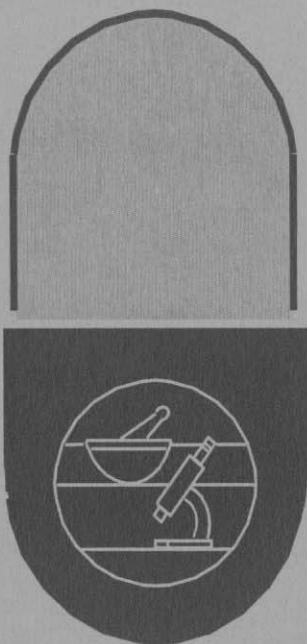


CUMULATIVE
SUPPLEMENT 10
OCTOBER 2001



APPROVED
DRUG PRODUCTS

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

21ST EDITION

Department of Health and Human Services

Food and Drug Administration
Center for Drug Evaluation and Research
Office of Information Technology
Division of Data Management and Services

2001

RM
301.45
.A66
2001
Oct.
suppl.

50
Juse

APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

21ST EDITION

Cumulative Supplement 10

October 2001

CONTENTS

	<i>PAGE</i>
1.0 INTRODUCTION	iii
1.1 How to Use the Cumulative Supplement	iii
1.2 Applicant Name Changes.....	iv
1.3 Availability of the Edition.....	vi
1.4 Report of Counts for the Prescription Drug Product List	viii
1.5 Cumulative Supplement Change Legend.....	x
1.6 Change of a Therapeutic Equivalent Code for a Drug Entity.....	xi
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 <i>in vivo</i> 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**

21ST EDITION

**CUMULATIVE SUPPLEMENT 10
OCTOBER 2001**

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, 21st 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and products that have never been marketed, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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 place an asterisk (*) 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. [Strength(s) which already exist in the List will not be repeated for context.]

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, 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 21st Edition List will then be added to the "Discontinued Drug Product List" appearing in the 22nd Edition. The current edition Section 2. How To Use The Drug Product Lists describes the layout and usage of the List.

The Patent and Exclusivity Lists are arranged in alphabetical order by active ingredient name. For those products with multiple active ingredients, only the first active ingredient (in alphabetical sort) will appear. In addition, the trade name will be displayed to the right of the active ingredient name for each product. 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. 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 in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations.

New additions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >A>. The Patent and Exclusivity Data are indicated by the symbol >ADD> to the left of the line on which new information exists. The >ADD> 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.

1.2 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

APPLICANT NAME CHANGES

<u>FORMER APPLICANT NAME (FORMER ABBREVIATED NAME)</u>	<u>NEW APPLICANT NAME (NEW ABBREVIATED NAME)</u>
BAXTER PHARMACEUTICAL PRODUCTS INC (BAXTER PHARM PROD)	BAXTER HEALTHCARE CORPORATION ANESTHESIA & CRITICAL CARE (BAXTER HLTHCARE CORP)
CAMALL CO INC (CAMALL)	ABC HOLDING CORPORATION (ABC HOLDING)
CIBA VISION CORP DIV NOVARTIS CO (CIBA)	NOVARTIS OPHTHALMICS INC (NOVARTIS)
CIBA VISION OPHTHALMICS (CIBA VISION OPHTHLMC)	NOVARTIS OPHTHALMICS INC (NOVARTIS)
DEY LABORATORIES INC (DEY)	DEY LP (DEY)
KNOLL PHARMACEUTICAL COMPANY (KNOLL PHARM)	ABBOTT LABORATORIES PHARMACEUTICAL PRODUCTS (ABBOTT)
LOTUS BIOCHEMICAL CORPORATION (LOTUS BIOCHEM)	NEW RIVER PHARMACEUTICALS INC (NEW RIVER)
MARSAM PHARMACEUTICALS INC (MARSAM PHARMS)	MARSAM PHARMACEUTICALS LLC (MARSAM PHARMS)
MEDEVA AMERICAS INC (MEDEVA)	CELLTECH PHARMACEUTICALS INC (CELLTECH PHARMS)
MEDEVA PHARMACEUTICALS INC (MEDEVA)	CELLTECH PHARMACEUTICALS INC (CELLTECH PHARMS)
MEDEVA INC (MEDEVA)	CELLTECH PHARMACEUTICALS INC (CELLTECH PHARMS)
MEDEVA PHARMACEUTICALS CA INC (MEDEVA PHARMS CA)	CELLTECