ 0.25% BASE

N78309 001 Mar 20, 2009 Mar NEWA

XOPENEX

AN + SEPRACOR EQ 0.25% BASE

N20837 004 Jul 18, 2003 Mar CFTG

LEVETIRACETAM

SOLUTION; ORAL

LEVETIRACETAM

AA SILARX 100MG/ML

N90263 001 Apr 03, 2009 Mar NEWA

AA TARO 100MG/ML

N78774 001 Feb 10, 2009 Jan NEWA

TABLET; ORAL

LEVETIRACETAM

AB APOTEX INC 250MG

N78869 001 Mar 13, 2009 Mar NEWA

AB 500MG

N78869 002 Mar 13, 2009 Mar NEWA

AB 750MG

N78869 003 Mar 13, 2009 Mar NEWA

AB 1GM

N78869 004 Mar 13, 2009 Mar NEWA

AB CIPLA LTD 250MG

N77319 001 Mar 20, 2009 Mar NEWA

AB 500MG

N77319 002 Mar 20, 2009 Mar NEWA

AB 750MG

N77319 003 Mar 20, 2009 Mar NEWA

AB GENPHARM ULC 250MG

N78731 001 Feb 10, 2009 Jan NEWA

AB 500MG

N78731 002 Feb 10, 2009 Jan NEWA

AB 750MG

N78731 003 Feb 10, 2009 Jan NEWA

AB 1GM

N78731 004 Feb 10, 2009 Jan NEWA

AB SOLCO HLTHCARE 250MG

N78106 001 Feb 10, 2009 Jan NEWA

AB 500MG

N78106 002 Feb 10, 2009 Jan NEWA

AB 750MG

N78106 003 Feb 10, 2009 Jan NEWA

AB 1GM

N78106 004 Feb 10, 2009 Jan NEWA

AB WATSON LABS FLORIDA 250MG

N77408 001 Mar 02, 2009 Feb NEWA

AB 500MG

N77408 002 Mar 02, 2009 Feb NEWA

AB 750MG

N77408 003 Mar 02, 2009 Feb NEWA

AB ZYDUS PHARMS USA INC 250MG

N78918 001 Apr 29, 2009 Apr NEWA

AB 1GM

N78918 002 Apr 29, 2009 Apr NEWA

TABLET, EXTENDED RELEASE; ORAL

KEPPRA XR

UCB INC 500MG

N22285 001 Sep 12, 2008 Feb CRLD

+ 750MG

N22285 002 Feb 12, 2009 Feb NEWA

LEVOFLOXACIN

TABLET; ORAL

LEVAQUIN

AB ORTHO MCNEIL JANSSEN 250MG

N20634 001 Dec 20, 1996 Jun CTEC

AB 250MG

N20634 001 Dec 20, 1996 Apr CFTG

AB 500MG

N20634 002 Dec 20, 1996 Jun CTEC

AB 500MG

N20634 002 Dec 20, 1996 Apr CFTG

+ 750MG

N20634 003 Sep 08, 2000 Jun CTEC

TABLET; ORAL

LEVAQUIN

AB	+	ORTHO MCNEIL JANSSEN	750MG	N20634 003	Sep 08, 2000	Apr	CFTG
----	---	----------------------	-------	------------	--------------	-----	------

LEVONORGESTREL

>D> IMPLANT; IMPLANTATION

>D> NORPLANT II

>D>	+	POPULATION COUNCIL	75MG/IMPLANT	N20544 001	Nov 01, 1996	Jul	DISC
-----	---	--------------------	--------------	------------	--------------	-----	------

>A>	@		75MG/IMPLANT	N20544 001	Nov 01, 1996	Jul	DISC
-----	---	--	--------------	------------	--------------	-----	------

TABLET; ORAL

LEVONORGESTREL

AB		WATSON LABS	0.75MG	N78666 001	Jun 24, 2009	Jun	NEWA
----	--	-------------	--------	------------	--------------	-----	------

>A> PLAN B ONE-STEP

>A>	+	DURAMED	1.5MG	N21998 001	Jul 10, 2009	Jul	NEWA
-----	---	---------	-------	------------	--------------	-----	------

LEVOTHYROXINE SODIUM**

**Refer to Annual Edition Preface Section 1.8 Levothyroxine Sodium for amplifying information

TABLET; ORAL

LEVOTHYROXINE SODIUM

AB2,		MERCK KGAA	0.025MG	N76752 001	Jun 16, 2005	Apr	CAHN
------	--	------------	---------	------------	--------------	-----	------

AB3			0.05MG	N76752 002	Jun 16, 2005	Apr	CAHN
-----	--	--	--------	------------	--------------	-----	------

AB2,			0.075MG	N76752 003	Jun 16, 2005	Apr	CAHN
------	--	--	---------	------------	--------------	-----	------

AB3			0.088MG	N76752 004	Jun 16, 2005	Apr	CAHN
-----	--	--	---------	------------	--------------	-----	------

AB2,			0.1MG	N76752 005	Jun 16, 2005	Apr	CAHN
------	--	--	-------	------------	--------------	-----	------

AB3			0.112MG	N76752 006	Jun 16, 2005	Apr	CAHN
-----	--	--	---------	------------	--------------	-----	------

AB2,			0.125MG	N76752 007	Jun 16, 2005	Apr	CAHN
------	--	--	---------	------------	--------------	-----	------

AB3			0.15MG	N76752 008	Jun 16, 2005	Apr	CAHN
-----	--	--	--------	------------	--------------	-----	------

AB2,			0.175MG	N76752 009	Jun 16, 2005	Apr	CAHN
------	--	--	---------	------------	--------------	-----	------

AB3			0.2MG	N76752 010	Jun 16, 2005	Apr	CAHN
-----	--	--	-------	------------	--------------	-----	------

AB2,			0.3MG	N76752 011	Jun 16, 2005	Apr	CAHN
------	--	--	-------	------------	--------------	-----	------

AB3

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

@	HOSPIRA	10%	N88367 001	Jul 31, 1984	Apr	DISC
---	---------	-----	------------	--------------	-----	------

XYLOCAINE

AP	+	APP PHARMS	2%	N06488 002		Apr	CMFD
----	---	------------	----	------------	--	-----	------

XYLOCAINE DENTAL

AP	+	DENTSPLY PHARM	2%	N21380 001		Apr	CTNA
----	---	----------------	----	------------	--	-----	------

SYSTEM; INTRADERMAL

ZINGO

@	ANESIVA	0.5MG	N22114 001	Aug 16, 2007	May	DISC
---	---------	-------	------------	--------------	-----	------

LINDANE

LOTION; TOPICAL

LINDANE

@	OLTA PHARMS	1%	N87313 001		Apr	CAHN
---	-------------	----	------------	--	-----	------

SHAMPOO; TOPICAL									
LINDANE									
AT	+	OLTA PHARMS	1%		N87266 001		Jan	CAHN	
<u>LIOTHYRONINE SODIUM</u>									
TABLET; ORAL									
CYTOMEL									
AB		KING PHARMS	EQ 0.005MG BASE		N10379 001		Mar	CFTG	
AB			EQ 0.025MG BASE		N10379 002		Mar	CTEC	
AB	+		EQ 0.05MG BASE		N10379 003		Mar	CTEC	
LIOTHYRONINE SODIUM									
AB		COASTAL PHARMS	EQ 0.005MG BASE		N90097 001	Mar 20, 2009	Mar	NEWA	
AB			EQ 0.025MG BASE		N90097 002	Mar 20, 2009	Mar	NEWA	
AB			EQ 0.05MG BASE		N90097 003	Mar 20, 2009	Mar	NEWA	
AB		MYLAN	EQ 0.005MG BASE		N90326 001	Jul 14, 2009	Jun	NEWA	
AB			EQ 0.025MG BASE		N90326 002	Jul 14, 2009	Jun	NEWA	
AB			EQ 0.05MG BASE		N90326 003	Jul 14, 2009	Jun	NEWA	
<u>LITHIUM CARBONATE</u>									
CAPSULE; ORAL									
LITHIUM CARBONATE									
AB		GLENMARK GENERICS	150MG		N79139 001	Feb 03, 2009	Jan	NEWA	
AB			300MG		N79139 002	Feb 03, 2009	Jan	NEWA	
AB			600MG		N79139 003	Feb 03, 2009	Jan	NEWA	
<u>LOPERAMIDE HYDROCHLORIDE</u>									
CAPSULE; ORAL									
LOPERAMIDE HYDROCHLORIDE									
		@ SANDOZ	2MG		N72993 001	Aug 28, 1992	Mar	DISC	
<u>LORAZEPAM</u>									
CONCENTRATE; ORAL									
LORAZEPAM									
AA		PADDOCK LABS	2MG/ML		N79244 001	Apr 28, 2009	Apr	NEWA	
LORAZEPAM INTENSOL									
AA	+	ROXANE	2MG/ML		N72755 001	Jun 28, 1991	Apr	CFTG	
SOLUTION; ORAL									
LORAZEPAM									
		@ ROXANE	0.5MG/5ML		N74648 001	Mar 18, 1997	Mar	DISC	
<u>MALATHION</u>									
LOTION; TOPICAL									
MALATHION									
AT		SYNERX PHARMA	0.5%		N78743 001	Mar 06, 2009	Feb	NEWA	
OVIDE									
AT	+	TARO PHARMS NORTH	0.5%		N18613 001	Aug 02, 1982	Feb	CFTG	
<u>MECHLORETHAMINE HYDROCHLORIDE</u>									
INJECTABLE; INJECTION									
MUSTARGEN									
	+	LUNDBECK INC	10MG/VIAL		N06695 001		Apr	CAHN	
<u>MECLIZINE HYDROCHLORIDE</u>									
TABLET; ORAL									
ANTIVERT									
AA	+	PFIZER	12.5MG		N10721 006		Jan	CAHN	

TABLET; ORAL									
ANTIVERT									
AA	+	PFIZER	25MG	N10721	004		Jan	CAHN	
AA	+		50MG	N10721	001	Jan 20, 1982	Jan	CAHN	
<u>MECLOFENAMATE SODIUM</u>									
CAPSULE; ORAL									
MECLOFENAMATE SODIUM									
		@ SANDOZ	EQ 50MG BASE	N72262	001	Nov 29, 1988	Mar	DISC	
		@	EQ 100MG BASE	N72263	001	Nov 29, 1988	Mar	DISC	
<u>MEDROXYPROGESTERONE ACETATE</u>									
INJECTABLE; INJECTION									
MEDROXYPROGESTERONE ACETATE									
AB		SANDOZ	150MG/ML	N78711	001	May 20, 2009	May	NEWA	
<u>MELOXICAM</u>									
TABLET; ORAL									
MELOXICAM									
AB		BEJING YABAO	7.5MG	N77933	001	Jul 19, 2006	Apr	CAHN	
AB			15MG	N77933	002	Jul 19, 2006	Apr	CAHN	
AB		STRIDES ARCOLAB LTD	7.5MG	N77928	001	May 13, 2009	May	NEWA	
AB			15MG	N77928	002	May 13, 2009	May	NEWA	
<u>MELPHALAN HYDROCHLORIDE</u>									
INJECTABLE; INJECTION									
ALKERAN									
AP	+	GLAXOSMITHKLINE	EQ 50MG BASE/VIAL	N20207	001	Nov 18, 1992	Jun	CFTG	
MELPHALAN HYDROCHLORIDE									
AP		SYNERX	EQ 50MG BASE/VIAL	N90270	001	Jun 09, 2009	Jun	NEWA	
<u>MEPERIDINE HYDROCHLORIDE</u>									
TABLET; ORAL									
MEPERIDINE HYDROCHLORIDE									
AA		MIKART	50MG	N40893	001	Jun 24, 2009	Jun	NEWA	
			75MG	N40893	002	Jun 24, 2009	Jun	NEWA	
AA			100MG	N40893	003	Jun 24, 2009	Jun	NEWA	
			150MG	N40893	004	Jun 24, 2009	Jun	NEWA	
<u>MEPIVACAINE HYDROCHLORIDE</u>									
INJECTABLE; INJECTION									
POLOCAINE									
AP		APP PHARMS	1%	N89407	001	Dec 01, 1986	Jun	CAHN	
AP			2%	N89410	001	Dec 01, 1986	Jun	CAHN	
POLOCAINE-MPF									
AP		APP PHARMS	1%	N89406	001	Dec 01, 1986	Jun	CAHN	
AP			1.5%	N89408	001	Dec 01, 1986	Jun	CAHN	
AP			2%	N89409	001	Dec 01, 1986	Jun	CAHN	
<u>MEPROBAMATE</u>									
TABLET; ORAL									
MEPROBAMATE									
AA		ALEMBIC LTD	200MG	N90122	001	Feb 18, 2009	Feb	NEWA	
AA			400MG	N90122	002	Feb 18, 2009	Feb	NEWA	
		@ IVC INDS	400MG	N84153	001		Apr	CAHN	

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL

SOLAGE

+	STIEFEL LABS INC	2%;0.01%	N20922	001	Dec 10, 1999	Jan	CAHN
---	------------------	----------	--------	-----	--------------	-----	------

MESALAMINE

TABLET, DELAYED RELEASE; ORAL

>D>	ASACOL						
>D>	+ PROCTER AND GAMBLE	800MG	N21830	001	May 29, 2008	Jul	CTNA
	+	800MG	N21830	001	May 29, 2008	May	CTNA
>A>	ASACOL HD						
>A>	+ PROCTER AND GAMBLE	800MG	N21830	001	May 29, 2008	Jul	CTNA

MESTRANOL; NORETHINDRONE

TABLET; ORAL-21

NORETHIN 1/50M-21

@	WATSON LABS	0.05MG;1MG	N71539	001	Apr 12, 1988	Apr	DISC
---	-------------	------------	--------	-----	--------------	-----	------

NORETHINDRONE AND MESTRANOL

@	WATSON LABS	0.05MG;1MG	N70758	001	Jul 01, 1988	Apr	DISC
---	-------------	------------	--------	-----	--------------	-----	------

NORINYL 1+50 21-DAY

@	WATSON LABS	0.05MG;1MG	N13625	002		Apr	DISC
---	-------------	------------	--------	-----	--	-----	------

TABLET; ORAL-28

NORETHIN 1/50M-28

@	WATSON LABS	0.05MG;1MG	N71540	001	Apr 12, 1988	Apr	DISC
---	-------------	------------	--------	-----	--------------	-----	------

NORETHINDRONE AND MESTRANOL

@	WATSON LABS	0.05MG;1MG	N70759	001	Jul 01, 1988	Apr	DISC
---	-------------	------------	--------	-----	--------------	-----	------

NORINYL 1+50 28-DAY

+	WATSON LABS	0.05MG;1MG	N16659	001		Apr	CRLD
---	-------------	------------	--------	-----	--	-----	------

ORTHO-NOVUM 1/50 28

@	ORTHO MCNEIL JANSSEN	0.05MG;1MG	N16709	001		Apr	DISC
---	----------------------	------------	--------	-----	--	-----	------

METAPROTERENOL SULFATE

SOLUTION; INHALATION

ALUPENT

@	BOEHRINGER INGELHEIM	0.4%	N18761	002	Oct 10, 1986	Feb	DISC
---	----------------------	------	--------	-----	--------------	-----	------

@		0.6%	N18761	001	Jun 30, 1983	Feb	DISC
---	--	------	--------	-----	--------------	-----	------

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

AB	ALVOGEN	500MG	N76033	001	Jan 24, 2002	Jan	CAHN
AB		850MG	N76033	002	Jan 24, 2002	Jan	CAHN
AB		1GM	N76033	003	Jan 24, 2002	Jan	CAHN
AB	PROVIDENT PHARM	500MG	N77853	001	Jul 28, 2006	Apr	CAHN
AB		850MG	N77853	002	Jul 28, 2006	Apr	CAHN
AB		1GM	N77853	003	Jul 28, 2006	Apr	CAHN

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

@	SANDOZ	500MG	N76223	001	Dec 14, 2004	Mar	DISC
---	--------	-------	--------	-----	--------------	-----	------

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ACTOPLUS MET XR

	TAKEDA GLOBAL	1GM;EQ 15MG BASE	N22024	001	May 12, 2009	May	NEWA
--	---------------	------------------	--------	-----	--------------	-----	------

+		1GM;EQ 30MG BASE	N22024	002	May 12, 2009	May	NEWA
---	--	------------------	--------	-----	--------------	-----	------

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DESOXYN

+	LUNDBECK INC	5MG	N05378 002		Apr	CAHN
---	--------------	-----	------------	--	-----	------

TABLET, EXTENDED RELEASE; ORAL

DESOXYN

@	LUNDBECK INC	5MG	N05378 004		Apr	CAHN
---	--------------	-----	------------	--	-----	------

@		10MG	N05378 003		Apr	CAHN
---	--	------	------------	--	-----	------

@		15MG	N05378 005		Apr	CAHN
---	--	------	------------	--	-----	------

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

AB	GENPHARM	10MG	N40350 002	Mar 29, 2000	May	CRLD
----	----------	------	------------	--------------	-----	------

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

@	SOLCO HLTHCARE	500MG	N86989 001		Jan	CAHN
---	----------------	-------	------------	--	-----	------

@		750MG	N86988 001		Jan	CAHN
---	--	-------	------------	--	-----	------

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE SODIUM PRESERVATIVE FREE

AP	EBEWE PARENTA	EQ 50MG BASE/2ML (EQ 25MG BASE/ML)	N90039 001	Mar 31, 2009	Mar	NEWA
----	---------------	------------------------------------	------------	--------------	-----	------

AP		EQ 250MG BASE/10ML (EQ 25MG BASE/ML)	N90039 002	Mar 31, 2009	Mar	NEWA
----	--	--------------------------------------	------------	--------------	-----	------

AP		EQ 1GM BASE/40ML (EQ 25MG BASE/ML)	N90029 001	Mar 31, 2009	Mar	NEWA
----	--	------------------------------------	------------	--------------	-----	------

METHYCLOTHIAZIDE

TABLET; ORAL

ENDURON

ABBOTT	2.5MG	N12524 001		Mar	CTEC
--------	-------	------------	--	-----	------

METHYCLOTHIAZIDE

@ SANDOZ	2.5MG	N89835 001	Aug 18, 1988	Mar	DISC
----------	-------	------------	--------------	-----	------

@	5MG	N89837 001	Aug 18, 1988	Mar	DISC
---	-----	------------	--------------	-----	------

METHYLDOPA

TABLET; ORAL

METHYLDOPA

@ SANDOZ	125MG	N71700 001	Mar 02, 1988	Mar	DISC
----------	-------	------------	--------------	-----	------

@	250MG	N18934 001	Jun 29, 1984	Mar	DISC
---	-------	------------	--------------	-----	------

@	500MG	N18934 002	Jun 29, 1984	Mar	DISC
---	-------	------------	--------------	-----	------

METHYLPREDNISOLONE

TABLET; ORAL

MEDROL

@ PHARMACIA AND UPJOHN	24MG	N11153 005		Jun	DISC
------------------------	------	------------	--	-----	------

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE ACETATE

AB	SANDOZ	40MG/ML	N40719 001	Jan 29, 2009	Jan	NEWA
----	--------	---------	------------	--------------	-----	------

AB		40MG/ML	N40794 001	Mar 05, 2009	Feb	NEWA
----	--	---------	------------	--------------	-----	------

INJECTABLE; INJECTION

METHYLPREDNISOLONE ACETATE

AB	SANDOZ	80MG/ML	N40719 002	Jan 29, 2009	Jan	NEWA
AB		80MG/ML	N40794 002	Mar 05, 2009	Feb	NEWA

METOCLOPRAMIDE HYDROCHLORIDE

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

AA	VISTAPHARM	EQ 5MG BASE/5ML	N75051 001	Jan 26, 2001	Mar	CMFD
----	------------	-----------------	------------	--------------	-----	------

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

@ SANDOZ	EQ 5MG BASE	N74478 001	Oct 05, 1995	Mar	DISC
@	EQ 10MG BASE	N72215 001	Jan 30, 1990	Mar	DISC

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

>A>	AB	WATSON LABS FLORIDA	EQ 25MG TARTRATE	N77118 001	Aug 03, 2009	Jul	NEWA
>A>	AB		EQ 50MG TARTRATE	N76862 001	Aug 03, 2009	Jul	NEWA

METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE

AB	MUTUAL PHARM	25MG	N73654 002	Jul 15, 2009	Jun	NEWA
	@ SOLCO HLTHCARE	50MG	N74453 001	Apr 27, 1995	Jan	CAHN
	@	100MG	N74453 002	Apr 27, 1995	Jan	CAHN

METRONIDAZOLE

CAPSULE; ORAL

METRONIDAZOLE

AB	ALEMBIC LTD	375MG	N79065 001	Jun 23, 2009	Jun	NEWA
----	-------------	-------	------------	--------------	-----	------

TABLET; ORAL

METRONIDAZOLE

AB	ALEMBIC LTD	250MG	N79067 001	Mar 13, 2009	Mar	NEWA
AB		500MG	N79067 002	Mar 13, 2009	Mar	NEWA
	@ SANDOZ	250MG	N18740 001	Oct 22, 1982	Mar	DISC
	@	500MG	N18740 002	Oct 22, 1982	Mar	DISC
AB	WATSON LABS	250MG	N70035 001	Dec 20, 1984	Jun	CAHN

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

@ SANDOZ	150MG	N74450 001	May 16, 1996	Mar	DISC
@	200MG	N74450 002	May 16, 1996	Mar	DISC
@	250MG	N74450 003	May 16, 1996	Mar	DISC

MICONAZOLE NITRATE

CREAM; TOPICAL

MONISTAT-DERM

@ ORTHONEUTROGENA	2%	N17494 001		Jun	DISC
-------------------	----	------------	--	-----	------

MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+	STIEFEL LABS INC	0.25%;81.35%;15%	N21026 001	Feb 16, 2006	Jan	CAHN
---	------------------	------------------	------------	--------------	-----	------

MILNACIPRAN HYDROCHLORIDE

TABLET; ORAL

SAVELLA

CYPRESS BIOSCIENCE

12.5MG

N22256 001 Jan 14, 2009 Jan NEWA

25MG

N22256 002 Jan 14, 2009 Jan NEWA

50MG

N22256 003 Jan 14, 2009 Jan NEWA

+

100MG

N22256 004 Jan 14, 2009 Jan NEWA

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

AP

+

BEDFORD

EQ 1MG BASE/ML

N75660 001 May 28, 2002 Jun CRLD

MINOCYCLINE HYDROCHLORIDE

INJECTABLE; INJECTION

MINOCIN

+

TRIAx PHARMS LLC

EQ 100MG BASE/VIAL

N50444 001

May CMFD

TABLET, EXTENDED RELEASE; ORAL

MINOCYCLINE HYDROCHLORIDE

AB

BARR

EQ 45MG BASE

N65485 001 Mar 17, 2009 Mar NEWA

AB

EQ 90MG BASE

N65485 002 Mar 17, 2009 Mar NEWA

AB

EQ 135MG BASE

N65485 003 Mar 17, 2009 Mar NEWA

AB

IMPAX LABS INC

EQ 45MG BASE

N90024 001 Feb 03, 2009 Jan NEWA

AB

EQ 90MG BASE

N90024 002 Feb 03, 2009 Jan NEWA

AB

EQ 135MG BASE

N90024 003 Feb 03, 2009 Jan NEWA

SOLODYN

AB

MEDICIS

EQ 45MG BASE

N50808 001 May 08, 2006 Jan CFTG

>A>

EQ 65MG BASE

N50808 004 Jul 23, 2009 Jul NEWA

AB

EQ 90MG BASE

N50808 002 May 08, 2006 Jan CFTG

>A>

EQ 115MG BASE

N50808 005 Jul 23, 2009 Jul NEWA

AB

+

EQ 135MG BASE

N50808 003 May 08, 2006 Jan CFTG

MITOMYCIN

INJECTABLE; INJECTION

MUTAMYCIN

>D>

+

BRISTOL MYERS

40MG/VIAL

N62336 003 Mar 10, 1988 Jul CTEC

>A>

AP

+

40MG/VIAL

N62336 003 Mar 10, 1988 Jul CTEC

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

AP

GENERAMEDIX

EQ 20MG BASE/10ML (EQ 2MG
BASE/ML)

N78980 001 Apr 13, 2009 Mar NEWA

AP

EQ 30MG BASE/15ML (EQ 2MG
BASE/ML)

N78980 002 Apr 13, 2009 Mar NEWA

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACURIUM CHLORIDE

+

EBEWE PARENTA

EQ 2MG BASE/ML

N78562 001 Apr 30, 2009 Apr NEWA

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

KADIAN

+

ACTAVIS ELIZABETH

10MG

N20616 008 Apr 20, 2007 Feb CRLD

CAPSULE, EXTENDED RELEASE; ORAL						
KADIAN						
	+	ACTAVIS ELIZABETH	80MG	N20616	006	Oct 27, 2006 Feb CRLD
INJECTABLE; INJECTION						
ASTRAMORPH PF						
AP		APP PHARMS	0.5MG/ML	N71050	001	Oct 07, 1986 Jun CAHN
AP			0.5MG/ML	N71051	001	Oct 07, 1986 Jun CAHN
AP			1MG/ML	N71052	001	Oct 07, 1986 Jun CAHN
AP			1MG/ML	N71053	001	Oct 07, 1986 Jun CAHN
TABLET, EXTENDED RELEASE; ORAL						
MORPHINE SULFATE						
	@	AB GENERICS	15MG	N74862	001	Jul 07, 1998 Apr DISC
	@		30MG	N74862	002	Jul 07, 1998 Apr DISC
	@		60MG	N74862	003	Jul 07, 1998 Apr DISC
	@		100MG	N74769	001	Jul 02, 1998 Apr DISC
	@		200MG	N74769	002	Jul 02, 1998 Apr DISC
<u>MOXIFLOXACIN HYDROCHLORIDE</u>						
INJECTABLE; IV (INFUSION)						
AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER						
	+	BAYER HLTHCARE	160MG/100ML	N21277	001	Nov 30, 2001 Jun CAHN
TABLET; ORAL						
AVELOX						
	+	BAYER HLTHCARE	EQ 400MG BASE	N21085	001	Dec 10, 1999 Jun CAHN
<u>MYCOPHENOLATE MOFETIL</u>						
CAPSULE; ORAL						
MYCOPHENOLATE MOFETIL						
AB		ACCORD HLTHCARE INC	250MG	N90253	001	May 04, 2009 Apr NEWA
AB		APOTEX CORP	250MG	N90419	001	Apr 22, 2009 Apr NEWA
AB		MYLAN	250MG	N65520	001	May 04, 2009 Apr NEWA
AB		TEVA PHARMS	250MG	N65491	001	May 06, 2009 Apr NEWA
AB		ZYDUS PHARMS USA INC	250MG	N65433	001	May 04, 2009 Apr NEWA
TABLET; ORAL						
MYCOPHENOLATE MOFETIL						
AB		ACCORD HLTHCARE	500MG	N65416	001	May 04, 2009 Apr NEWA
AB		APOTEX	500MG	N90499	001	Apr 22, 2009 Apr NEWA
AB		MYLAN	500MG	N65521	001	May 04, 2009 Apr NEWA
AB		TEVA PHARMS	500MG	N65457	001	May 04, 2009 Apr NEWA
AB		ZYDUS PHARMS USA INC	500MG	N65477	001	May 04, 2009 Apr NEWA
<u>NABUMETONE</u>						
TABLET; ORAL						
NABUMETONE						
AB		ACTAVIS ELIZABETH	500MG	N79093	001	Feb 27, 2009 Feb NEWA
AB			750MG	N79093	002	Feb 27, 2009 Feb NEWA
	@	SANDOZ	500MG	N75590	001	Feb 25, 2002 Mar DISC
	@		750MG	N75590	002	Feb 25, 2002 Mar DISC
<u>NALTREXONE HYDROCHLORIDE</u>						
TABLET; ORAL						
NALTREXONE HYDROCHLORIDE						
		MALLINCKRODT	100MG	N76264	003	Mar 22, 2002 May CRLD

NAPROXEN SODIUM

TABLET, EXTENDED RELEASE; ORAL

NAPRELAN

+	STAT TRADE	EQ 375MG BASE	N20353 001	Jan 05, 1996	Mar	CTEC
+		EQ 500MG BASE	N20353 002	Jan 05, 1996	Mar	CTEC

NAPROXEN SODIUM

@	WATSON LABS FLORIDA	EQ 375MG BASE	N75416 002	Apr 23, 2003	Mar	DISC
@		EQ 500MG BASE	N75416 001	Aug 27, 2002	Mar	DISC

NIFEDIPINE

CAPSULE; ORAL

PROCARDIA

AB	+	PFIZER	10MG	N18482 001		Apr	CRLD
		@	20MG	N18482 002	Jul 24, 1986	Apr	DISC

TABLET, EXTENDED RELEASE; ORAL

ADALAT CC

AB1		BAYER HLTHCARE	30MG	N20198 001	Apr 21, 1993	Jun	CAHN
AB1	+		60MG	N20198 002	Apr 21, 1993	Jun	CAHN
AB1	+		90MG	N20198 003	Apr 21, 1993	Jun	CAHN

NIMODIPINE

CAPSULE; ORAL

NIMODIPINE

AB	+	BARR	30MG	N77811 001	May 02, 2007	Mar	CRLD
		NIMOTOP					
		@ BAYER PHARMS	30MG	N18869 001	Dec 28, 1988	Feb	DISC

NITISINONE

CAPSULE; ORAL

ORFADIN

		RARE DIS	2MG	N21232 001	Jan 18, 2002	Mar	CAHN
			5MG	N21232 002	Jan 18, 2002	Mar	CAHN
	+		10MG	N21232 003	Jan 18, 2002	Mar	CAHN

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

		@ SANDOZ	25MG	N74336 001	Jan 25, 1995	Mar	DISC
		@	50MG	N74336 002	Jan 25, 1995	Mar	DISC
		@	100MG	N74336 003	Jan 25, 1995	Mar	DISC

OFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

OFLOXACIN

AT		FDC LTD	0.3%	N78559 001	Feb 25, 2009	Feb	NEWA
----	--	---------	------	------------	--------------	-----	------

SOLUTION/DROPS; OTIC

OFLOXACIN

>A>	AT	PHARMAFORCE	0.3%	N90395 001	Aug 11, 2009	Jul	NEWA
-----	----	-------------	------	------------	--------------	-----	------

TABLET; ORAL

FLOXIN

		@ ORTHO MCNEIL JANSSEN	200MG	N19735 001	Dec 28, 1990	Mar	DISC
		@	300MG	N19735 002	Dec 28, 1990	Mar	DISC
		@	400MG	N19735 003	Dec 28, 1990	Mar	DISC

OFLOXACIN

>D>	AB	RANBAXY	200MG	N76220 001	Sep 02, 2003	Jul	DISC
-----	----	---------	-------	------------	--------------	-----	------

TABLET; ORAL

	OFLOXACIN							
>A>	@ RANBAXY	200MG	N76220 001	Sep 02, 2003	Jul	DISC		
>D>	AB	300MG	N76220 002	Sep 02, 2003	Jul	DISC		
>A>	@	300MG	N76220 002	Sep 02, 2003	Jul	DISC		
>D>	AB	400MG	N76220 003	Sep 02, 2003	Jul	DISC		
>A>	@	400MG	N76220 003	Sep 02, 2003	Jul	DISC		

OLSALAZINE SODIUM

CAPSULE; ORAL

DIPENTUM

+	ALAVEN PHARM	250MG	N19715 001	Jul 31, 1990	Apr	CAHN
+	UCB INC	250MG	N19715 001	Jul 31, 1990	Jun	CAHN

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE

AB	DR REDDYS LABS	40MG	N78490 001	Apr 17, 2009	Mar	NEWA
AB	DR REDDYS LABS LTD	10MG	N78693 001	Mar 16, 2009	Mar	NEWA
AB		20MG	N78693 002	Mar 16, 2009	Mar	NEWA
AB	KREMERS URBAN DEV	40MG	N75410 003	Jan 23, 2009	Jan	NEWA

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE; ORAL

OMEPRAZOLE MAGNESIUM

	DR REDDYS LABS LTD	20MG	N78878 001	Jun 05, 2009	May	NEWA
--	--------------------	------	------------	--------------	-----	------

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

AP	BEDFORD LABS	EQ 0.64MG BASE/ML	N78291 001	Apr 13, 2009	Mar	NEWA
----	--------------	-------------------	------------	--------------	-----	------

ORLISTAT

CAPSULE; ORAL

XENICAL

>D>	+	HLR	120MG	N20766 001	Apr 23, 1999	Jul	CAHN
>A>	+	HOFFMANN LA ROCHE	120MG	N20766 001	Apr 23, 1999	Jul	CAHN

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

@	WATSON LABS	EQ 250MG BASE/VIAL	N62856 001	Oct 26, 1988	Apr	DISC
@		EQ 500MG BASE/VIAL	N62856 002	Oct 26, 1988	Apr	DISC
@		EQ 1GM BASE/VIAL	N62856 003	Oct 26, 1988	Apr	DISC
@		EQ 2GM BASE/VIAL	N62856 004	Oct 26, 1988	Apr	DISC
@		EQ 4GM BASE/VIAL	N62856 005	Oct 26, 1988	Apr	DISC

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

>D>	+	SANOFI AVENTIS US	50MG/10ML (5MG/ML)	N21759 001	Jan 31, 2005	Jul	CFTG
>A>	AP	+	50MG/10ML (5MG/ML)	N21759 001	Jan 31, 2005	Jul	CFTG
>D>	+		100MG/20ML (5MG/ML)	N21759 002	Jan 31, 2005	Jul	CFTG
>A>	AP	+	100MG/20ML (5MG/ML)	N21759 002	Jan 31, 2005	Jul	CFTG
>A>		OXALIPLATIN					
>A>	AP	EBEWE PHARMA	50MG/10ML (5MG/ML)	N78812 001	Aug 07, 2009	Jul	NEWA

INJECTABLE; IV (INFUSION)

>A> OXALIPLATIN

>A>	AP	EBEWE PHARMA	100MG/20ML (5MG/ML)	N78812 002	Aug 07, 2009	Jul	NEWA
>A>	AP	FRESENIUS KABI ONCOL	50MG/VIAL	N78810 001	Aug 07, 2009	Jul	NEWA
>A>	AP		100MG/VIAL	N78810 002	Aug 07, 2009	Jul	NEWA
>A>	AP	HOSPIRA WORLDWIDE	50MG/10ML (5MG/ML)	N78813 001	Aug 07, 2009	Jul	NEWA
>A>	AP		100MG/20ML (5MG/ML)	N78813 002	Aug 07, 2009	Jul	NEWA
>A>	AP	SUN PHARMA GLOBAL	50MG/VIAL	N78818 001	Aug 07, 2009	Jul	NEWA
>A>	AP		100MG/VIAL	N78818 002	Aug 07, 2009	Jul	NEWA

OXAPROZIN

TABLET; ORAL

OXAPROZIN

@ SANDOZ

600MG

N75850 001 Apr 27, 2001 Mar DISC

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

AB		RANBAXY	300MG/5ML	N78734 001	Jun 26, 2009	Jun	NEWA
----	--	---------	-----------	------------	--------------	-----	------

TRILEPTAL

AB	+	NOVARTIS	300MG/5ML	N21285 001	May 25, 2001	Jun	CFTG
----	---	----------	-----------	------------	--------------	-----	------

OXTRIPHYLLINE

TABLET, EXTENDED RELEASE; ORAL

CHOLEDYL SA

+ WARNER CHILCOTT 400MG

N87863 001 May 24, 1983 Jun CAHN

OXYBUTYNIN CHLORIDE

GEL; TRANSDERMAL

GELNIQUE

+		WATSON LABS	10%(100MG/PACKET)	N22204 001	Jan 27, 2009	Mar	CTNA
---	--	-------------	-------------------	------------	--------------	-----	------

OXYBUTYNIN CHLORIDE

+		WATSON LABS	10%(100MG/PACKET)	N22204 001	Jan 27, 2009	Jan	NEWA
---	--	-------------	-------------------	------------	--------------	-----	------

TABLET, EXTENDED RELEASE; ORAL

OXYBUTYNIN CHLORIDE

AB		OSMOTICA PHARM	5MG	N78503 001	Feb 04, 2009	Jan	NEWA
----	--	----------------	-----	------------	--------------	-----	------

AB			10MG	N78503 002	Feb 04, 2009	Jan	NEWA
----	--	--	------	------------	--------------	-----	------

AB			15MG	N78503 003	Feb 04, 2009	Jan	NEWA
----	--	--	------	------------	--------------	-----	------

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

AB		SUN PHARM INDS INC	5MG	N90659 001	Apr 10, 2009	Mar	NEWA
----	--	--------------------	-----	------------	--------------	-----	------

AB			15MG	N90659 002	Apr 10, 2009	Mar	NEWA
----	--	--	------	------------	--------------	-----	------

AB			30MG	N90659 003	Apr 10, 2009	Mar	NEWA
----	--	--	------	------------	--------------	-----	------

AB		VINTAGE PHARMS	5MG	N77712 003	Mar 02, 2009	Mar	NEWA
----	--	----------------	-----	------------	--------------	-----	------

ROXICODONE

AB		XANODYNE PHARMS	5MG	N21011 003	May 15, 2009	May	NEWA
----	--	-----------------	-----	------------	--------------	-----	------

>A> PALIPERIDONE PALMITATE

>A> SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

>A> INVEGA SUSTENNA

>A>		JOHNSON AND JOHNSON	39MG/0.25ML (39MG/0.25ML)	N22264 001	Jul 31, 2009	Jul	NEWA
-----	--	---------------------	---------------------------	------------	--------------	-----	------

>A>			78MG/0.5ML (78MG/0.5ML)	N22264 002	Jul 31, 2009	Jul	NEWA
-----	--	--	-------------------------	------------	--------------	-----	------

>A>			117MG/0.75ML (117MG/0.75ML)	N22264 003	Jul 31, 2009	Jul	NEWA
-----	--	--	-----------------------------	------------	--------------	-----	------

>A>			156MG/ML (156MG/ML)	N22264 004	Jul 31, 2009	Jul	NEWA
-----	--	--	---------------------	------------	--------------	-----	------

>A> SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR
 >A> INVEGA SUSTENNA
 >A> JOHNSON AND JOHNSON 234MG/1.5ML (156MG/ML) N22264 005 Jul 31, 2009 Jul NEWA

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION
 PAMIDRONATE DISODIUM
 AP GENERAMEDIX 30MG/VIAL N78300 001 Mar 10, 2009 Feb NEWA
 AP 90MG/VIAL N78300 002 Mar 10, 2009 Feb NEWA

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

>D> CAPSULE; ORAL
 >D> CREON
 >D> SOLVAY 30,000USP UNITS;6,000USP UNITS;19,000USP UNITS N20725 001 Apr 30, 2009 Jul CDFR
 30,000USP UNITS;6,000USP UNITS;19,000USP UNITS N20725 001 Apr 30, 2009 Apr NEWA
 >D> 60,000USP UNITS;12,000USP UNITS;38,000USP UNITS N20725 002 Apr 30, 2009 Jul CDFR
 60,000USP UNITS;12,000USP UNITS;38,000USP UNITS N20725 002 Apr 30, 2009 Apr NEWA
 >D> + 120,000USP UNITS;24,000USP UNITS;76,000USP UNITS N20725 003 Apr 30, 2009 Jul CDFR
 + 120,000USP UNITS;24,000USP UNITS;76,000USP UNITS N20725 003 Apr 30, 2009 Apr NEWA
 >A> CAPSULE, DELAYED RELEASE; ORAL
 >A> CREON
 >A> SOLVAY 30,000USP UNITS;6,000USP UNITS;19,000USP UNITS N20725 001 Apr 30, 2009 Jul CDFR
 >A> 60,000USP UNITS;12,000USP UNITS;38,000USP UNITS N20725 002 Apr 30, 2009 Jul CDFR
 >A> + 120,000USP UNITS;24,000USP UNITS;76,000USP UNITS N20725 003 Apr 30, 2009 Jul CDFR

PANCURONIUM BROMIDE

INJECTABLE; INJECTION
 PANCURONIUM BROMIDE
 @ HOSPIRA 2MG/ML N72321 001 Jan 19, 1989 Apr DISC

PANTOPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL
 PANTOPRAZOLE SODIUM
 AB KUDCO IRELAND EQ 20MG BASE N78281 001 Mar 17, 2009 Mar NEWA
 AB EQ 40MG BASE N78281 002 Mar 17, 2009 Mar NEWA

PAROMOMYCIN SULFATE

CAPSULE; ORAL
 PAROMOMYCIN SULFATE
 >A> AA HERITAGE PHARMS INC EQ 250MG BASE N65173 001 Dec 14, 2007 Jul CAHN
 >D> AA X-GEN PHARMS EQ 250MG BASE N65173 001 Dec 14, 2007 Jul CAHN

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION
 PENICILLIN G POTASSIUM
 >A> HANFORD GC 1,000,000 UNITS/VIAL N65149 001 Jul 23, 2009 Jul NEWA
 >A> AP 5,000,000 UNITS/VIAL N65149 002 Jul 23, 2009 Jul NEWA
 >A> AP 20,000,000 UNITS/VIAL N65149 003 Jul 23, 2009 Jul NEWA

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN-VK

AA	+	TEVA	EQ 250MG BASE/5ML	N60456 002	Apr	CRLD
		VEETIDS				
		@ APOTHECON	EQ 125MG BASE/5ML	N61410 001	Mar	DISC
		@	EQ 250MG BASE/5ML	N61410 002	Mar	DISC

PENTAMIDINE ISETHIONATE

FOR SOLUTION; INHALATION

NEBUPENT

@ APP PHARMS	600MG/VIAL	N19887 002	Mar 22, 1996	May	DISC
--------------	------------	------------	--------------	-----	------

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUM

+	LUNDBECK INC	50MG/ML	N83246 001	Jun	CAHN
---	--------------	---------	------------	-----	------

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENDIMETRAZINE TARTRATE

@ SANDOZ	35MG	N85695 001	May	DISC
----------	------	------------	-----	------

PHENYTOIN

SUSPENSION; ORAL

PHENYTOIN

AB		WOCKHARDT EU	125MG/5ML	N40420 001	Apr 19, 2002	Apr	CAHN
----	--	--------------	-----------	------------	--------------	-----	------

PHENYTOIN SODIUM

CAPSULE; ORAL

PROMPT PHENYTOIN SODIUM

@ IVAX PHARMS	100MG PROMPT	N80259 001	Jan	DISC
---------------	--------------	------------	-----	------

PILOCARPINE HYDROCHLORIDE

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

AB		LANNETT	7.5MG	N77220 002	May 06, 2009	Apr	NEWA
----	--	---------	-------	------------	--------------	-----	------

PINDOLOL

TABLET; ORAL

PINDOLOL

@ SANDOZ	5MG	N73608 001	Mar 29, 1993	Mar	DISC
----------	-----	------------	--------------	-----	------

@	10MG	N73609 001	Mar 29, 1993	Mar	DISC
---	------	------------	--------------	-----	------

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

POLYETHYLENE GLYCOL 3350

AA		BRECKENRIDGE PHARM	17GM/SC00PFUL	N77736 001	May 26, 2006	Jun	CAHN
AA		GAVIS PHARMS	17GM/SC00PFUL	N77736 001	May 26, 2006	Jan	CAHN
		@ TEVA PHARMS	17GM/SC00PFUL	N77445 001	May 04, 2006	Jan	DISC

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

FOR SOLUTION; ORAL

PEG-3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

AA	NOVEL LABS INC	420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 1.2GM/BOT	N90019 001	May 27, 2009	May	NEWA
----	----------------	--	------------	--------------	-----	------

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

GOLYTELY

	+	BRAINTREE	227.1GM/PACKET; 2.82GM/PACKET; 6.36GM/PACKET; 5.53GM/PACKET; 21.5GM/PACKET	N19011 002	Jun 02, 1992	May	CTEC
AA	+		236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	N19011 001	Jul 13, 1984	May	CFTG
	+		236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	N19011 001	Jul 13, 1984	Mar	CTEC

PEG 3350 AND ELECTROLYTES

AA	NOVEL LABS INC	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	N90231 001	Jun 01, 2009	May	NEWA
AA		240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT	N90186 001	Jun 01, 2009	May	NEWA

FOR SUSPENSION; ORAL

CO-LAV

	@ BOCA PHARMA	240GM/BOT; 2.98GM/BOT; 6.72GM/BOT; 5.84GM/BOT; 22.72GM/BOT	N73428 001	Jan 28, 1992	Mar	DISC
--	---------------	--	------------	--------------	-----	------

GO-EVAC

	@ BOCA PHARMA	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	N73433 001	Apr 28, 1992	Mar	DISC
--	---------------	--	------------	--------------	-----	------

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MICRO-K

	@ KV PHARM	8MEQ	N18238 001		Feb	DISC
--	------------	------	------------	--	-----	------

MICRO-K 10

	@ KV PHARM	10MEQ	N18238 002	May 14, 1984	Feb	DISC
--	------------	-------	------------	--------------	-----	------

POTASSIUM CHLORIDE

	@ KV PHARM	10MEQ	N70980 001	Feb 17, 1987	Feb	DISC
--	------------	-------	------------	--------------	-----	------

WATSON LABS FLORIDA

		8MEQ	N77419 001	Jun 02, 2008	Feb	CTEC
--	--	------	------------	--------------	-----	------

+ 10MEQ

			N77419 002	Jun 02, 2008	Feb	CRLD
--	--	--	------------	--------------	-----	------

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

	@ HOSPIRA	1.5MEQ/ML	N83345 001		Apr	DISC
--	-----------	-----------	------------	--	-----	------

TABLET, EXTENDED RELEASE; ORAL

POTASSIUM CHLORIDE

AB	WATSON LABS FLORIDA	10MEQ	N75604 001	Apr 10, 2002	May	CRLD
----	---------------------	-------	------------	--------------	-----	------

AB	+	10MEQ	N75604 001	Apr 10, 2002	Jan	CRLD
----	---	-------	------------	--------------	-----	------

>A> PRASUGREL HYDROCHLORIDE

>A> TABLET; ORAL

>A> EFFIENT

>A>	ELI LILLY AND CO	EQ 5MG BASE	N22307 001	Jul 10, 2009	Jul	NEWA
-----	------------------	-------------	------------	--------------	-----	------

>A>	+	EQ 10MG BASE	N22307 002	Jul 10, 2009	Jul	NEWA
-----	---	--------------	------------	--------------	-----	------

PRAZQUANTEL

TABLET; ORAL

BILTRICIDE

+	BAYER HLTHCARE	600MG	N18714 001	Dec 29, 1982	Jun	CAHN
---	----------------	-------	------------	--------------	-----	------

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

@ SANDOZ	EQ 1MG BASE	N72576 001	May 16, 1989	Mar	DISC
@	EQ 2MG BASE	N72577 001	May 16, 1989	Mar	DISC
@	EQ 5MG BASE	N72578 001	May 16, 1989	Mar	DISC

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

ORAPRED

>D>	AA	+	ASCENT PEDS	EQ 15MG BASE/5ML	N75117 001	Dec 14, 2000	Jul	CAHN
>A>	AA	+	SCIELE PHARMA INC	EQ 15MG BASE/5ML	N75117 001	Dec 14, 2000	Jul	CAHN

PREDNISOLONE SODIUM PHOSPHATE

AA	AMNEAL PHARMS	EQ 15MG BASE/5ML	N78345 001	Mar 10, 2009	Feb	NEWA
AA	VINTAGE	EQ 15MG BASE/5ML	N79010 001	May 26, 2009	May	NEWA

SOLUTION/DROPS; OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE

@ BAUSCH AND LOMB	EQ 0.11% PHOSPHATE	N40065 001	Jul 29, 1994	Apr	DISC
-------------------	--------------------	------------	--------------	-----	------

TABLET, ORALLY DISINTEGRATING; ORAL

ORAPRED ODT

>D>		MEDICIS	EQ 10MG BASE	N21959 001	Jun 01, 2006	Jul	CAHN
>D>			EQ 15MG BASE	N21959 002	Jun 01, 2006	Jul	CAHN
>D>		+	EQ 30MG BASE	N21959 003	Jun 01, 2006	Jul	CAHN
>A>		SCIELE PHARMA INC	EQ 10MG BASE	N21959 001	Jun 01, 2006	Jul	CAHN
>A>			EQ 15MG BASE	N21959 002	Jun 01, 2006	Jul	CAHN
>A>		+	EQ 30MG BASE	N21959 003	Jun 01, 2006	Jul	CAHN

PREDNISONE

TABLET; ORAL

PREDNISONE

@ SANDOZ	10MG	N89983 001	Jan 12, 1989	Mar	DISC
@	50MG	N89984 001	Jan 12, 1989	Mar	DISC

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST PLAIN DENTAL

+	DENTSPLY PHARM	4%	N21382 001		Apr	CTNA
---	----------------	----	------------	--	-----	------

PROGESTERONE

INJECTABLE; INJECTION

PROGESTERONE

>D>	AP	PHARMAFORCE	50MG/ML	N90845 001	Jun 22, 2009	Jul	CTEC
>A>	AO		50MG/ML	N90845 001	Jun 22, 2009	Jul	CTEC
	AP		50MG/ML	N90845 001	Jun 22, 2009	Jun	NEWA

PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

AA	SUN PHARM INDS INC	6.25MG/5ML	N40891 001	Mar 13, 2009	Mar	NEWA
----	--------------------	------------	------------	--------------	-----	------

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

>D>	AB	SANDOZ	12.5MG	N84176 002	May 22, 2009	Jul	DISC
>A>		@	12.5MG	N84176 002	May 22, 2009	Jul	DISC
	AB		12.5MG	N84176 002	May 22, 2009	May	NEWA

PROPANTHELINE BROMIDE

TABLET; ORAL

PROPANTHELINE BROMIDE

>D>	+	ROXANE	7.5MG	N80927 001		Jul	DISC
>A>	@		7.5MG	N80927 001		Jul	DISC

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

AB	+	APP PHARMS	10MG/ML	N19627 002	Jun 11, 1996	Jun	CAHN
	@		10MG/ML	N19627 001	Oct 02, 1989	Jun	CAHN

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DOLENE

AA		HERITAGE PHARMS INC	65MG	N80530 001		Mar	CMFD
AA		VINTAGE PHARMS	65MG	N40908 001	Jul 17, 2009	Jun	NEWA

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

AB		UPSHER SMITH	60MG	N78311 001	Mar 06, 2009	Feb	NEWA
AB			80MG	N78311 002	Mar 06, 2009	Feb	NEWA
AB			120MG	N78311 003	Mar 06, 2009	Feb	NEWA
AB			160MG	N78311 004	Mar 06, 2009	Feb	NEWA

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

	@	SANDOZ	10MG	N70663 001	Jun 13, 1986	Mar	DISC
	@		20MG	N70664 001	Jun 13, 1986	Mar	DISC
	@		40MG	N70665 001	Jun 13, 1986	Mar	DISC
	@		60MG	N70666 001	Oct 10, 1986	Mar	DISC
	@		80MG	N70667 001	Jun 13, 1986	Mar	DISC

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL

CORPHED

	@	SANDOZ	60MG;2.5MG	N88602 001	Apr 11, 1985	Mar	DISC
--	---	--------	------------	------------	--------------	-----	------

PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE

+		SANDOZ	60MG;2.5MG	N88193 001	May 17, 1983	Mar	CTEC
---	--	--------	------------	------------	--------------	-----	------

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE

AB		SUN PHARM INDS LTD	EQ 5MG BASE	N90800 001	Jun 18, 2009	Jun	NEWA
AB			EQ 10MG BASE	N90800 002	Jun 18, 2009	Jun	NEWA
AB			EQ 20MG BASE	N90800 003	Jun 18, 2009	Jun	NEWA
AB			EQ 40MG BASE	N90800 004	Jun 18, 2009	Jun	NEWA

RAMIPRIL

CAPSULE; ORAL

RAMIPRIL

AB		RANBAXY	5MG	N78849 001	Mar 06, 2009	Feb	NEWA
AB			10MG	N78849 002	Mar 06, 2009	Feb	NEWA

RANITIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

RANITIDINE HYDROCHLORIDE

@ BEDFORD

EQ 25MG BASE/ML

N74764 001 Nov 19, 2004 Apr DISC

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

AA AMNEAL PHARMS EQ 15MG BASE/ML

N78312 001 Sep 02, 2008 Feb CTNA

AA DR REDDYS LABS LTD EQ 15MG BASE/ML

N90102 001 May 26, 2009 May NEWA

AA WOCKHARDT EQ 15MG BASE/ML

N79211 001 May 26, 2009 May NEWA

AA EQ 15MG BASE/ML

N79212 001 Feb 23, 2009 Feb NEWA

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

>D> AB RANBAXY EQ 150MG BASE

N75439 001 Apr 19, 2000 Jul DISC

>A> @ EQ 150MG BASE

N75439 001 Apr 19, 2000 Jul DISC

>D> AB EQ 300MG BASE

N75439 002 Apr 19, 2000 Jul DISC

>A> @ EQ 300MG BASE

N75439 002 Apr 19, 2000 Jul DISC

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

GILEAD

500MG

N21526 002 Jan 27, 2006 Jun CAHN

+

1GM

N21526 001 Feb 12, 2007 Jun CAHN

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

@ PROCTER AND GAMBLE

75MG

N20835 004 Apr 16, 2007 May DISC

RISPERIDONE

SOLUTION; ORAL

RISPERDAL

AA + ORTHO MCNEIL JANSSEN 1MG/ML

N20588 001 Jun 10, 1996 Jan CFTG

RISPERIDONE

>A> AA APOTEX INC 1MG/ML

N77719 001 Jul 29, 2009 Jul NEWA

>A> AA DR REDDYS LABS LTD 1MG/ML

N78909 001 Jul 29, 2009 Jul NEWA

>A> AA ROXANE 1MG/ML

N76904 001 Jul 29, 2009 Jul NEWA

AA TEVA 1MG/ML

N76440 001 Jan 30, 2009 Jan NEWA

TABLET; ORAL

RISPERIDONE

AB ACTAVIS TOTOWA 0.25MG

N78071 001 Jun 17, 2009 Jun NEWA

AB 0.5MG

N78071 002 Jun 17, 2009 Jun NEWA

AB 1MG

N78071 003 Jun 17, 2009 Jun NEWA

AB 2MG

N78071 004 Jun 17, 2009 Jun NEWA

AB 3MG

N78071 005 Jun 17, 2009 Jun NEWA

AB 4MG

N78071 006 Jun 17, 2009 Jun NEWA

AB CADISTA PHARMS 0.25MG

N78828 001 Mar 23, 2009 Mar NEWA

AB 0.5MG

N78828 002 Mar 23, 2009 Mar NEWA

AB 1MG

N78828 003 Mar 23, 2009 Mar NEWA

AB 2MG

N78828 004 Mar 23, 2009 Mar NEWA

AB 3MG

N78828 005 Mar 23, 2009 Mar NEWA

AB 4MG

N78828 006 Mar 23, 2009 Mar NEWA

AB WEST WARD PHARMS 0.25MG

N78740 001 May 29, 2009 May NEWA

AB 0.5MG

N78740 002 May 29, 2009 May NEWA

AB 1MG

N78740 003 May 29, 2009 May NEWA

AB 2MG

N78740 004 May 29, 2009 May NEWA

TABLET; ORAL

RISPERIDONE

AB	WEST WARD PHARMS	3MG	N78740 005	May 29, 2009	May	NEWA
AB		4MG	N78740 006	May 29, 2009	May	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

AB	ORTHO MCNEIL JANSSEN	0.5MG	N21444 001	Apr 02, 2003	Feb	CFTG
AB		2MG	N21444 003	Apr 02, 2003	Feb	CFTG
AB		3MG	N21444 004	Dec 23, 2004	Apr	CFTG
AB		4MG	N21444 005	Dec 23, 2004	Apr	CFTG

RISPERIDONE

AB	DR REDDYS LABS LTD	0.5MG	N77328 001	Feb 24, 2009	Feb	NEWA
AB		2MG	N77328 003	Feb 24, 2009	Feb	NEWA
	KALI LABS	0.25MG	N77494 001	Apr 30, 2009	Apr	NEWA
AB		0.5MG	N77494 002	Apr 30, 2009	Apr	NEWA
AB		2MG	N77494 004	Apr 30, 2009	Apr	NEWA
AB		3MG	N77494 005	Apr 30, 2009	Apr	NEWA
AB		4MG	N77494 006	Apr 30, 2009	Apr	NEWA
	PAR PHARM	0.25MG	N77494 001	Apr 30, 2009	Jun	CAHN
AB		0.5MG	N77494 002	Apr 30, 2009	Jun	CAHN
AB		2MG	N77494 004	Apr 30, 2009	Jun	CAHN
AB		3MG	N77494 005	Apr 30, 2009	Jun	CAHN
AB		4MG	N77494 006	Apr 30, 2009	Jun	CAHN
AB	ZYDUS PHARMS USA	0.5MG	N78516 001	May 01, 2009	Apr	NEWA
AB		2MG	N78516 003	May 01, 2009	Apr	NEWA

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION

RITODRINE HYDROCHLORIDE

@	HOSPIRA	10MG/ML	N71618 001	Feb 28, 1991	Apr	DISC
@		15MG/ML	N71619 001	Feb 28, 1991	Apr	DISC
	RITODRINE HYDROCHLORIDE	IN DEXTROSE 5% IN PLASTIC CONTAINER				
@	HOSPIRA	30MG/100ML	N71438 001	Jan 22, 1991	Apr	DISC

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

AB	HUAHAI US INC	EQ 0.25MG BASE	N78110 001	May 05, 2008	Apr	CAHN
AB		EQ 0.5MG BASE	N78110 002	May 05, 2008	Apr	CAHN
AB		EQ 1MG BASE	N78110 003	May 05, 2008	Apr	CAHN
AB		EQ 2MG BASE	N78110 004	May 05, 2008	Apr	CAHN
AB		EQ 3MG BASE	N78110 005	May 05, 2008	Apr	CAHN
AB		EQ 4MG BASE	N78110 006	May 05, 2008	Apr	CAHN
AB	ZYDUS PHARMS USA INC	EQ 0.25MG BASE	N90411 001	Jun 01, 2009	May	NEWA
AB		EQ 0.5MG BASE	N90411 002	Jun 01, 2009	May	NEWA
AB		EQ 1MG BASE	N90411 003	Jun 01, 2009	May	NEWA
AB		EQ 2MG BASE	N90411 004	Jun 01, 2009	May	NEWA
AB		EQ 3MG BASE	N90411 005	Jun 01, 2009	May	NEWA
AB		EQ 4MG BASE	N90411 006	Jun 01, 2009	May	NEWA
AB		EQ 5MG BASE	N90411 007	Jun 01, 2009	May	NEWA

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

	SMITHKLINE BEECHAM	EQ 6MG BASE	N22008 006	Apr 10, 2009	Apr	NEWA
--	--------------------	-------------	------------	--------------	-----	------

SAMARIUM SM 153 LEXIDRONAM PENTASODIUM

INJECTABLE; INJECTION

QUADRAMET

+	EUSA PHARMA USA	50mCi/ML	N20570 001	Mar 28, 1997	May	CAHN
---	-----------------	----------	------------	--------------	-----	------

>A> SAXAGLIPTIN HYDROCHLORIDE

>A> TABLET; ORAL

>A> ONGLYZA

>A>	BRISTOL MYERS SQUIBB	EQ 2.5MG BASE	N22350 001	Jul 31, 2009	Jul	NEWA
-----	----------------------	---------------	------------	--------------	-----	------

>A>	+	EQ 5MG BASE	N22350 002	Jul 31, 2009	Jul	NEWA
-----	---	-------------	------------	--------------	-----	------

SELEGILINE HYDROCHLORIDE

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

@ ENDO PHARMS 5MG

N74565 001 Aug 02, 1996 Apr DISC

@ SIEGFRIED 5MG

N74672 001 Apr 01, 1997 Apr DISC

SERTACONAZOLE NITRATE

CREAM; TOPICAL

ERTACZO

>A>	+	ORTHO DERMATOLOGICS	2%	N21385 001	Dec 10, 2003	Jul	CAHN
-----	---	---------------------	----	------------	--------------	-----	------

>D>	+	ORTHONEUTROGENA	2%	N21385 001	Dec 10, 2003	Jul	CAHN
-----	---	-----------------	----	------------	--------------	-----	------

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB	AUSTARPHARMA LLC	EQ 25MG BASE	N78677 001	Mar 04, 2009	Feb	NEWA
----	------------------	--------------	------------	--------------	-----	------

AB		EQ 50MG BASE	N78677 002	Mar 04, 2009	Feb	NEWA
----	--	--------------	------------	--------------	-----	------

AB		EQ 100MG BASE	N78677 003	Mar 04, 2009	Feb	NEWA
----	--	---------------	------------	--------------	-----	------

>A>	AB	HIKMA PHARMS	EQ 25MG BASE	N77864 001	Aug 10, 2009	Jul	NEWA
-----	----	--------------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N77864 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

>A>	AB		EQ 100MG BASE	N77864 003	Aug 10, 2009	Jul	NEWA
-----	----	--	---------------	------------	--------------	-----	------

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

AB	LUPIN	5MG	N78103 005	Apr 14, 2009	Mar	NEWA
----	-------	-----	------------	--------------	-----	------

SODIUM PHOSPHATE, P-32

SOLUTION; INJECTION, ORAL

SODIUM PHOSPHATE P 32

@	MALLINCKRODT	0.67mCi/ML	N11777 001		May	DISC
---	--------------	------------	------------	--	-----	------

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KALEXATE

AA	KVK TECH	454GM/BOT	N40905 001	Mar 30, 2009	Mar	NEWA
----	----------	-----------	------------	--------------	-----	------

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

>D> ACCRETROPIN

>D>	+	CANGENE	5MG/ML (5MG/ML)	N21538 001	Jan 23, 2008	Jul	DISC
-----	---	---------	-----------------	------------	--------------	-----	------

>A>	@		5MG/ML (5MG/ML)	N21538 001	Jan 23, 2008	Jul	DISC
-----	---	--	-----------------	------------	--------------	-----	------

NORDITROPIN NORDIFLEX

	NOVO NORDISK INC	30MG/3ML	N21148 007	Mar 10, 2009	Mar	NEWA
--	------------------	----------	------------	--------------	-----	------

INJECTABLE; INJECTIONSEROSTIM

>D>	EMD SERONO	8.8MG/VIAL	N20604 004	Sep 06, 2001	Jul	DISC
>A>	@	8.8MG/VIAL	N20604 004	Sep 06, 2001	Jul	DISC

SORAFENIB TOSYLATETABLET; ORALNEXAVAR

+	BAYER HLTHCARE	EQ 200MG BASE	N21923 001	Dec 20, 2005	Jun	CAHN
---	----------------	---------------	------------	--------------	-----	------

SOTALOL HYDROCHLORIDE

>A>	SOLUTION; INTRAVENOUS					
>A>	SOTALOL HYDROCHLORIDE					
>A>	+	ACADEMIC PHARMS	150MG/10ML (15MG/ML)	N22306 001	Jul 02, 2009	Jul NEWA

STANZOLOLTABLET; ORALWINSTROL

@	LUNDBECK INC	2MG	N12885 001	May 14, 1984	Mar	CAHN
---	--------------	-----	------------	--------------	-----	------

STAVUDINEFOR SOLUTION; ORALSTAVUDINE

AA	CIPLA LTD	1MG/ML	N78030 001	Mar 20, 2009	Mar	NEWA
----	-----------	--------	------------	--------------	-----	------

SUCCIMERCAPSULE; ORALCHEMET

+	LUNDBECK INC	100MG	N19998 002	Jan 30, 1991	Mar	CAHN
---	--------------	-------	------------	--------------	-----	------

SULFACETAMIDE SODIUMLOTION; TOPICALSULFACETAMIDE SODIUM

AB	PERRIGO CO TENNESSEE	10%	N78649 001	Mar 23, 2009	Mar	NEWA
----	----------------------	-----	------------	--------------	-----	------

AB	TARO	10%	N78668 001	May 20, 2009	May	NEWA
----	------	-----	------------	--------------	-----	------

SOLUTION/DROPS; OPHTHALMICSULF-10

@	NOVARTIS	10%	N80025 001		Apr	DISC
---	----------	-----	------------	--	-----	------

SULFACETAMIDE SODIUM

@	ALCON	30%	N89068 001	May 05, 1987	Apr	DISC
---	-------	-----	------------	--------------	-----	------

SULFAMETHOXAZOLE; TRIMETHOPRIMTABLET; ORALSULFAMETHOPRIM

AB	PAR PHARM	400MG;80MG	N70022 001	Feb 15, 1985	Mar	CMFD
----	-----------	------------	------------	--------------	-----	------

SULFAMETHOPRIM-DS

AB	PAR PHARM	800MG;160MG	N70032 001	Feb 15, 1985	Mar	CMFD
----	-----------	-------------	------------	--------------	-----	------

SULFAMETHOXAZOLE AND TRIMETHOPRIM

@	SANDOZ	800MG;160MG	N70890 001	Nov 13, 1986	Mar	DISC
---	--------	-------------	------------	--------------	-----	------

SULFISOXAZOLE ACETYLSUSPENSION; ORALGANTRISIN PEDIATRIC

@	ROCHE	EQ 500MG BASE/5ML	N09182 004		Jun	DISC
---	-------	-------------------	------------	--	-----	------

SULINDAC

TABLET; ORAL

SULINDAC

@ SANDOZ

150MG

N72712 001 Aug 30, 1991 Mar DISC

@

200MG

N72713 001 Aug 30, 1991 Mar DISC

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX

AP	+	GLAXOSMITHKLINE	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N20080 001	Dec 28, 1992	Jan	CFTG
----	---	-----------------	-------------------------------------	------------	--------------	-----	------

SUMATRIPTAN SUCCINATE

AP		APP PHARMS	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N79242 001	Mar 02, 2009	Feb	NEWA
----	--	------------	-------------------------------------	------------	--------------	-----	------

AP		BEDFORD	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N79123 001	Feb 06, 2009	Jan	NEWA
----	--	---------	-------------------------------------	------------	--------------	-----	------

AP		JHP PHARMS	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N77871 001	Jul 09, 2009	Jun	NEWA
----	--	------------	-------------------------------------	------------	--------------	-----	------

AP		SANDOZ	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	N78067 002	Feb 06, 2009	Jan	NEWA
----	--	--------	------------------------------------	------------	--------------	-----	------

AP			EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N78067 001	Feb 06, 2009	Jan	NEWA
----	--	--	-------------------------------------	------------	--------------	-----	------

AP		TEVA PARENTERAL	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)	N78318 001	Feb 06, 2009	Jan	NEWA
----	--	-----------------	------------------------------------	------------	--------------	-----	------

AP			EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N78318 002	Feb 06, 2009	Jan	NEWA
----	--	--	-------------------------------------	------------	--------------	-----	------

AP			EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N77907 001	Feb 06, 2009	Jan	NEWA
----	--	--	-------------------------------------	------------	--------------	-----	------

AP		WOCKHARDT	EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)	N78593 001	Feb 06, 2009	Jan	NEWA
----	--	-----------	-------------------------------------	------------	--------------	-----	------

>A> SUMAVEL DOSEPRO

>A>	+	ZOGENIX INC	EQ 6MG BASE/0.5ML (EQ 6MG BASE/0.5ML)	N22239 001	Jul 15, 2009	Jul	NEWA
-----	---	-------------	---------------------------------------	------------	--------------	-----	------

TABLET; ORAL

IMITREX

AB		GLAXOSMITHKLINE	EQ 25MG BASE	N20132 002	Jun 01, 1995	Jan	CFTG
----	--	-----------------	--------------	------------	--------------	-----	------

AB			EQ 50MG BASE	N20132 003	Jun 01, 1995	Jan	CFTG
----	--	--	--------------	------------	--------------	-----	------

AB	+		EQ 100MG BASE	N20132 001	Jun 01, 1995	Jan	CFTG
----	---	--	---------------	------------	--------------	-----	------

SUMATRIPTAN SUCCINATE

>A>	AB	AUROBINDO PHARMA	EQ 25MG BASE	N78327 001	Aug 10, 2009	Jul	NEWA
-----	----	------------------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N78327 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

>A>	AB		EQ 100MG BASE	N78327 003	Aug 10, 2009	Jul	NEWA
-----	----	--	---------------	------------	--------------	-----	------

>A>	AB	COBALT LABS INC	EQ 25MG BASE	N76933 001	Aug 10, 2009	Jul	NEWA
-----	----	-----------------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N76933 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

>A>	AB		EQ 100MG BASE	N76933 003	Aug 10, 2009	Jul	NEWA
-----	----	--	---------------	------------	--------------	-----	------

>A>	AB	DR REDDYS LABS INC	EQ 25MG BASE	N76847 001	Aug 10, 2009	Jul	NEWA
-----	----	--------------------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N76847 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

>A>	AB		EQ 100MG BASE	N76847 003	Aug 10, 2009	Jul	NEWA
-----	----	--	---------------	------------	--------------	-----	------

>A>	AB	MYLAN	EQ 25MG BASE	N77744 001	Aug 10, 2009	Jul	NEWA
-----	----	-------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N77744 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

>A>	AB		EQ 100MG BASE	N77744 003	Aug 10, 2009	Jul	NEWA
-----	----	--	---------------	------------	--------------	-----	------

>A>	AB	ORCHID HLTHCARE	EQ 25MG BASE	N78284 001	Aug 10, 2009	Jul	NEWA
-----	----	-----------------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N78284 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

>A>	AB		EQ 100MG BASE	N78284 003	Aug 10, 2009	Jul	NEWA
-----	----	--	---------------	------------	--------------	-----	------

>A>	AB	RANBAXY	EQ 25MG BASE	N76554 001	Aug 10, 2009	Jul	NEWA
-----	----	---------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N76554 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

	AB		EQ 100MG BASE	N76572 001	Feb 09, 2009	Jan	NEWA
--	----	--	---------------	------------	--------------	-----	------

>A>	AB	ROXANE	EQ 25MG BASE	N78241 001	Aug 10, 2009	Jul	NEWA
-----	----	--------	--------------	------------	--------------	-----	------

>A>	AB		EQ 50MG BASE	N78241 002	Aug 10, 2009	Jul	NEWA
-----	----	--	--------------	------------	--------------	-----	------

TABLET; ORAL

SUMATRIPTAN SUCCINATE

>A>	AB	ROXANE	EQ 100MG BASE	N78241 003	Aug 10, 2009	Jul	NEWA
>A>	AB	SANDOZ	EQ 25MG BASE	N76976 001	Aug 10, 2009	Jul	NEWA
>A>	AB		EQ 50MG BASE	N76976 002	Aug 10, 2009	Jul	NEWA
>A>	AB		EQ 100MG BASE	N76976 003	Aug 10, 2009	Jul	NEWA
>A>	AB	SUN PHARM INDS	EQ 25MG BASE	N78295 001	Aug 10, 2009	Jul	NEWA
>A>	AB		EQ 50MG BASE	N78295 002	Aug 10, 2009	Jul	NEWA
>A>	AB		EQ 100MG BASE	N78295 003	Aug 10, 2009	Jul	NEWA
	AB	TEVA	EQ 25MG BASE	N76840 001	Feb 09, 2009	Jan	NEWA
	AB		EQ 50MG BASE	N76840 002	Feb 09, 2009	Jan	NEWA
	AB		EQ 100MG BASE	N76840 003	Feb 09, 2009	Jan	NEWA

SUNITINIB MALATE

CAPSULE; ORAL

SUTENT

CPPI CV

EQ 37.5MG BASE	N21938 004	Mar 31, 2009	Mar	NEWA
----------------	------------	--------------	-----	------

TACROLIMUS

CAPSULE; ORAL

PROGRAF

>D>		ASTELLAS	EQ 0.5MG BASE	N50708 003	Aug 24, 1998	Jul	CFTG
>A>	AB		EQ 0.5MG BASE	N50708 003	Aug 24, 1998	Jul	CFTG
>D>			EQ 1MG BASE	N50708 001	Apr 08, 1994	Jul	CFTG
>A>	AB		EQ 1MG BASE	N50708 001	Apr 08, 1994	Jul	CFTG
>D>	+		EQ 5MG BASE	N50708 002	Apr 08, 1994	Jul	CFTG
>A>	AB	+	EQ 5MG BASE	N50708 002	Apr 08, 1994	Jul	CFTG
>A>		TACROLIMUS					
>A>	AB	SANDOZ	EQ 0.5MG BASE	N65461 001	Aug 10, 2009	Jul	NEWA
>A>	AB		EQ 1MG BASE	N65461 002	Aug 10, 2009	Jul	NEWA
>A>	AB		EQ 5MG BASE	N65461 003	Aug 10, 2009	Jul	NEWA

TADALAFIL

TABLET; ORAL

ADCIRCA

+ ELI LILLY CO	20MG	N22332 001	May 22, 2009	May	NEWA
----------------	------	------------	--------------	-----	------

TAMOXIFEN CITRATE

TABLET; ORAL

TAMOXIFEN CITRATE

@ ROXANE

EQ 10MG BASE	N76027 001	Feb 20, 2003	Mar	DISC
--------------	------------	--------------	-----	------

@

EQ 20MG BASE	N76027 002	Feb 20, 2003	Mar	DISC
--------------	------------	--------------	-----	------

TAPENTADOL HYDROCHLORIDE

TABLET; ORAL

TAPENTADOL HYDROCHLORIDE

ORTHO MCNEIL JANSSEN

EQ 50MG BASE	N22304 001	Nov 20, 2008	Jun	CPOT
EQ 75MG BASE	N22304 002	Nov 20, 2008	Jun	CPOT
+ EQ 100MG BASE	N22304 003	Nov 20, 2008	Jun	CPOT

TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION

NEUROLITE

LANTHEUS MEDCL

N/A	N20256 001	Nov 23, 1994	Apr	CPOT
-----	------------	--------------	-----	------

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

DRAXIMAGE MDP-10

	+	DRAXIMAGE	N/A	N18035 001		Apr	CTEC
--	---	-----------	-----	------------	--	-----	------

>D>		DRAXIMAGE MDP-25					
-----	--	------------------	--	--	--	--	--

>D>	+	DRAXIMAGE	N/A	N18035 002	Feb 27, 2004	Jul	DISC
-----	---	-----------	-----	------------	--------------	-----	------

>A>	@		N/A	N18035 002	Feb 27, 2004	Jul	DISC
-----	---	--	-----	------------	--------------	-----	------

	+		N/A	N18035 002	Feb 27, 2004	Jun	CMFD
--	---	--	-----	------------	--------------	-----	------

	@		N/A	N18035 002	Feb 27, 2004	Apr	DISC
--	---	--	-----	------------	--------------	-----	------

TECHNETIUM TC 99M MPI MDP

	@	GE HEALTHCARE	N/A	N18141 002	Jun 12, 1989	Apr	DISC
--	---	---------------	-----	------------	--------------	-----	------

	@		N/A	N18141 001		Apr	DISC
--	---	--	-----	------------	--	-----	------

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

TECHNETIUM TC 99M SESTAMIBI

AP		CARDINAL HEALTH 414	N/A	N78809 001	Apr 28, 2009	Apr	NEWA
----	--	---------------------	-----	------------	--------------	-----	------

AP		DRAXIMAGE	N/A	N78806 001	Apr 29, 2009	Apr	NEWA
----	--	-----------	-----	------------	--------------	-----	------

AP		PHARMALUCENCE	N/A/VIAL	N79157 001	Jul 10, 2009	Jun	NEWA
----	--	---------------	----------	------------	--------------	-----	------

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW 30ML

>D>	@	GE HEALTHCARE	N/A	N20372 002	Jul 07, 2005	Jul	CMFD
-----	---	---------------	-----	------------	--------------	-----	------

>A>	+		N/A	N20372 002	Jul 07, 2005	Jul	CMFD
-----	---	--	-----	------------	--------------	-----	------

TELIVUDINE

SOLUTION; ORAL

TYZEKA

	+	NOVARTIS	100MG/5ML	N22154 001	Apr 28, 2009	Apr	NEWA
--	---	----------	-----------	------------	--------------	-----	------

TEMAZEPAM

CAPSULE; ORAL

RESTORIL

AB		TYCO HLTHCARE	22.5MG	N18163 004	Nov 02, 2004	Jun	CFTG
----	--	---------------	--------	------------	--------------	-----	------

TEMAZEPAM

AB		MYLAN	22.5MG	N70920 003	Jun 12, 2009	Jun	NEWA
----	--	-------	--------	------------	--------------	-----	------

	@	NOVEL LABS INC	30MG	N71457 001	Apr 21, 1987	Mar	CAHN
--	---	----------------	------	------------	--------------	-----	------

TEMOZOLOMIDE

POWDER; INTRAVENOUS

TEMODAR

	+	SCHERING	100MG/VIAL	N22277 001	Feb 27, 2009	Feb	NEWA
--	---	----------	------------	------------	--------------	-----	------

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

AB		IVAX PHARMS	EQ 1MG BASE	N75614 002	Jan 30, 2001	May	CMFD
----	--	-------------	-------------	------------	--------------	-----	------

AB			EQ 2MG BASE	N75614 001	Jan 30, 2001	May	CMFD
----	--	--	-------------	------------	--------------	-----	------

AB			EQ 5MG BASE	N75614 003	Jan 30, 2001	May	CMFD
----	--	--	-------------	------------	--------------	-----	------

AB			EQ 10MG BASE	N75614 004	Jan 30, 2001	May	CMFD
----	--	--	--------------	------------	--------------	-----	------

TABLET; ORAL

HYTRIN

		ABBOTT	EQ 1MG BASE	N19057 001	Aug 07, 1987	Mar	CTEC
--	--	--------	-------------	------------	--------------	-----	------

	+		EQ 2MG BASE	N19057 002	Aug 07, 1987	Mar	CTEC
--	---	--	-------------	------------	--------------	-----	------

TABLET; ORAL									
HYTRIN									
ABBOTT				EQ 5MG BASE	N19057 003	Aug 07, 1987	Mar	CTEC	
				EQ 10MG BASE	N19057 004	Aug 07, 1987	Mar	CTEC	
TERAZOSIN HYDROCHLORIDE									
@ SANDOZ				EQ 1MG BASE	N74315 001	Dec 31, 1998	Mar	DISC	
@				EQ 2MG BASE	N74315 002	Dec 31, 1998	Mar	DISC	
@				EQ 5MG BASE	N74315 003	Dec 31, 1998	Mar	DISC	
@				EQ 10MG BASE	N74315 004	Dec 31, 1998	Mar	DISC	
TERBUTALINE SULFATE									
INJECTABLE; INJECTION									
TERBUTALINE SULFATE									
AP		HIKMA FARMACEUTICA	1MG/ML		N78630 001	May 20, 2009	May	NEWA	
TERIPARATIDE RECOMBINANT HUMAN									
INJECTABLE; SUBCUTANEOUS									
FORTEO									
LILLY				0.6MG/2.4ML (0.25MG/ML)	N21318 002	Jun 25, 2008	Feb	NEWA	
+				0.75MG/3ML (0.25MG/ML)	N21318 001	Nov 26, 2002	Feb	CPOT	
TESTOSTERONE									
GEL; TRANSDERMAL									
ANDROGEL									
UNIMED PHARMS				1% (2.5GM/PACKET)	N21015 001	Feb 28, 2000	Mar	CTEC	
BX	+			1% (5GM/PACKET)	N21015 002	Feb 28, 2000	Mar	CTEC	
TESTOSTERONE									
@ WATSON LABS				1% (2.5GM/PACKET)	N76737 001	Jan 27, 2006	Mar	DISC	
@				1% (5GM/PACKET)	N76737 002	Jan 27, 2006	Mar	DISC	
TESTOSTERONE ENANTHATE									
INJECTABLE; INJECTION									
DELATESTYL									
AO	+	ENDO PHARM	200MG/ML		N09165 003		Apr	CAHN	
		@	200MG/ML		N09165 001		Apr	CAHN	
AO	+	ENDO PHARMS	200MG/ML		N09165 003		Mar	CAHN	
		@	200MG/ML		N09165 001		Mar	CAHN	
TETRAHYDROZOLINE HYDROCHLORIDE									
SOLUTION; NASAL									
TYZINE									
+ NYCOMED US				0.05%	N86576 002		Apr	CAHN	
				0.1%	N86576 001		Apr	CAHN	
SPRAY; NASAL									
TYZINE									
+ NYCOMED US				0.1%	N86576 003		Apr	CAHN	
THEOPHYLLINE									
ELIXIR; ORAL									
ELIXOPHYLLIN									
+ FOREST LABS				80MG/15ML	N85186 001		Jun	CRLD	
THEOPHYLLINE									
@ MORTON GROVE				80MG/15ML	N86748 001		Jun	DISC	
@ PRECISION DOSE				80MG/15ML	N85863 001		Jun	DISC	
AA				80MG/15ML	N85863 001		Apr	CAHN	
@ TARO				80MG/15ML	N89626 001	Oct 28, 1988	Jun	DISC	

SOLUTION; ORAL
THEOPHYLLINE
@ ROXANE

80MG/15ML

N87449 001 Sep 15, 1983 Mar DISC

THIABENDAZOLE

TABLET, CHEWABLE; ORAL
MINTEZOL
@ MERCK

500MG

N16096 001 Jun DISC

THIOTHIXENE

CAPSULE; ORAL
NAVANE
@ PFIZER

20MG

N16584 005 Jun DISC

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL
TICLID

@ ROCHE PALO

250MG

N19979 002 Oct 31, 1991 Jun DISC

TICLOPIDINE HYDROCHLORIDE

AB + TEVA 250MG

N75149 001 Aug 20, 1999 Jun CRLD

TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC
TIMOPTIC

AT + ATON EQ 0.25% BASE

N18086 001 Feb CAHN

AT + EQ 0.5% BASE

N18086 002 Feb CAHN

TIMOPTIC IN OCUDOSE

+ ATON EQ 0.25% BASE

N19463 001 Nov 05, 1986 Feb CAHN

+ EQ 0.5% BASE

N19463 002 Nov 05, 1986 Feb CAHN

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC
TIMOPTIC-XE

AB + ATON EQ 0.25% BASE

N20330 001 Nov 04, 1993 Feb CAHN

AB + EQ 0.5% BASE

N20330 002 Nov 04, 1993 Feb CAHN

TABLET; ORAL

TIMOLOL MALEATE

MYLAN

5MG

N72666 001 Jun 08, 1990 Mar CTEC

10MG

N72667 001 Jun 08, 1990 Mar CTEC

+ 20MG

N72668 001 Jun 08, 1990 Mar CTEC

@ SANDOZ

5MG

N72550 001 Apr 13, 1989 Mar DISC

@

10MG

N72551 001 Apr 13, 1989 Mar DISC

@

20MG

N72552 001 Apr 13, 1989 Mar DISC

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

IVAX PHARMS

100MG

N18894 001 Nov 02, 1984 Mar CTEC

>D> IVAX SUB TEVA PHARMS 100MG

N18894 001 Nov 02, 1984 Jul DISC

>A> @ 100MG

N18894 001 Nov 02, 1984 Jul DISC

>D> AB 250MG

N18894 002 Nov 02, 1984 Jul DISC

>A> @ 250MG

N18894 002 Nov 02, 1984 Jul DISC

>D> AB + 500MG

N18894 003 Nov 02, 1984 Jul DISC

>A> @ 500MG

N18894 003 Nov 02, 1984 Jul DISC

>D> AB MYLAN 500MG

N70913 001 Mar 17, 1986 Jul CRLD

>A> AB + 500MG

N70913 001 Mar 17, 1986 Jul CRLD

@ SANDOZ

100MG

N71633 001 Dec 09, 1987 Mar DISC

@

250MG

N70289 001 Mar 13, 1986 Mar DISC

TABLET; ORAL									
TOLAZAMIDE									
@ SANDOZ	500MG			N70290 001	Mar 13, 1986	Mar	DISC		
<u>TOLBUTAMIDE</u>									
TABLET; ORAL									
TOLBUTAMIDE									
@ SANDOZ	500MG			N86574 001		Mar	DISC		
<u>TOLMETIN SODIUM</u>									
TABLET; ORAL									
TOLMETIN SODIUM									
@ SANDOZ	EQ 200MG BASE			N73588 001	Jul 31, 1992	Mar	DISC		
@	EQ 600MG BASE			N74002 001	Sep 27, 1993	Mar	DISC		
<u>TOLVAPTAN</u>									
TABLET; ORAL									
SAMSCA									
OTSUKA AMERICA PHARM	15MG			N22275 001	May 19, 2009	May	NEWA		
+	30MG			N22275 002	May 19, 2009	May	NEWA		
@	60MG			N22275 003	May 19, 2009	May	DISC		
<u>TOPIRAMATE</u>									
CAPSULE; ORAL									
TOPAMAX									
AB	ORTHO MCNEIL JANSSEN	15MG		N20844 001	Oct 26, 1998	Mar	CFTG		
AB	+	25MG		N20844 002	Oct 26, 1998	Mar	CFTG		
TOPIRAMATE									
AB	BARR	15MG		N76448 001	Apr 15, 2009	Mar	NEWA		
AB		25MG		N76448 002	Apr 15, 2009	Mar	NEWA		
AB	COBALT LABS INC	15MG		N77868 001	Apr 15, 2009	Mar	NEWA		
AB		25MG		N77868 002	Apr 15, 2009	Mar	NEWA		
AB	TEVA	15MG		N76575 001	Apr 17, 2009	Mar	NEWA		
AB		25MG		N76575 002	Apr 17, 2009	Mar	NEWA		
TABLET; ORAL									
TOPAMAX									
AB	+	ORTHO MCNEIL JANSSEN	25MG	N20505 004	Dec 24, 1996	Mar	CFTG		
AB			50MG	N20505 005	Dec 24, 1996	Mar	CFTG		
AB			100MG	N20505 001	Dec 24, 1996	Mar	CFTG		
AB			200MG	N20505 002	Dec 24, 1996	Mar	CFTG		
TOPIRAMATE									
AB	APOTEX INC	25MG		N77733 001	Mar 27, 2009	Mar	NEWA		
AB		50MG		N77733 002	Mar 27, 2009	Mar	NEWA		
AB		100MG		N77733 003	Mar 27, 2009	Mar	NEWA		
AB		200MG		N77733 004	Mar 27, 2009	Mar	NEWA		
AB	AUROBINDO PHARMA	25MG		N78462 001	Mar 27, 2009	Mar	NEWA		
AB		50MG		N78462 002	Mar 27, 2009	Mar	NEWA		
AB		100MG		N78462 003	Mar 27, 2009	Mar	NEWA		
AB		200MG		N78462 004	Mar 27, 2009	Mar	NEWA		
AB	BARR	25MG		N76315 001	Mar 27, 2009	Mar	NEWA		
AB		100MG		N76315 002	Mar 27, 2009	Mar	NEWA		
AB		200MG		N76315 003	Mar 27, 2009	Mar	NEWA		
AB	CIPLA LTD	25MG		N76343 001	Mar 27, 2009	Mar	NEWA		
AB		50MG		N76343 002	Mar 27, 2009	Mar	NEWA		
AB		100MG		N76343 003	Mar 27, 2009	Mar	NEWA		
AB		200MG		N76343 004	Mar 27, 2009	Mar	NEWA		

TABLET; ORAL

TOPIRAMATE

AB	COBALT LABS INC	25MG	N77643 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N77643 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N77643 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N77643 004	Mar 27, 2009	Mar	NEWA
AB	GLENMARK GENERICS	25MG	N77627 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N77627 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N77627 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N77627 004	Mar 27, 2009	Mar	NEWA
AB	INVAGEN PHARMS	25MG	N79162 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N79162 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N79162 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N79162 004	Mar 27, 2009	Mar	NEWA
AB	MYLAN	25MG	N76314 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N76314 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N76314 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N76314 004	Mar 27, 2009	Mar	NEWA
AB	PAR PHARM	25MG	N76311 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N76311 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N76311 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N76311 004	Mar 27, 2009	Mar	NEWA
AB	PLIVA HRVATSKA DOO	25MG	N77905 001	Mar 30, 2009	Mar	NEWA
AB		50MG	N77905 002	Mar 30, 2009	Mar	NEWA
AB		100MG	N77905 003	Mar 30, 2009	Mar	NEWA
AB		200MG	N77905 004	Mar 30, 2009	Mar	NEWA
AB	RANBAXY	25MG	N76327 001	Mar 27, 2009	Mar	NEWA
AB		100MG	N76327 002	Mar 27, 2009	Mar	NEWA
AB		200MG	N76327 003	Mar 27, 2009	Mar	NEWA
AB	ROXANE	25MG	N76306 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N76306 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N76306 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N76306 004	Mar 27, 2009	Mar	NEWA
AB	SUN PHARM INDS LTD	25MG	N90278 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N90278 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N90278 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N90278 004	Mar 27, 2009	Mar	NEWA
AB	TEVA	25MG	N76317 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N76317 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N76317 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N76317 004	Mar 27, 2009	Mar	NEWA
AB	TORRENT PHARMS	25MG	N79153 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N79153 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N79153 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N79153 004	Mar 27, 2009	Mar	NEWA
AB	UNICHEM	25MG	N90162 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N90162 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N90162 003	Mar 27, 2009	Mar	NEWA
AB	ZYDUS PHARMS USA INC	25MG	N78235 001	Mar 27, 2009	Mar	NEWA
AB		50MG	N78235 002	Mar 27, 2009	Mar	NEWA
AB		100MG	N78235 003	Mar 27, 2009	Mar	NEWA
AB		200MG	N78235 004	Mar 27, 2009	Mar	NEWA

TORSEMIDE

TABLET; ORAL

TORSEMIDE

AB	HETERO DRUGS	5MG	N79234 001	Jan 27, 2009	Jan	NEWA
AB		10MG	N79234 002	Jan 27, 2009	Jan	NEWA
AB		20MG	N79234 003	Jan 27, 2009	Jan	NEWA
AB		100MG	N79234 004	Jan 27, 2009	Jan	NEWA

TRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

RYZOLT

BC	+	PURDUE PHARMA	100MG	N21745 001	Dec 30, 2008	Mar	CAHN
BC			200MG	N21745 002	Dec 30, 2008	Mar	CAHN
BC			300MG	N21745 003	Dec 30, 2008	Mar	CAHN

TABLET, ORALLY DISINTEGRATING; ORAL

TRAMADOL HYDROCHLORIDE

@ ETHYPHARM NORTH

50MG

N21693 001 May 05, 2005 Mar CAHN

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

AB	ALVOGEN	50MG	N71636 001	Apr 18, 1988	Feb	CAHN
AB		100MG	N71514 001	Apr 18, 1988	Feb	CAHN
AB	APOTEX INC	300MG	N71196 003	Apr 26, 1999	May	CTEC
AB	MATRIX LABS LTD	50MG	N90514 001	Jun 02, 2009	May	NEWA
AB		100MG	N90514 002	Jun 02, 2009	May	NEWA
AB		150MG	N90514 003	Jun 02, 2009	May	NEWA
AB		300MG	N90514 004	Jun 02, 2009	May	NEWA
	@ SANDOZ	50MG	N72484 001	Apr 30, 1990	Mar	DISC
	@	100MG	N72483 001	Apr 30, 1990	Mar	DISC

TREPROSTINIL SODIUM

>A> SOLUTION; INHALATION

>A> TYVASO

>A>	+	UNITED THERAP	EQ 0.6MG BASE/ML	N22387 001	Jul 30, 2009	Jul	NEWA
-----	---	---------------	------------------	------------	--------------	-----	------

TRETINOIN

CREAM; TOPICAL

RENOVA

>D>	+	JOHNSON AND JOHNSON	0.02%	N21108 001	Aug 31, 2000	Jul	CAHN
>D>	AB2	+	0.05%	N19963 001	Dec 29, 1995	Jul	CAHN
>A>	+	ORTHO DERMATOLOGICS	0.02%	N21108 001	Aug 31, 2000	Jul	CAHN
>A>	AB2	+	0.05%	N19963 001	Dec 29, 1995	Jul	CAHN

GEL; TOPICAL

ATRALIN

+ DOW PHARM SCIENCES 0.05%

N22070 001 Jul 26, 2007 Jun CAHN

RETIN-A MICRO

>D>	+	JOHNSON AND JOHNSON	0.04%	N20475 002	May 10, 2002	Jul	CAHN
>D>	+		0.1%	N20475 001	Feb 07, 1997	Jul	CAHN
>A>	+	ORTHO DERMATOLOGICS	0.04%	N20475 002	May 10, 2002	Jul	CAHN
>A>	+		0.1%	N20475 001	Feb 07, 1997	Jul	CAHN

TRIAMCINOLONE ACETONIDE

INJECTABLE; INJECTION

KENALOG-10

AB		APOTHECON	10MG/ML	N12041 001		May	CFTG
----	--	-----------	---------	------------	--	-----	------

KENALOG-40

AB	+	APOTHECON	40MG/ML	N14901 001		May	CFTG
----	---	-----------	---------	------------	--	-----	------

TRIAMCINOLONE ACETONIDE

AB		SANDOZ	10MG/ML	N90166 001	May 27, 2009	May	NEWA
----	--	--------	---------	------------	--------------	-----	------

AB			40MG/ML	N90164 001	May 27, 2009	May	NEWA
----	--	--	---------	------------	--------------	-----	------

SPRAY, METERED; NASAL

NASACORT AQ

>D>	+	SANOFI AVENTIS US	0.055MG/SPRAY	N20468 001	May 20, 1996	Jul	CFTG
-----	---	-------------------	---------------	------------	--------------	-----	------

>A>	AB	+		N20468 001	May 20, 1996	Jul	CFTG
-----	----	---	--	------------	--------------	-----	------

>A>		TRIAMCINOLONE ACETONIDE					
-----	--	-------------------------	--	--	--	--	--

>A>	AB	BARR	0.055MG/SPRAY	N78104 001	Jul 30, 2009	Jul	NEWA
-----	----	------	---------------	------------	--------------	-----	------

TROSPIMUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

SANCTURA XR

+	ALLERGAN	60MG	N22103 001	Aug 03, 2007	Apr	CAHN
---	----------	------	------------	--------------	-----	------

+	ENDO PHARMS	60MG	N22103 001	Aug 03, 2007	Mar	CAHN
---	-------------	------	------------	--------------	-----	------

TABLET; ORAL

SANCTURA

+	ALLERGAN	20MG	N21595 001	May 28, 2004	Apr	CAHN
---	----------	------	------------	--------------	-----	------

+	ENDO PHARMS	20MG	N21595 001	May 28, 2004	Mar	CAHN
---	-------------	------	------------	--------------	-----	------

TRYPAN BLUE

SOLUTION; OPHTHALMIC

MEMBRANEBLUE

+	DORC	0.15%	N22278 001	Feb 20, 2009	Feb	NEWA
---	------	-------	------------	--------------	-----	------

URSODIOL

TABLET; ORAL

URSO 250

AB		AXCAN	250MG	N20675 001	Dec 10, 1997	May	CFTG
----	--	-------	-------	------------	--------------	-----	------

URSO FORTE

AB	+	AXCAN	500MG	N20675 002	Jul 21, 2004	May	CFTG
----	---	-------	-------	------------	--------------	-----	------

URSODIOL

AB		TEVA PHARMS	250MG	N79184 001	May 13, 2009	May	NEWA
----	--	-------------	-------	------------	--------------	-----	------

AB			500MG	N79184 002	May 13, 2009	May	NEWA
----	--	--	-------	------------	--------------	-----	------

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

+	ENDO PHARM	40MG/ML	N20892 001	Sep 25, 1998	Apr	CAHN
---	------------	---------	------------	--------------	-----	------

+	ENDO PHARMS	40MG/ML	N20892 001	Sep 25, 1998	Mar	CAHN
---	-------------	---------	------------	--------------	-----	------

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

AP	+	ABRAXIS PHARM	EQ 10GM BASE/VIAL	N62663 004	Nov 28, 1997	Apr	CTEC
----	---	---------------	-------------------	------------	--------------	-----	------

AP		HOSPIRA	EQ 750MG BASE/VIAL	N62912 002	Jan 07, 2009	May	CTEC
----	--	---------	--------------------	------------	--------------	-----	------

AP			EQ 750MG BASE/VIAL	N62933 002	May 27, 2009	May	NEWA
----	--	--	--------------------	------------	--------------	-----	------

AP		HOSPIRA INC	EQ 10GM BASE/VIAL	N65455 001	Apr 29, 2009	Apr	NEWA
----	--	-------------	-------------------	------------	--------------	-----	------

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

AP	SUN PHARMA GLOBAL	10MG/VIAL	N79001 001	Jun 17, 2009	Jun	NEWA
AP		20MG/VIAL	N79001 002	Jun 17, 2009	Jun	NEWA

VERAPAMIL HYDROCHLORIDE

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

@ BEDFORD 2.5MG/ML

N72888 001 Jul 28, 1995 Apr DISC

TABLET; ORAL

VERAPAMIL HYDROCHLORIDE

@ SANDOZ 40MG

N73168 001 Jul 31, 1992 Mar DISC

@ 80MG

N71423 001 May 24, 1988 Mar DISC

@ 120MG

N71424 001 May 25, 1988 Mar DISC

VINORELBINE TARTRATE

INJECTABLE; INJECTION

VINORELBINE TARTRATE

>A>	AP	ACTAVIS TOTOWA	EQ 10MG BASE/ML	N78011 001	Jul 22, 2009	Jul	NEWA
-----	----	----------------	-----------------	------------	--------------	-----	------

ZIDOVUDINE

TABLET; ORAL

ZIDOVUDINE

>A>		@ AUROBINDO PHARMA	60MG	N22294 001	Jul 23, 2009	Jul	DISC
-----	--	--------------------	------	------------	--------------	-----	------

ZOLPIDEM TARTRATE

TABLET; SUBLINGUAL

EDLUAR

>A>		MEDA PHARMS	5MG	N21997 001	Mar 13, 2009	Jul	CAHN
-----	--	-------------	-----	------------	--------------	-----	------

>A>	+		10MG	N21997 002	Mar 13, 2009	Jul	CAHN
-----	---	--	------	------------	--------------	-----	------

>D>		OREXO AB	5MG	N21997 001	Mar 13, 2009	Jul	CAHN
-----	--	----------	-----	------------	--------------	-----	------

			5MG	N21997 001	Mar 13, 2009	Jun	CAHN
--	--	--	-----	------------	--------------	-----	------

			5MG	N21997 001	Mar 13, 2009	Mar	NEWA
--	--	--	-----	------------	--------------	-----	------

>D>	+		10MG	N21997 002	Mar 13, 2009	Jul	CAHN
-----	---	--	------	------------	--------------	-----	------

	+		10MG	N21997 002	Mar 13, 2009	Jun	CAHN
--	---	--	------	------------	--------------	-----	------

	+		10MG	N21997 002	Mar 13, 2009	Mar	NEWA
--	---	--	------	------------	--------------	-----	------

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

@ MUTUAL PHARM 25MG

N77635 001 Dec 22, 2005 May DISC

@ 50MG

N77635 002 Dec 22, 2005 May DISC

@ 100MG

N77635 003 Dec 22, 2005 May DISC

OTC DRUG PRODUCT LIST - 29TH EDITION

OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

2-1

ACETAMINOPHEN

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

OHM LABS	650MG	N76200 001	Mar 19, 2002	Mar	CAHN
----------	-------	------------	--------------	-----	------

CETIRIZINE HYDROCHLORIDE

CAPSULE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

>A>	BANNER PHARMACAPS	5MG	N22429 001	Jul 23, 2009	Jul	NEWA
>A>		5MG	N22429 003	Jul 23, 2009	Jul	NEWA

CETIRIZINE HYDROCHLORIDE HIVES

>A>	+ BANNER PHARMACAPS	10MG	N22429 002	Jul 23, 2009	Jul	NEWA
>A>		10MG	N22429 004	Jul 23, 2009	Jul	NEWA

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

DR REDDYS LABS LTD	5MG/5ML	N90474 002	Mar 30, 2009	Mar	NEWA
--------------------	---------	------------	--------------	-----	------

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

DR REDDYS LABS LTD	5MG/5ML	N90474 001	Mar 30, 2009	Mar	NEWA
--------------------	---------	------------	--------------	-----	------

TABLET, CHEWABLE; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

>A>	CARACO	5MG	N77631 004	Jan 11, 2008	Jul	NEWA
>A>		10MG	N77631 003	Jan 11, 2008	Jul	NEWA

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

>A>	CARACO	5MG	N77631 001	Jan 11, 2008	Jul	CDFR
>A>		10MG	N77631 002	Jan 11, 2008	Jul	CDFR

TABLET; ORAL

CETIRIZINE HYDROCHLORIDE

>D>	CARACO	5MG	N77631 001	Jan 11, 2008	Jul	CDFR
>D>		10MG	N77631 002	Jan 11, 2008	Jul	CDFR

CETIRIZINE HYDROCHLORIDE ALLERGY

	ORCHID HLTHCARE	5MG	N78862 001	Feb 19, 2009	Feb	NEWA
		10MG	N78862 002	Feb 19, 2009	Feb	NEWA
>A>	TARO	5MG	N78072 003	Jul 22, 2009	Jul	NEWA
>A>		5MG	N78072 001	Jul 22, 2009	Jul	NEWA
	UNICHEM	5MG	N78680 003	Jun 26, 2009	Jun	NEWA
		10MG	N78680 004	Jun 26, 2009	Jun	NEWA

CETIRIZINE HYDROCHLORIDE HIVES

	ORCHID HLTHCARE	5MG	N78862 003	Feb 19, 2009	Feb	NEWA
		10MG	N78862 004	Feb 19, 2009	Feb	NEWA
	UNICHEM	5MG	N78680 001	Jun 26, 2009	Jun	NEWA
		10MG	N78680 002	Jun 26, 2009	Jun	NEWA

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

>A>	TARO	10MG	N78072 002	Jul 22, 2009	Jul	NEWA
>A>		10MG	N78072 004	Jul 22, 2009	Jul	NEWA

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE

AVANTHI INC	12MG	N40829 001	May 13, 2009	Apr	NEWA
-------------	------	------------	--------------	-----	------

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

DELSYM

+	RECKITT BENCKISER	EQ 30MG HBR/5ML	N18658 001	Oct 08, 1982	Feb	CAHN
---	-------------------	-----------------	------------	--------------	-----	------

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET; ORAL

IBUPROFEN AND DIPHENHYDRAMINE CITRATE

DR REDDYS LABS LTD 38MG;200MG

N90619 001	Jul 08, 2009	Jun	NEWA
------------	--------------	-----	------

EPINEPHRINE

AEROSOL, METERED; INHALATION

PRIMATENE MIST

@ WYETH CONS 0.2MG/INH

N16126 001		Jun	DISC
------------	--	-----	------

EPINEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

BRONITIN MIST

@ WYETH CONS 0.3MG/INH

N16126 002		Jun	DISC
------------	--	-----	------

IBUPROFEN

CAPSULE; ORAL

IBUPROFEN

BANNER PHARMACAPS EQ 200MG FREE ACID AND POTASSIUM SALT

N78682 001	Mar 24, 2009	Mar	NEWA
------------	--------------	-----	------

>A>	DR REDDYS LABS LTD	EQ 200MG FREE ACID AND POTASSIUM SALT	N77338 001	Jul 10, 2009	Jul	NEWA
-----	--------------------	---------------------------------------	------------	--------------	-----	------

>A>		EQ 200MG FREE ACID AND POTASSIUM SALT	N77338 001	Jul 10, 2009	Jul	NEWA
-----	--	---------------------------------------	------------	--------------	-----	------

	MARKSANS PHARMA	EQ 200MG FREE ACID AND POTASSIUM SALT	N77338 001	Jul 10, 2009	Jun	NEWA
		EQ 200MG FREE ACID AND POTASSIUM SALT	N79205 001	Jun 26, 2009	Jun	NEWA

MIDOL LIQUID GELS

+ BANNER PHARMACAPS 200MG

N21472 001	Oct 18, 2002	Feb	CTNA
------------	--------------	-----	------

TABLET; ORAL

IBUPROFEN

@ SANDOZ 200MG

N70733 001	Sep 19, 1986	Mar	DISC
------------	--------------	-----	------

MEDIPREN

@ MCNEIL 200MG

N70475 001	Feb 06, 1986	Apr	DISC
------------	--------------	-----	------

@ 200MG

N71215 001	Jun 26, 1986	Apr	DISC
------------	--------------	-----	------

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN 50/50

@ LILLY 50 UNITS/ML;50 UNITS/ML

N20100 001	Apr 29, 1992	Jan	DISC
------------	--------------	-----	------

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

PREVACID 24 HR

+ NOVARTIS 15MG

N22327 001	May 18, 2009	May	NEWA
------------	--------------	-----	------

LEVONORGESTREL

TABLET; ORAL

PLAN B ONE-STEP

>A>						
>A>	+	DURAMED	1.5MG			

N21998 001	Jul 10, 2009	Jul	NEWA
------------	--------------	-----	------

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

PROSTEP

@ AVEVA 11MG/24HR

N19983 003 Dec 23, 1998 Jun DISC

@ 22MG/24HR

N19983 004 Dec 23, 1998 Jun DISC

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICOTINE POLACRILEX

IVAX PHARMS EQ 2MG BASE

N76880 001 Feb 18, 2009 Feb NEWA

EQ 4MG BASE

N77850 001 Feb 18, 2009 Feb NEWA

WATSON LABS EQ 2MG BASE

N79044 001 Jul 08, 2009 Jun NEWA

EQ 2MG BASE

N79216 001 Jul 08, 2009 Jun NEWA

EQ 4MG BASE

N79038 001 Jul 08, 2009 Jun NEWA

EQ 4MG BASE

N79219 001 Jul 08, 2009 Jun NEWA

TROCHE/LOZENGE; ORAL

NICORETTE

GLAXOSMITHKLINE CONS EQ 2MG BASE

N22360 001 May 18, 2009 May NEWA

+ EQ 4MG BASE

N22360 002 May 18, 2009 May NEWA

NICOTINE POLACRILEX

PERRIGO R AND D EQ 2MG BASE

N90711 001 Jul 10, 2009 Jun NEWA

EQ 2MG BASE

N90821 001 Jul 10, 2009 Jun NEWA

EQ 4MG BASE

N90711 002 Jul 10, 2009 Jun NEWA

EQ 4MG BASE

N90821 002 Jul 10, 2009 Jun NEWA

POTASSIUM IODIDE

SOLUTION; ORAL

>D> POTASSIUM IODIDE

>D> @ ROXANE 1GM/ML

N18551 001 Feb 19, 1982 Jul DISC

>A> @ 1GM/ML

N18551 001 Feb 19, 1982 Jul DISC

THYROSHIELD

+ FLEMING 65MG/ML

N77218 001 Jan 12, 2005 Apr CRLD

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE

>D> RANBAXY EQ 75MG BASE

N75132 001 Jan 14, 2000 Jul DISC

>A> @ EQ 75MG BASE

N75132 001 Jan 14, 2000 Jul DISC

@ SANDOZ EQ 75MG BASE

N75519 001 Sep 26, 2002 Mar DISC

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

CUMULATIVE SUPPLEMENT NUMBER 07 JULY 2009

NO JULY 2009 APPROVALS

ORPHAN PRODUCT DESIGNATIONS AND APPROVALS LIST

The list of List of Orphan Designations and Approvals is available at:

<http://www.fda.gov/orphan/designat/list.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

NO JULY 2009 ADDITIONS

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST - 29TH EDITION

A - 1

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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>						
022320 001	4717720	May 31, 2010	DS DP			
	RE34440	May 31, 2010	U-818			
<u>ALBUTEROL SULFATE - VENTOLIN HFA</u>						
020983 001	6161724	Jan 16, 2018	DP			
	6161724*PED	Jul 16, 2018				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6431168	Jun 08, 2018	DP			
	6431168*PED	Dec 08, 2018				
	6435372	Jan 16, 2018	DP			
	6435372*PED	Jul 16, 2018				
	6596260	Aug 10, 2014	DP			
	6596260*PED	Feb 10, 2015				
	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>ALISKIREN FUMARATE; HYDROCHLOROTHIAZIDE - TEKTURN HCT</u>						
022107 001					>A> I-600	Jul 16, 2012
<u>ALISKIREN FUMARATE; HYDROCHLOROTHIAZIDE - TEKTURN HCT</u>						
022107 002					>A> I-600	Jul 16, 2012
<u>ALISKIREN FUMARATE; HYDROCHLOROTHIAZIDE - TEKTURN HCT</u>						
022107 003					>A> I-600	Jul 16, 2012
<u>ALISKIREN FUMARATE; HYDROCHLOROTHIAZIDE - TEKTURN HCT</u>						
022107 004					>A> I-600	Jul 16, 2012
<u>ALMOTRIPTAN MALATE - AXERT</u>						
021001 001	5565447	May 07, 2015	DS DP U-969			
	5565447*PED	Nov 07, 2015				
<u>ALMOTRIPTAN MALATE - AXERT</u>						
021001 002	5565447	May 07, 2015	DS DP U-969			
	5565447*PED	Nov 07, 2015				
<u>ALPRAZOLAM - ALPRAZOLAM</u>						
078088 001					PC	Jul 13, 2009
<u>ALPRAZOLAM - ALPRAZOLAM</u>						
078088 002					PC	Jul 13, 2009

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALPRAZOLAM - ALPRAZOLAM</u>						
078088 003					PC	Jul 13, 2009
<u>ALPRAZOLAM - ALPRAZOLAM</u>						
078088 004					PC	Jul 13, 2009
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u>						
022325 001	5134127	Jan 23, 2010	DP			
	5376645	Jan 23, 2010	DP			
	6869939	May 04, 2022	DP			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 001	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 002	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 003	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 004	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 005	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 006	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 007	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 008	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 009	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 010	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u>						
021540 011	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMLODIPINE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
022314 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; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
022314 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; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
022314 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; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
022314 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; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u>						
022314 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>AMMONIA, N-13 - AMMONIA N 13</u>						
022119 001					NCE W	Aug 23, 2012 Aug 23, 2012
<u>AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE - PREVPAC</u>						
050757 001	4628098	May 10, 2009	DS			
	4628098*PED	Nov 10, 2009				
	5013743	Feb 12, 2010			U-452	
	5013743*PED	Aug 12, 2010				
	5045321	Sep 03, 2008	DP			
	5045321*PED	Mar 03, 2009				
	5093132	Sep 03, 2008	DP			
	5093132*PED	Mar 03, 2009				
	5433959	Sep 03, 2008	DP			
	5433959*PED	Mar 03, 2009				
<u>ANASTROZOLE - ARIMIDEX</u>						
020541 001	RE36617	Dec 27, 2009	DS DP U-946			
<u>ANIDULAFUNGIN - ERAXIS</u>						
021632 001	5965525	Feb 17, 2020	DS DP U-540			
<u>ANIDULAFUNGIN - ERAXIS</u>						
021632 002	5965525	Feb 17, 2020	DS DP U-540			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARIPIRAZOLE - ABILIFY</u>						
021866 001	>A> 7550445	Jul 21, 2024	DP			
	>A> 7550445*PED	Jan 21, 2025				
<u>ARMODAFINIL - NUVIGIL</u>						
021875 001	4927855	Apr 22, 2010	DS DP U-820			
	4927855*PED	Oct 22, 2010				
<u>ARMODAFINIL - NUVIGIL</u>						
021875 002	4927855	Apr 22, 2010	DS DP U-820		NP	Jun 15, 2010
	4927855*PED	Oct 22, 2010				
	7297346	Nov 29, 2023	DP			
	7297346*PED	May 29, 2024				
<u>ARMODAFINIL - NUVIGIL</u>						
021875 003	4927855	Apr 22, 2010	DS DP U-820			
	4927855*PED	Oct 22, 2010				
<u>ARMODAFINIL - NUVIGIL</u>						
021875 004	4927855	Apr 22, 2010	DS DP U-820			
	4927855*PED	Oct 22, 2010				
<u>ARMODAFINIL - NUVIGIL</u>						
021875 005	4927855	Apr 22, 2010	DS DP U-820		NP	Jun 15, 2010
	4927855*PED	Oct 22, 2010				
	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>ARSENIC TRIOXIDE - TRISENOX</u>						
021248 001	6982096	Nov 10, 2018	U-651			
<u>ARTEMETHER; LUMEFANTRINE - COARTEM</u>						
022268 001	5677331	Oct 14, 2014	DP U-977		NCE ODE	Apr 07, 2014 Apr 07, 2016
<u>ATORVASTATIN CALCIUM - LIPITOR</u>						
020702 001	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>ATORVASTATIN CALCIUM - LIPITOR</u>						
020702 002	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>ATORVASTATIN CALCIUM - LIPITOR</u>						
020702 003	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				
<u>ATORVASTATIN CALCIUM - LIPITOR</u>						
020702 004	RE40667	Dec 28, 2010	DS DP U-162			
	RE40667*PED	Jun 28, 2011				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ATOVAQUONE; PROGUANIL HYDROCHLORIDE - MALARONE</u>						
021078 001	>A> 5998449	Nov 25, 2013	U-990			
	>A> 5998449*PED	May 25, 2014				
<u>ATOVAQUONE; PROGUANIL HYDROCHLORIDE - MALARONE PEDIATRIC</u>						
021078 002	>A> 5998449	Nov 25, 2013	U-990			
	>A> 5998449*PED	May 25, 2014				
<u>AZITHROMYCIN - AZASITE</u>						
050810 001	5192535	Mar 09, 2010	DP U-709			
	>A> 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>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u>						
022249 002					I-580 NCE ODE	Oct 31, 2011 Mar 20, 2013 Mar 20, 2015
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u>						
050819 001	5733886	Mar 31, 2015	DP U-124			
	6117843	Feb 18, 2012	DP			
<u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - DUAC</u>						
050741 001	5466446	Feb 16, 2014	DS DP			
<u>BENZYL ALCOHOL - ULESFIA</u>						
022129 001	5858383	Aug 11, 2017	U-970		NCE	Apr 09, 2014
	6139859	Aug 11, 2017	U-970			
	6793931	Jul 11, 2022	DP U-970			
	7294342	May 19, 2024	U-970			
<u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u>						
022308 001	5447926	Sep 05, 2012	DS DP U-80		NCE	May 28, 2014
	6685958	Jun 29, 2021	DP U-80			
	6699492	Mar 31, 2019	DP U-80			
<u>BETAMETHASONE VALERATE - LUXIQ</u>						
020934 001	7078058	May 24, 2017	DP			
<u>BIMATOPROST - LATISSE</u>						
022369 001					NP	Dec 24, 2011
<u>BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE - PYLERA</u>						
050786 001	5196205	Mar 23, 2010	U-933			
	5476669	Mar 23, 2010	U-933			
	6350468	Dec 14, 2018	U-956			
	6350468	Dec 14, 2018	U-932			
<u>BIVALIRUDIN - ANGIOMAX</u>						
020873 001	5196404	Mar 23, 2010				
	5196404*PED	Sep 23, 2010				
<u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u>						
021398 001	>A> 7323463	Jan 19, 2023	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BROMOCRIPTINE MESYLATE - CYCLOSET</u>						
020866 001	5468755	Nov 21, 2012	U-976		NP	May 05, 2012
	5679685	Oct 21, 2014	DP			
	5716957	Feb 10, 2015	U-976			
	5756513	Nov 21, 2012	U-976			
	5866584	Nov 21, 2012	U-976			
<u>BUDESONIDE - PULMICORT RESPULES</u>						
020929 001	7524834	Nov 11, 2018	DP U-966			
	7524834*PED	May 11, 2019				
<u>BUDESONIDE - PULMICORT RESPULES</u>						
020929 002	7524834	Nov 11, 2018	DP U-966			
	7524834*PED	May 11, 2019				
<u>BUDESONIDE - PULMICORT RESPULES</u>						
020929 003	7524834	Nov 11, 2018	DP U-966			
	7524834*PED	May 11, 2019				
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
021929 001					I-582	Feb 27, 2012
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>						
021929 002					I-582	Feb 27, 2012
<u>CALCITONIN SALMON RECOMBINANT - FORTICAL</u>						
021406 001	RE40812	Feb 02, 2021	DP			
<u>CALCITRIOL - VECTICAL</u>						
022087 001					NDF	Jan 23, 2012
<u>CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE - PEPICID COMPLETE</u>						
020958 001	5075114	May 23, 2010	DP			
	5075114*PED	Nov 23, 2010				
	6814978	Aug 26, 2021	DP			
	6814978*PED	Feb 26, 2022				
<u>CALCIUM CARBONATE; RISEDRONATE SODIUM - ACTONEL WITH CALCIUM (COPACKAGED)</u>						
021823 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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
020838 001	>A> 5196444	Jun 04, 2012	DS DP U-660			
	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-660			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
020838 002	>A> 5196444	Jun 04, 2012	DS DP U-660			
	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-660			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
020838 003	>A> 5196444	Jun 04, 2012	DS DP U-660			
	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-660			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL - ATACAND</u>						
020838 004	>A> 5196444	Jun 04, 2012	DS DP U-660			
	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-660			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE - ATACAND HCT</u>						
021093 001	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-3			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE - ATACAND HCT</u>						
021093 002	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-3			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				
<u>CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE - ATACAND HCT</u>						
021093 003	>A> 5196444	Jun 04, 2012	DS DP U-3			
	>A> 5196444*PED	Dec 04, 2012				
	>A> 5534534	Jul 09, 2013	DP			
	>A> 5534534*PED	Jan 09, 2014				
	>A> 5705517	Apr 18, 2011	DS DP U-3			
	>A> 5705517*PED	Oct 18, 2011				
	>A> 7538133	Apr 18, 2011	DS			
	>A> 7538133*PED	Oct 18, 2011				
<u>CICLESONIDE - ALVESCO</u>						
021658 002					NDF NCE	Jan 10, 2011 Oct 20, 2011
<u>CICLESONIDE - ALVESCO</u>						
021658 003					NDF NCE	Jan 10, 2011 Oct 20, 2011
<u>CICLESONIDE - OMNARIS</u>						
022004 001	>A> 5482934	Oct 24, 2017	DS DP U-557			
<u>CIPROFLOXACIN HYDROCHLORIDE - CETRAXAL</u>						
021918 001					NDF	May 01, 2012
<u>CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE - CIPRO HC</u>						
020805 001	5843930	Jun 06, 2015	U-646			
<u>CLARITHROMYCIN - BIAXIN XL</u>						
050775 001	6551616	Jul 15, 2017	U-924			
<u>CLOBETASOL PROPIONATE - OLUX</u>						
021142 001	6126920	Mar 01, 2016	U-484			
<u>CLOPIDOGREL BISULFATE - PLAVIX</u>						
020839 002	4847265	Nov 17, 2011	DS DP			
	6429210	Jun 10, 2019	DS DP			
	6504030	Jun 10, 2019	DS			
<u>COLCHICINE - COLCRYS</u>						
022352 001					>A> I-603	Jul 30, 2012

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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>						
021141 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>						
021176 001	5607669	Jun 10, 2014	U-323		I-553	Jan 18, 2011
	5607669*PED	Dec 10, 2014			PED	Jul 18, 2011
	5679717	Apr 29, 2014	U-323			
	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				
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>						
021777 001	7544372	Nov 14, 2023	U-979			
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>						
021777 002	7544372	Nov 14, 2023	U-979			
<u>CYCLOSPORINE - SANDIMMUNE</u>						
050625 001	7511014	Feb 16, 2010	DP			
<u>CYCLOSPORINE - SANDIMMUNE</u>						
050625 002	7511014	Feb 16, 2010	DP			
<u>CYCLOSPORINE - SANDIMMUNE</u>						
050625 003	7511014	Feb 16, 2010	DP			
<u>DASATINIB - SPRYCEL</u>						
021986 001	7491725	Oct 13, 2025	DS DP		D-120	May 21, 2012

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DASATINIB - SPRYCEL</u>						
021986 002	7491725	Oct 13, 2025	DS DP		D-120	May 21, 2012
<u>DASATINIB - SPRYCEL</u>						
021986 003	7491725	Oct 13, 2025	DS DP		D-120	May 21, 2012
<u>DASATINIB - SPRYCEL</u>						
021986 004	6596746	Jun 28, 2020	DS DP U-780		D-120	May 21, 2012
	6596746	Jun 28, 2020	DS DP U-748			
	7125875	Apr 13, 2020	U-780			
	7125875	Apr 13, 2020	U-779			
	7153856	Apr 28, 2020	U-780			
	7491725	Oct 13, 2025	DS DP			
<u>DEGARELIX ACETATE - FIRMAGON</u>						
022201 001	5925730	Apr 11, 2017	DS DP U-943			
<u>DEGARELIX ACETATE - FIRMAGON</u>						
022201 002	5925730	Apr 11, 2017	DS DP U-943			
<u>DESONIDE - DESONATE</u>						
021844 001					NDF	Oct 20, 2009
<u>DEXAMETHASONE - OZURDEX</u>						
022315 001	6726918	Oct 20, 2020	DP U-985		NDF	Jun 17, 2012
	6899717	Nov 01, 2023	U-985			
	7033605	Oct 20, 2020	DP			
<u>DEXAMETHASONE; TOBRAMYCIN - TOBRADEX ST</u>						
050818 001	5149694	Sep 22, 2009	U-953			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEXLANSOPRAZOLE - KAPIDEX</u>						
022287 001	5045321	Sep 03, 2008	DP		NP	Jan 30, 2012
	5045321*PED	Mar 03, 2009			PED	Jul 30, 2012
	5093132	Sep 03, 2008	DP U-949			
	5093132	Sep 03, 2008	DP U-950			
	5093132	Sep 03, 2008	DP U-951			
	5093132*PED	Mar 03, 2009				
	5433959	Sep 03, 2008	DP U-949			
	5433959	Sep 03, 2008	DP U-950			
	5433959	Sep 03, 2008	DP U-951			
	5433959*PED	Mar 03, 2009				
	6462058	Jun 15, 2020	DS DP U-951			
	6462058	Jun 15, 2020	DS DP U-950			
	6462058	Jun 15, 2020	DS DP U-949			
	6462058*PED	Dec 15, 2020				
	6664276	Jun 15, 2020	DS DP U-949			
	6664276	Jun 15, 2020	DS DP U-950			
	6664276	Jun 15, 2020	DS DP U-951			
	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				
<u>DEXLANSOPRAZOLE - KAPIDEX</u>						
022287 002	5045321	Sep 03, 2008	DP		NP	Jan 30, 2012
	5045321*PED	Mar 03, 2009			PED	Jul 30, 2012
	5093132	Sep 03, 2008	DP U-949			
	5093132	Sep 03, 2008	DP U-950			
	5093132	Sep 03, 2008	DP U-951			
	5093132*PED	Mar 03, 2009				
	5433959	Sep 03, 2008	DP U-949			
	5433959	Sep 03, 2008	DP U-950			
	5433959	Sep 03, 2008	DP U-951			
	5433959*PED	Mar 03, 2009				
	6462058	Jun 15, 2020	DS DP U-951			
	6462058	Jun 15, 2020	DS DP U-950			
	6462058	Jun 15, 2020	DS DP U-949			
	6462058*PED	Dec 15, 2020				
	6664276	Jun 15, 2020	DS DP U-949			
	6664276	Jun 15, 2020	DS DP U-950			
	6664276	Jun 15, 2020	DS DP U-951			
	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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DICLOFENAC POTASSIUM - CAMBIA</u>						
022165 001	>A> 6974595	May 15, 2017	U-436			
<u>DICLOFENAC POTASSIUM - ZIPSOR</u>						
022202 001	6365180	Jul 16, 2019	DP U-980		NDF	Jun 16, 2012
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM CD</u>						
020062 005	5286497	May 20, 2011	DP			
	5439689	Aug 08, 2012	DP U-107			
	5470584	May 20, 2011	DP			
<u>DINOPROSTONE - CERVIDIL</u>						
020411 001	5269321	Jul 14, 2012	DP U-110			
<u>DIVALPROEX SODIUM - DIVALPROEX SODIUM</u>						
077567 002					PC	Aug 01, 2009
<u>DOXERCALCIFEROL - HECTOROL</u>						
020862 001	>A> 5602116	Feb 11, 2014	U-987			
	>A> 5602116	Feb 11, 2014	U-278			
	>A> 6903083	Jul 18, 2021	DS DP	Y		
<u>DOXERCALCIFEROL - HECTOROL</u>						
020862 002	>A> 5602116	Feb 11, 2014	U-987			
	>A> 5602116	Feb 11, 2014	U-278			
	>A> 6903083	Jul 18, 2021	DS DP	Y		
<u>DOXERCALCIFEROL - HECTOROL</u>						
020862 003	>A> 5602116	Feb 11, 2014	U-987			
<u>DOXERCALCIFEROL - HECTOROL</u>						
021027 001	5707980	Aug 17, 2010	U-321	Y		
	>A> 6903083	Jul 18, 2021	DS DP	Y		
<u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u>						
022425 001	>A> 5223510	Jul 26, 2011	DS DP U-992		>A> NCE	Jul 01, 2014
	>A> 7323493	Jun 19, 2018	DP			
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
022291 001	7473686	Jul 24, 2021	DS DP U-930			
	7547719	Mar 04, 2024	DS DP U-930			
<u>ELTROMBOPAG OLAMINE - PROMACTA</u>						
022291 002	7473686	Jul 24, 2021	DS DP U-930			
	7547719	Mar 04, 2024	DS DP U-930			
<u>EPINEPHRINE - EPIPEN</u>						
019430 001	7449012	Sep 11, 2025	DP			
<u>EPINEPHRINE - EPIPEN JR.</u>						
019430 002	7449012	Sep 11, 2025	DP			
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
021323 001					NPP	Mar 19, 2012
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
021323 002					NPP	Mar 19, 2012
<u>ESCITALOPRAM OXALATE - LEXAPRO</u>						
021323 003					NPP	Mar 19, 2012

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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>						
021365	001				NPP	Mar 19, 2012
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
021153	001				NPP PED	Apr 28, 2009 Oct 28, 2009
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
021153	002				NPP PED	Apr 28, 2009 Oct 28, 2009
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
021957	001				M-86 NPP PED PED	Jun 18, 2012 Apr 28, 2009 Dec 18, 2012 Oct 28, 2009
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
021957	002				M-86 NPP PED PED	Jun 18, 2012 Apr 28, 2009 Dec 18, 2012 Oct 28, 2009
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>						
022101	001				NPP PED	Feb 27, 2011 Aug 27, 2011
<u>ESTRADIOL - ELESTRIN</u>						
021813	001	7470433	Aug 03, 2021	DP		
<u>ESTRADIOL - EVAMIST</u>						
022014	001	6818226	Feb 19, 2017	DP U-889		
		6818226	Feb 19, 2017	DP U-888		
		6923983	Feb 19, 2017	DP U-889		
		6923983	Feb 19, 2017	DP U-888		
		6978945	Nov 30, 2021	DP		
<u>ETHINYL ESTRADIOL; NORGESTIMATE - TRI LO SPRINT</u>						
076784	001				>A> PC	Dec 29, 2009
<u>EVEROLIMUS - AFINITOR</u>						
022334	001	5665772	Sep 09, 2014	DS DP	NCE	Mar 30, 2014
		6004973	Jul 12, 2016	DP		
		7297703	Dec 06, 2019	DP		
<u>EVEROLIMUS - AFINITOR</u>						
022334	002	5665772	Sep 09, 2014	DS DP	NCE	Mar 30, 2014
		6004973	Jul 12, 2016	DP		
		7297703	Dec 06, 2019	DP		
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
021773	001	7521423	Oct 15, 2017	DP		
<u>EXENATIDE SYNTHETIC - BYETTA</u>						
021773	002	7521423	Oct 15, 2017	DP		
<u>FEBUXOSTAT - ULORIC</u>						
021856	001	5614520	Mar 25, 2014	DS DP U-954	NCE	Feb 13, 2014
		6225474	Jun 18, 2019	DS		
		7361676	Mar 08, 2024	DP		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FEBUXOSTAT - ULORIC</u>						
021856 002	5614520	Mar 25, 2014	DS DP U-954		NCE	Feb 13, 2014
	6225474	Jun 18, 2019	DS			
	7361676	Mar 08, 2024	DP			
<u>FENTANYL CITRATE - ONSOLIS</u>						
022266 001					>A> NP	Jul 16, 2012
<u>FENTANYL CITRATE - ONSOLIS</u>						
022266 002					>A> NP	Jul 16, 2012
<u>FENTANYL CITRATE - ONSOLIS</u>						
022266 003					>A> NP	Jul 16, 2012
<u>FENTANYL CITRATE - ONSOLIS</u>						
022266 004					>A> NP	Jul 16, 2012
<u>FENTANYL CITRATE - ONSOLIS</u>						
022266 005					>A> NP	Jul 16, 2012
<u>FERUMOXYTOL - FERAHEME</u>						
022180 001	>A> 6599498	Mar 08, 2020	DS DP		NP	Jun 30, 2012
	>A> 7553479	Mar 08, 2020	DS DP			
<u>FLUDARABINE PHOSPHATE - OFORTA</u>						
022273 001	7148207	Dec 20, 2022	DP U-944		NDF	Dec 18, 2011
<u>FLUOXETINE HYDROCHLORIDE - PROZAC</u>						
018936 001	6960577	Nov 01, 2017	U-963		I-589	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE - PROZAC</u>						
018936 003	6960577	Nov 01, 2017	U-963		I-589	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE - PROZAC</u>						
018936 006	6960577	Nov 01, 2017	U-963		I-589	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>						
021520 001	6960577	Nov 01, 2017	U-962		I-593	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>						
021520 002	6960577	Nov 01, 2017	U-962		I-593	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>						
021520 003	6960577	Nov 01, 2017	U-962		I-593	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>						
021520 004	6960577	Nov 01, 2017	U-962		I-593	Mar 19, 2012
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>						
021520 005	6960577	Nov 01, 2017	U-962		I-593	Mar 19, 2012
<u>FLUTICASONE FUROATE - VERAMYST</u>						
022051 001	>A> 7541350	Aug 03, 2021	DP U-988			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - FLOVENT DISKUS 100</u>						
020833 002	5590645	Mar 01, 2011	DP			
	5590645*PED	Sep 01, 2011				
	5860419	Mar 01, 2011	DP			
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666	Mar 01, 2011	DP			
	6032666*PED	Sep 01, 2011				
	6378519	Mar 01, 2011	DP			
	6378519*PED	Sep 01, 2011				
	6536427	Mar 01, 2011	DP			
	6536427*PED	Sep 01, 2011				
	6792945	Mar 01, 2011	DP			
	6792945*PED	Sep 01, 2011				
	7225808	Mar 01, 2011	DP			
	7225808*PED	Sep 01, 2011				
	7389775	Mar 01, 2011	DP			
	7389775*PED	Sep 01, 2011				
<u>FLUTICASONE PROPIONATE - FLOVENT DISKUS 250</u>						
020833 003	5590645	Mar 01, 2011	DP			
	5590645*PED	Sep 01, 2011				
	5860419	Mar 01, 2011	DP			
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666	Mar 01, 2011	DP			
	6032666*PED	Sep 01, 2011				
	6378519	Mar 01, 2011	DP			
	6378519*PED	Sep 01, 2011				
	6536427	Mar 01, 2011	DP			
	6536427*PED	Sep 01, 2011				
	6792945	Mar 01, 2011	DP			
	6792945*PED	Sep 01, 2011				
	7225808	Mar 01, 2011	DP			
	7225808*PED	Sep 01, 2011				
	7389775	Mar 01, 2011	DP			
	7389775*PED	Sep 01, 2011				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE - FLOVENT DISKUS 50						
020833 001	5590645	Mar 01, 2011	DP			
	5590645*PED	Sep 01, 2011				
	5860419	Mar 01, 2011	DP			
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666	Mar 01, 2011	DP			
	6032666*PED	Sep 01, 2011				
	6378519	Mar 01, 2011	DP			
	6378519*PED	Sep 01, 2011				
	6536427*PED	Sep 01, 2011				
	6792945	Mar 01, 2011	DP			
	6792945*PED	Sep 01, 2011				
	7225808	Mar 01, 2011	DP			
	7225808*PED	Sep 01, 2011				
	7389775	Mar 01, 2011	DP			
	7389775*PED	Sep 01, 2011				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>						
021433 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				
	6253762	Apr 14, 2015	DP U-582			
	6253762*PED	Oct 14, 2015				
	6315173	Jun 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				
	6546928	Apr 14, 2015	DP U-583			
	6546928*PED	Oct 14, 2015				
	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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE - FLOVENT HFA						
021433 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				
	6253762	Apr 14, 2015	DP U-582			
	6253762*PED	Oct 14, 2015				
	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				
	6546928	Apr 14, 2015	DP U-583			
	6546928*PED	Oct 14, 2015				
	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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE - FLOVENT HFA						
021433 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				
	6253762	Apr 14, 2015	DP U-582			
	6253762*PED	Oct 14, 2015	U-582			
	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			
	6546928	Apr 14, 2015	DP U-583			
	6546928*PED	Oct 14, 2015	U-583			
	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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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 100/50								
021077 001	5590645	Mar	01, 2011	DP		M-84	Mar	31, 2012
	5590645*PED	Sep	01, 2011					
	5860419	Mar	01, 2011	DP				
	5860419*PED	Sep	01, 2011					
	5873360	Feb	23, 2016	DP				
	5873360*PED	Aug	23, 2016					
	6032666	Mar	01, 2011	DP				
	6032666*PED	Sep	01, 2011					
	6378519	Mar	01, 2011	DP				
	6378519*PED	Sep	01, 2011					
	6792945	Mar	01, 2011	DP				
	6792945*PED	Sep	01, 2011					
	7225808	Mar	01, 2011	DP				
	7225808*PED	Sep	01, 2011					
	7389775	Mar	01, 2011	DP				
	7389775*PED	Sep	01, 2011					
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50								
021077 002	5590645	Mar	01, 2011	DP		M-84	Mar	31, 2012
	5590645*PED	Sep	01, 2011					
	5860419	Mar	01, 2011	DP				
	5860419*PED	Sep	01, 2011					
	5873360	Feb	23, 2016	DP				
	5873360*PED	Aug	23, 2016					
	6032666	Mar	01, 2011	DP				
	6032666*PED	Sep	01, 2011					
	6378519	Mar	01, 2011	DP				
	6378519*PED	Sep	01, 2011					
	6792945	Mar	01, 2011	DP				
	6792945*PED	Sep	01, 2011					
	7225808	Mar	01, 2011	DP				
	7225808*PED	Sep	01, 2011					
	7389775	Mar	01, 2011	DP				
	7389775*PED	Sep	01, 2011					

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50</u>						
021077 003	5590645	Mar 01, 2011	DP		M-84	Mar 31, 2012
	5590645*PED	Sep 01, 2011				
	5860419	Mar 01, 2011	DP			
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666	Mar 01, 2011	DP			
	6032666*PED	Sep 01, 2011				
	6378519	Mar 01, 2011	DP			
	6378519*PED	Sep 01, 2011				
	6792945	Mar 01, 2011	DP			
	6792945*PED	Sep 01, 2011				
	7225808	Mar 01, 2011	DP			
	7225808*PED	Sep 01, 2011				
	7389775	Mar 01, 2011	DP			
	7389775*PED	Sep 01, 2011				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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						
021254 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				
	6253762	Apr 14, 2015	DP		U-738	
	6253762*PED	Oct 14, 2015			U-738	
	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				
	6546928	Apr 14, 2015	DP			
	6546928*PED	Oct 14, 2015				
	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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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						
021254 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			
	6253762	Apr 14, 2015	DP U-738			
	6253762*PED	Oct 14, 2015	DP U-738			
	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			
	6546928	Apr 14, 2015	DP			
	6546928*PED	Oct 14, 2015	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				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List 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>						
021254 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				
	6170717	Dec 23, 2017	DP			
	6170717*PED	Jun 23, 2018				
	6251368	Dec 04, 2012	DP			
	6251368*PED	Jun 04, 2013	DP			
	6253762	Apr 14, 2015	DP U-738			
	6253762*PED	Oct 14, 2015	DP U-738			
	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			
	6546928	Apr 14, 2015	DP			
	6546928*PED	Oct 14, 2015	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				
<u>FLUVOXAMINE MALEATE - LUVOX</u>						
021519 001					M-83	Apr 14, 2011
<u>FLUVOXAMINE MALEATE - LUVOX</u>						
021519 002					M-83	Apr 14, 2011
<u>FLUVOXAMINE MALEATE - LUVOX</u>						
021519 003					M-83	Apr 14, 2011
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
020378 004	>A> 7563763	Aug 23, 2019	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FOLLITROPIN ALFA/BETA - GONAL-F</u>						
020378 005	>A> 7563763	Aug 23, 2019	DP			
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF PEN</u>						
021684 001	>A> 7446090	Aug 23, 2019	DP			
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF PEN</u>						
021684 002	>A> 7446090	Aug 23, 2019	DP			
<u>FOLLITROPIN ALFA/BETA - GONAL-F RFF PEN</u>						
021684 003	>A> 7446090	Aug 23, 2019	DP			
<u>FOSPROPOFOL DISODIUM - LUSEDRA</u>						
022244 001	6204257	Jun 07, 2018	DS DP U-945		NCE	Dec 12, 2013
<u>GADODIAMIDE - OMNISCAN</u>						
022066 002	5362475	Nov 08, 2011	DS			
<u>GATIFLOXACIN - ZYMAR</u>						
021493 001	4980470	Dec 15, 2009	DS DP			
	4980470*PED	Jun 15, 2010				
	5880283	Dec 05, 2015				
	5880283*PED	Jun 05, 2016				
	6333045	Aug 20, 2019	DP			
	6333045*PED	Feb 20, 2020				
<u>GLATIRAMER ACETATE - COPAXONE</u>						
020622 001					I-594	Feb 27, 2012
<u>GLATIRAMER ACETATE - COPAXONE</u>						
020622 002					I-594	Feb 27, 2012
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>						
021925 001	7538125	Jun 19, 2016	DP			
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>						
021925 002	7538125	Jun 19, 2016	DP			
<u>GOSERELIN ACETATE - ZOLADEX</u>						
019726 001	7500964	Feb 26, 2021	DP			
<u>GOSERELIN ACETATE - ZOLADEX</u>						
020578 001	7500964	Feb 26, 2021	DP			
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u>						
021162 003	5591762	Jan 07, 2014	DS DP U-3			
<u>IBUPROFEN - CALDOLOR</u>						
022348 001	6727286	Nov 27, 2021	DP U-981		NP	Jun 11, 2012
<u>IBUPROFEN - CALDOLOR</u>						
022348 002	6727286	Nov 27, 2021	DP U-981		NP	Jun 11, 2012
<u>ILOPERIDONE - FANAPT</u>						
022192 001	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>						
022192 002	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ILOPERIDONE - FANAPT</u>						
022192 003	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>						
022192 004	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>						
022192 005	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>						
022192 006	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014
<u>ILOPERIDONE - FANAPT</u>						
022192 007	RE39198	Nov 15, 2011	DS DP U-971		NCE	May 06, 2014
<u>IMATINIB MESYLATE - GLEEVEC</u>						
021588 001					I-583	Dec 19, 2011
<u>IMATINIB MESYLATE - GLEEVEC</u>						
021588 002					I-583	Dec 19, 2011
<u>INSULIN DETEMIR RECOMBINANT - LEVEMIR</u>						
021536 001	5750497	May 16, 2019	DS DP U-668			
<u>INSULIN GLARGINE RECOMBINANT - LANTUS</u>						
021081 001	5656722	Aug 12, 2014	DS DP U-948			
	5656722*PED	Feb 12, 2015				
	7476652	Jul 23, 2023	DP			
	7476652*PED	Jan 23, 2024				
<u>INSULIN GLULISINE RECOMBINANT - APIDRA</u>						
021629 002					NCE	Apr 16, 2009
<u>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u>						
021629 003					NPP NCE	Oct 24, 2011 Apr 16, 2009
<u>IODIXANOL - VISIPAQUE 270</u>						
020351 001	5366722	Nov 22, 2011	DP			
	RE36418	Jul 12, 2011	DP			
<u>IODIXANOL - VISIPAQUE 270</u>						
020808 001	5366722	Nov 22, 2011	DP			
	RE36418	Jul 12, 2011	DP			
<u>IODIXANOL - VISIPAQUE 320</u>						
020351 002	RE36418	Jul 12, 2011	DP			
<u>IODIXANOL - VISIPAQUE 320</u>						
020808 002	RE36418	Jul 12, 2011	DP			
<u>IXABEPILONE - IXEMPRA KIT</u>						
022065 001	6605599	May 26, 2018	DS DP U-961			
	6670384	Jan 23, 2022	DP U-960			
	6670384	Jan 23, 2022	DP U-959			
	7022330	Jan 23, 2022	DP U-958			
	7125899	May 26, 2018	DS DP U-957			
	7312237	Aug 21, 2024	U-965			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IXABEPILONE - IXEMPRA KIT</u>						
022065 002	6605599	May 26, 2018	DS DP U-961			
	6670384	Jan 23, 2022	DP U-960			
	6670384	Jan 23, 2022	DP U-959			
	7022330	Jan 23, 2022	DP U-958			
	7125899	May 26, 2018	DS DP U-957			
	7312237	Aug 21, 2024	U-965			
<u>KETOCONAZOLE - EXTINA</u>						
021738 001	>A> 7553835	Oct 19, 2018	DP U-245			
<u>KETOROLAC TROMETHAMINE - ACUVAIL</u>						
022427 001					>A> NP	Jul 22, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
020406 001					M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
020406 002					M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
021281 001					M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
021281 002					M-85 PED	Oct 28, 2011 Apr 28, 2012
<u>LANSOPRAZOLE - PREVACID</u>						
021428 001	7431942	May 17, 2019	DP		M-85 PED	Oct 28, 2011 Apr 28, 2012
	7431942*PED	Nov 17, 2019				
<u>LANSOPRAZOLE - PREVACID</u>						
021428 002	7431942	May 17, 2019	DP		M-85 PED	Oct 28, 2011 Apr 28, 2012
	7431942*PED	Nov 17, 2019				
<u>LANSOPRAZOLE - PREVACID 24 HR</u>						
022327 001					NP	May 18, 2012
<u>LANSOPRAZOLE - PREVACID IV</u>						
021566 001	4628098	May 10, 2009	DS			
	4628098*PED	Nov 10, 2009				
	7396841	Aug 17, 2021	DP U-947			
	7396841*PED	Feb 17, 2022				
<u>LANSOPRAZOLE; NAPROXEN - PREVACID NAPRAPAC 500 (COPACKAGED)</u>						
021507 004	4628098	May 10, 2009	DS			
	4628098*PED	Nov 10, 2009				
	5045321	Sep 03, 2008	DP			
	5045321*PED	Mar 03, 2009				
	5093132	Sep 03, 2008	DP			
	5093132*PED	Mar 03, 2009				
	5433959	Sep 03, 2008	DP			
	5433959*PED	Mar 03, 2009				
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
021468 001	5968976	Oct 26, 2018	DP U-613			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
021468 002	5968976	Oct 26, 2018	DP U-613			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
021468 003	5968976	Oct 26, 2018	DP U-613			
<u>LANTHANUM CARBONATE - FOSRENOL</u>						
021468 004	5968976	Oct 26, 2018	DP U-613			
<u>LEVETIRACETAM - KEPRA XR</u>						
022285 002					NDF	Sep 12, 2011
<u>LEVONORGESTREL - PLAN B ONE-STEP</u>						
021998 001					>A> NP	Jul 10, 2012
<u>LIDOCAINE HYDROCHLORIDE - ZINGO</u>						
022114 001					NPP	Jan 08, 2012
<u>MALATHION - OVIDE</u>						
018613 001	7560445	Feb 01, 2027	DS DP U-986			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>						
021487 001	5061703	Apr 11, 2015	U-539			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>						
021487 002	5061703	Apr 11, 2015	U-539			
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>						
021627 001	5061703	Apr 11, 2015	U-539			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
022024 001	4687777	Jan 17, 2011	DS			
	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			
<u>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</u>						
022024 002	4687777	Jan 17, 2011	DS			
	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			
<u>MICONAZOLE NITRATE - MONISTAT 1 COMBINATION PACK</u>						
021308 001	5514698	Mar 21, 2014		Y		
	6153635	Nov 28, 2020		Y		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
022256 001	6602911	Nov 05, 2021	U-882		NCE	Jan 14, 2014
	6992110	Nov 05, 2021	U-882			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
022256 002	6602911	Nov 05, 2021	U-882		NCE	Jan 14, 2014
	6992110	Nov 05, 2021	U-882			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
022256 003	6602911	Nov 05, 2021	U-882		NCE	Jan 14, 2014
	6992110	Nov 05, 2021	U-882			
<u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u>						
022256 004	6602911	Nov 05, 2021	U-882		NCE	Jan 14, 2014
	6992110	Nov 05, 2021	U-882			
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
050808 002	7541347	Apr 02, 2027	U-917			
	7544373	Apr 02, 2027	DP			
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
050808 004	>A> 5908838	Feb 19, 2018	U-917			
<u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u>						
050808 005	>A> 5908838	Feb 19, 2018	U-917			
<u>MOMETASONE FUROATE - ASMANEX TWISTHALER</u>						
021067 002	5394868	Jun 25, 2012	DP		NPP	Feb 01, 2011
	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				
<u>MORPHINE SULFATE - AVINZA</u>						
021260 005	6066339	Nov 25, 2017	DP			
<u>MORPHINE SULFATE - AVINZA</u>						
021260 006	6066339	Nov 25, 2017	DP			
<u>NICOTINE POLACRILEX - NICORETTE</u>						
022360 001	5110605	Aug 21, 2010	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NICOTINE POLACRILEX - NICORETTE</u>						
022360 002	5110605	Aug 21, 2010	DP			
<u>NITROGLYCERIN - NITROMIST</u>						
021780 001	5869082	Apr 16, 2016	DP			
<u>OLANZAPINE - ZYPREXA</u>						
020592 001	6960577	Nov 01, 2017	U-963		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA</u>						
020592 002	6960577	Nov 01, 2017	U-963		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA</u>						
020592 003	6960577	Nov 01, 2017	U-963		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA</u>						
020592 004	6960577	Nov 01, 2017	U-963		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA</u>						
020592 005	6960577	Nov 01, 2017	U-963		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA</u>						
020592 006	6960577	Nov 01, 2017	U-963		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
021086 001	6960577	Nov 01, 2017	U-964		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
021086 002	6960577	Nov 01, 2017	U-964		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
021086 003	6960577	Nov 01, 2017	U-964		I-591	Mar 19, 2012
<u>OLANZAPINE - ZYPREXA ZYDIS</u>						
021086 004	6960577	Nov 01, 2017	U-964		I-591	Mar 19, 2012
<u>OLOPATADINE HYDROCHLORIDE - PATADAY</u>						
021545 001	>A> 5116863	Dec 18, 2010	DS DP			
	>A> 5116863*PED	Jun 18, 2011				
	>A> 5641805	Jun 06, 2015			U-765	
	>A> 5641805*PED	Dec 06, 2015				
	>A> 6995186	Nov 12, 2023	DP	U-765		
	>A> 6995186*PED	May 12, 2024				
	>A> 7402609	Jun 19, 2022	DP			
	>A> 7402609*PED	Dec 19, 2022				
<u>OLOPATADINE HYDROCHLORIDE - PATANASE</u>						
021861 001	>A> 5116863	Dec 18, 2010	DS DP		>A> NDF	Apr 15, 2011
	>A> 5116863*PED	Jun 18, 2011			>A> PED	Oct 15, 2011
<u>OLOPATADINE HYDROCHLORIDE - PATANOL</u>						
020688 001	>A> 5116863	Dec 18, 2010				
	>A> 5116863*PED	Jun 18, 2011				
	>A> 5641805	Jun 06, 2015		U-184		
	>A> 5641805*PED	Dec 06, 2015				
<u>OMEGA-3-ACID ETHYL ESTERS - LOVAZA</u>						
021654 001	5656667	Apr 10, 2017	DS DP	U-822		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYBUTYNIN CHLORIDE - GELNIQUE</u>						
022204 001	7029694	Apr 26, 2020	DP U-318		NDF	Jan 27, 2012
	7179483	Apr 26, 2020	U-318			
<u>PALIPERIDONE - INVEGA</u>						
021999 006	5158952	Oct 27, 2009	DP U-90			
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
020987 001	4758579	Jul 19, 2010				
	4758579*PED	Jan 19, 2011				
	5997903	Dec 07, 2016				
	5997903*PED	Jun 07, 2017				
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
020987 002	4758579	Jul 19, 2010				
	4758579*PED	Jan 19, 2011				
	5997903	Dec 07, 2016				
	5997903*PED	Jun 07, 2017				
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
022020 001	4758579	Jul 19, 2010	DS DP U-859			
	4758579*PED	Jan 19, 2011				
	7544370	Feb 07, 2026	DP			
	7544370*PED	Aug 07, 2026				
	>A> 7550153	Sep 30, 2024			U-859	
	>A> 7550153*PED	Mar 30, 2025				
	>A> 7553498	Sep 30, 2024			U-859	
	>A> 7553498*PED	Mar 30, 2025				
<u>PANTOPRAZOLE SODIUM - PROTONIX IV</u>						
020988 001	4758579	Jul 19, 2010				
	4758579*PED	Jan 19, 2011				
	6780881	Nov 17, 2021	DP			
	6780881*PED	May 17, 2022				
	7351723	Nov 17, 2021	DP			
	7351723*PED	May 17, 2022				
<u>PARICALCITOL - ZEMPLAR</u>						
021606 001					I-599	Jun 29, 2012
<u>PARICALCITOL - ZEMPLAR</u>						
021606 002					I-599	Jun 29, 2012
<u>PARICALCITOL - ZEMPLAR</u>						
021606 003					I-599	Jun 29, 2012
<u>PEMETREXED DISODIUM - ALIMTA</u>						
021462 001					>A> I-601	Jul 02, 2012
<u>PEMETREXED DISODIUM - ALIMTA</u>						
021462 002					>A> I-601	Jul 02, 2012
<u>PHENTOLAMINE MESYLATE - ORAVERSE</u>						
022159 001	6764678	May 11, 2021	U-967			
	6872390	May 11, 2021	DP			
	7229630	Jun 20, 2023	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>PRAMLINTIDE ACETATE - SYMLIN</u>								
021332 002	5814600	Sep	29, 2015	U-639				
	5814600	Sep	29, 2015	U-638				
	5814600	Sep	29, 2015	U-637				
	5998367	Mar	08, 2011	DS DP				
	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				
	6610824	Mar	03, 2011	DS				
	7407934	Mar	08, 2011	U-640				
	7407934	Mar	08, 2011	U-637				
<u>PRAMLINTIDE ACETATE - SYMLIN</u>								
021332 003	5814600	Sep	29, 2015	U-639				
	5814600	Sep	29, 2015	U-638				
	5814600	Sep	29, 2015	U-637				
	5998367	Mar	08, 2011	DS DP				
	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				
	6610824	Mar	03, 2011	DS				
	7407934	Mar	08, 2011	U-640				
	7407934	Mar	08, 2011	U-637				
<u>PRASUGREL HYDROCHLORIDE - EFFIENT</u>								
022307 001	>A> 5288726	Sep	08, 2012	DS DP U-991		>A> NCE	Jul	10, 2014
	>A> 6693115	Jul	03, 2021	DS DP U-991				
<u>PRASUGREL HYDROCHLORIDE - EFFIENT</u>								
022307 002	>A> 5288726	Sep	08, 2012	DS DP U-991		>A> NCE	Jul	10, 2014
	>A> 6693115	Jul	03, 2021	DS DP U-991				
<u>QUETIAPINE FUMARATE - SEROQUEL</u>								
020639 001	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010
<u>QUETIAPINE FUMARATE - SEROQUEL</u>								
020639 002	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010
<u>QUETIAPINE FUMARATE - SEROQUEL</u>								
020639 003	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
QUETIAPINE FUMARATE - SEROQUEL								
020639 004	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010
QUETIAPINE FUMARATE - SEROQUEL								
020639 005	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010
QUETIAPINE FUMARATE - SEROQUEL								
020639 006	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010
QUETIAPINE FUMARATE - SEROQUEL								
020639 007	4879288	Sep	26, 2011	DS DP U-550		I-560	May	13, 2011
	4879288*PED	Mar	26, 2012			I-503 PED PED	Oct Nov Apr	20, 2009 13, 2011 20, 2010
QUETIAPINE FUMARATE - SEROQUEL XR								
022047 001	4879288	Sep	26, 2011	DS DP U-814		D-117	Oct	08, 2011
	4879288	Sep	26, 2011	DS DP U-601		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		NDF	May	17, 2010
	5948437*PED	Nov	28, 2017			PED	Apr	08, 2012
	5948437*PED	Nov	28, 2017			PED	Nov	17, 2010
QUETIAPINE FUMARATE - SEROQUEL XR								
022047 002	4879288	Sep	26, 2011	DS DP U-814		D-117	Oct	08, 2011
	4879288	Sep	26, 2011	DS DP U-601		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		NDF	May	17, 2010
	5948437*PED	Nov	28, 2017			PED	Apr	08, 2012
	5948437*PED	Nov	28, 2017			PED	Nov	17, 2010
QUETIAPINE FUMARATE - SEROQUEL XR								
022047 003	4879288	Sep	26, 2011	DS DP U-814		D-117	Oct	08, 2011
	4879288	Sep	26, 2011	DS DP U-601		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		NDF	May	17, 2010
	5948437*PED	Nov	28, 2017			PED	Apr	08, 2012
	5948437*PED	Nov	28, 2017			PED	Nov	17, 2010
QUETIAPINE FUMARATE - SEROQUEL XR								
022047 004	4879288	Sep	26, 2011	DS DP U-814		D-117	Oct	08, 2011
	4879288	Sep	26, 2011	DS DP U-601		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		NDF	May	17, 2010
	5948437*PED	Nov	28, 2017			PED	Apr	08, 2012
	5948437*PED	Nov	28, 2017			PED	Nov	17, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>						
022047 005	4879288	Sep 26, 2011	DS DP U-814		D-117	Oct 08, 2011
	4879288	Sep 26, 2011	DS DP U-601		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		NDF	May 17, 2010
	5948437*PED	Nov 28, 2017			PED	Apr 08, 2012
					PED	Nov 17, 2010
<u>RASAGILINE MESYLATE - AZILECT</u>						
021641 001	5453446	Feb 07, 2017	U-219			
<u>RASAGILINE MESYLATE - AZILECT</u>						
021641 002	5453446	Feb 07, 2017	U-219			
<u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u>						
020630 001	5019583	Jul 12, 2010	DS DP U-952			
	5019583*PED	Jan 12, 2011				
<u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u>						
020630 002	5019583	Jul 12, 2010	DS DP U-952			
	5019583*PED	Jan 12, 2011				
<u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u>						
020630 003	5019583	Jul 12, 2010	DS DP U-952			
	5019583*PED	Jan 12, 2011				
<u>REPAGLINIDE - PRANDIN</u>						
020741 001	6677358	Jun 12, 2018	DS DP U-968			
<u>REPAGLINIDE - PRANDIN</u>						
020741 002	6677358	Jun 12, 2018	DS DP U-968			
<u>REPAGLINIDE - PRANDIN</u>						
020741 003	6677358	Jun 12, 2018	DS DP U-968			
<u>RISEDRONATE SODIUM - ACTONEL</u>						
020835 001	5583122	Dec 10, 2013	U-222		>A> M-61	Jul 23, 2012
	5583122*PED	Jun 10, 2014			>A> 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>						
020835 002	5583122	Dec 10, 2013	U-222		>A> M-61	Jul 23, 2012
	5583122*PED	Jun 10, 2014			>A> PED	Jan 23, 2013
	6096342	Nov 22, 2011				
	6096342*PED	May 22, 2012				
	6165513	Jun 10, 2018				
	6165513*PED	Dec 10, 2018				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RISEDRONATE SODIUM - ACTONEL</u>						
020835 003	5583122	Dec 10, 2013	DS DP U-756		I-309	Aug 11, 2009
	5583122	Dec 10, 2013	DS DP U-222		>A> M-61	Jul 23, 2012
	5583122*PED	Jun 10, 2014			>A> PED	Jan 23, 2013
	5994329	Jul 17, 2018	U-353		PED	Feb 11, 2010
	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>						
020835 004	5583122	Dec 10, 2013	DS DP U-353		D-105	Apr 16, 2010
	5583122*PED	Jun 10, 2014			>A> M-61	Jul 23, 2012
	6096342	Nov 22, 2011	DP U-353		>A> PED	Jan 23, 2013
	6096342*PED	May 22, 2012			PED	Oct 16, 2010
	6165513	Jun 10, 2018	DP			
	6165513*PED	Dec 10, 2018				
<u>RISEDRONATE SODIUM - ACTONEL</u>						
020835 005	5583122	Dec 10, 2013	DS DP U-353		>A> M-61	Jul 23, 2012
	5583122*PED	Jun 10, 2014			NS	Apr 22, 2011
	6165513	Jun 10, 2018	DP		>A> PED	Jan 23, 2013
	6165513*PED	Dec 10, 2018			PED	Oct 22, 2011
	7192938	May 06, 2023	U-353			
	7192938*PED	Nov 06, 2023				
<u>RISPERIDONE - RISPERDAL CONSTA</u>						
021346 001	5688801	Nov 18, 2014	U-972		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-543		I-596	May 15, 2012
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				
<u>RISPERIDONE - RISPERDAL CONSTA</u>						
021346 002	5688801	Nov 18, 2014	U-972		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-543		I-596	May 15, 2012
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				
<u>RISPERIDONE - RISPERDAL CONSTA</u>						
021346 003	5688801	Nov 18, 2014	U-972		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-543		I-596	May 15, 2012
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - RISPERDAL CONSTA</u>						
021346 004	5688801	Nov 18, 2014	U-972		I-597	May 15, 2012
	5688801	Nov 18, 2014	U-543		I-596	May 15, 2012
	7547452	Nov 19, 2013	DP			
	7547452*PED	May 19, 2014				
<u>RISPERIDONE - RISPERIDONE</u>						
076440 001					PC	Jul 29, 2009
<u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u>						
022008 006	5422123	Jun 06, 2012	DP		NDF	Jun 13, 2011
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
021366 002	6316460	Aug 04, 2020	DP		I-573	Nov 06, 2011
	6316460*PED	Feb 04, 2021			I-547	Nov 08, 2010
	6858618	Dec 17, 2021	U-618		PED	May 06, 2012
	6858618*PED	Jun 17, 2022			PED	May 08, 2011
	RE37314	Jan 08, 2016	DS			
	RE37314*PED	Jul 08, 2016				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
021366 003	6316460	Aug 04, 2020	DP		I-573	Nov 06, 2011
	6316460*PED	Feb 04, 2021			I-547	Nov 08, 2010
	6858618	Dec 17, 2021	U-618		PED	May 06, 2012
	6858618*PED	Jun 17, 2022			PED	May 08, 2011
	RE37314	Jan 08, 2016	DS			
	RE37314*PED	Jul 08, 2016				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
021366 004	6316460	Aug 04, 2020	DP		I-573	Nov 06, 2011
	6316460*PED	Feb 04, 2021			I-547	Nov 08, 2010
	6858618	Dec 17, 2021	U-618		PED	May 06, 2012
	6858618*PED	Jun 17, 2022			PED	May 08, 2011
	RE37314	Jan 08, 2016	DS			
	RE37314*PED	Jul 08, 2016				
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
021366 005	6316460	Aug 04, 2020	DP		I-573	Nov 06, 2011
	6316460*PED	Feb 04, 2021			I-547	Nov 08, 2010
	6858618	Dec 17, 2021	U-618		PED	May 06, 2012
	6858618*PED	Jun 17, 2022			PED	May 08, 2011
	RE37314	Jan 08, 2016	DS			
	RE37314*PED	Jul 08, 2016				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SALMETEROL XINAFOATE - SEREVENT</u>						
020692 001	5590645	Mar 01, 2011	DP			
	5590645*PED	Sep 01, 2011				
	5860419	Mar 01, 2011	DP			
	5860419*PED	Sep 01, 2011				
	5873360	Feb 23, 2016	DP			
	5873360*PED	Aug 23, 2016				
	6032666	Mar 01, 2011	DP			
	6032666*PED	Sep 01, 2011				
	6378519	Mar 01, 2011	DP			
	6378519*PED	Sep 01, 2011				
	6536427	Mar 01, 2011	DP			
	6536427*PED	Sep 01, 2011				
	6792945	Mar 01, 2011	DP			
	6792945*PED	Sep 01, 2011				
	7225808	Mar 01, 2011	DP			
	7225808*PED	Sep 01, 2011				
	7389775	Mar 01, 2011	DP			
	7389775*PED	Sep 01, 2011				
<u>SAPROTERIN DIHYDROCHLORIDE - KUVAN</u>						
022181 001	>A> 7566462	Nov 16, 2025	DP			
	>A> 7566714	Nov 17, 2024		U-989		
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
022350 001					>A> NCE	Jul 31, 2014
<u>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</u>						
022350 002					>A> NCE	Jul 31, 2014
<u>SILDENAFIL CITRATE - REVATIO</u>						
021845 001					I-598	May 07, 2009
<u>SILODOSIN - RAPAFL0</u>						
022206 001	5780485	Nov 13, 2012	U-902			
<u>SILODOSIN - RAPAFL0</u>						
022206 002	5780485	Nov 13, 2012	U-902			
<u>SINECATECHINS - VEREGEN</u>						
021902 001	5795911	Oct 31, 2020	U-172			
<u>SOMATROPIN RECOMBINANT - HUMATROPE</u>						
019640 001					I-585	Mar 12, 2012
<u>SOMATROPIN RECOMBINANT - HUMATROPE</u>						
019640 004	5612315	Mar 18, 2014	DP		I-585	Mar 12, 2012
<u>SOMATROPIN RECOMBINANT - HUMATROPE</u>						
019640 005	5612315	Mar 18, 2014	DP		I-585	Mar 12, 2012
<u>SOMATROPIN RECOMBINANT - HUMATROPE</u>						
019640 006	5612315	Mar 18, 2014	DP		I-585	Mar 12, 2012
<u>SOMATROPIN RECOMBINANT - HUMATROPE</u>						
019640 007	5612315	Mar 18, 2014	DP		I-585	Mar 12, 2012

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u>						
021148 004	5849700	Dec 15, 2015	U-340			
	5849704	Dec 15, 2015	DP U-340			
<u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u>						
021148 005	5849700	Dec 15, 2015	U-340			
	5849704	Dec 15, 2015	DP U-340			
<u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u>						
021148 006	5849700	Dec 15, 2015	U-340			
	5849704	Dec 15, 2015	DP U-340			
<u>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</u>						
021148 007	5849700	Dec 15, 2015	U-340		I-572	Oct 31, 2011
	5849704	Dec 15, 2015	DP U-340		I-551	Sep 20, 2010
					I-536	May 31, 2010
					ODE	May 31, 2014
<u>SOMATROPIN RECOMBINANT - SAIZEN</u>						
019764 002	>A> 5898030	Apr 27, 2016	DP			
<u>SOMATROPIN RECOMBINANT - SAIZEN</u>						
019764 003	>A> 5898030	Apr 27, 2016	DP			
<u>SOMATROPIN RECOMBINANT - SEROSTIM</u>						
020604 001	>A> 5898030	Apr 27, 2016	DP			
<u>SOMATROPIN RECOMBINANT - SEROSTIM</u>						
020604 002	>A> 5898030	Apr 27, 2016	DP			
<u>SOMATROPIN RECOMBINANT - SEROSTIM</u>						
020604 003	>A> 5898030	Apr 27, 2016	DP			
<u>SOMATROPIN RECOMBINANT - SEROSTIM</u>						
020604 004	>A> 5898030	Apr 27, 2016	DP			
<u>SOMATROPIN RECOMBINANT - ZORBTIVE</u>						
021597 004	>A> 5288703	Oct 07, 2011	U-898			
	>A> 5898030	Apr 27, 2016	DP			
<u>SUMATRIPTAN SUCCINATE - SUMATRIPTAN SUCCINATE</u>						
076572 001					PC	Aug 08, 2009
<u>SUMATRIPTAN SUCCINATE - SUMATRIPTAN SUCCINATE</u>						
076840 001					PC	Aug 08, 2009
<u>SUMATRIPTAN SUCCINATE - SUMATRIPTAN SUCCINATE</u>						
076840 002					PC	Aug 08, 2009
<u>SUMATRIPTAN SUCCINATE - SUMATRIPTAN SUCCINATE</u>						
076840 003					PC	Aug 08, 2009
<u>SUMATRIPTAN SUCCINATE - SUMAVEL DOSEPRO</u>						
022239 001	>A> 5891086	Jul 27, 2014	DP			
	>A> 5957886	Mar 08, 2016	DP			
	>A> 6135979	Mar 21, 2017	DP			
<u>SUNITINIB MALATE - SUTENT</u>						
021938 004	6573293	Feb 15, 2021	DS DP U-703		NCE	Jan 26, 2011
	7125905	Feb 15, 2021	DS DP			
	7211600	Dec 22, 2020	U-883			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TADALAFIL - ADCIRCA</u>						
022332 001	5859006	Nov 21, 2017	DS DP U-975		NP	May 22, 2012
	6821975	Nov 19, 2020	DS DP		ODE	May 22, 2016
	7182958	Apr 26, 2020	DP			
<u>TELBIVUDINE - TYZEKA</u>						
022154 001					NCE	Oct 25, 2011
<u>TEMOZOLOMIDE - TEMODAR</u>						
022277 001	5260291	Aug 11, 2013	DS DP U-619			
	5260291*PED	Feb 11, 2014				
	6987108	Sep 08, 2023	DP			
<u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u>						
021318 001	7550434	Dec 08, 2018	DP U-982		>A> I-602	Jul 22, 2012
<u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u>						
021318 002	6770623	Dec 08, 2018	DP U-982		>A> I-602	Jul 22, 2012
	6977077	Aug 19, 2019	U-982			
	7144861	Dec 08, 2018	DP			
	7163684	Aug 19, 2019	U-983			
	7351414	Aug 19, 2019	U-984			
	7550434	Dec 08, 2018	DP U-982			
<u>TIGECYCLINE - TYGACIL</u>						
021821 001					I-588 I-587 I-586	Mar 20, 2012 Mar 20, 2012 Mar 20, 2012
<u>TOLVAPTAN - SAMSCA</u>						
022275 001	5258510	Nov 02, 2010	DS		NCE	May 19, 2014
	5753677	May 19, 2015	U-978			
<u>TOLVAPTAN - SAMSCA</u>						
022275 002	5258510	Nov 02, 2010	DS		NCE	May 19, 2014
	5753677	May 19, 2015	U-978			
<u>TOLVAPTAN - SAMSCA</u>						
022275 003	5258510	Nov 02, 2010	DS		NCE	May 19, 2014
	5753677	May 19, 2015	U-978			
<u>TOPIRAMATE - TOPAMAX</u>						
020505 001	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>						
020505 002	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>						
020505 003	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>						
020505 004	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TOPIRAMATE - TOPAMAX</u>						
020505 005	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>						
020505 006	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>						
020844 001	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX</u>						
020844 002	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPAMAX SPRINKLE</u>						
020844 003	7498311	Oct 13, 2015	U-955			
	7498311*PED	Apr 13, 2016				
<u>TOPIRAMATE - TOPIRAMATE</u>						
076448 001					PC	Oct 12, 2009
<u>TOPIRAMATE - TOPIRAMATE</u>						
076448 002					PC	Oct 12, 2009
<u>TOPIRAMATE - TOPIRAMATE</u>						
077868 001					PC	Oct 12, 2009
<u>TOPIRAMATE - TOPIRAMATE</u>						
077868 002					PC	Oct 12, 2009
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
021745 001					NP	Dec 30, 2011
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
021745 002					NP	Dec 30, 2011
<u>TRAMADOL HYDROCHLORIDE - RYZOLT</u>						
021745 003					NP	Dec 30, 2011
<u>TREPROSTINIL SODIUM - TYVASO</u>						
022387 001					>A> NDF	Jul 30, 2012
<u>TRYPAN BLUE - MEMBRANEBLUE</u>						
022278 001					NCE ODE	Dec 16, 2009 Dec 16, 2011
<u>ZOLEDRONIC ACID - RECLAST</u>						
021817 001					I-595 I-584 I-581	May 29, 2012 Mar 15, 2012 Dec 19, 2011
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
021997 001	6761910	Sep 24, 2018	DP U-674			
<u>ZOLPIDEM TARTRATE - EDLUAR</u>						
021997 002	6761910	Sep 24, 2018	DP U-674			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 7 - July 2009

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
-----------------	-----------	------------------------------	-----------------	-------------------------------	------------------------	-----------------------------------

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

Due to space limitations in the patent and exclusivity columns, abbreviations and references have been developed. Refer to the APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 29th Edition for a full listing of patent and exclusivity terms (Abbreviations, Dosing Schedule, Indications, and Patent Use Codes).

The current complete list of patent terms is available at <http://www.accessdata.fda.gov/scripts/cder/ob/docs/pattermsall.cfm>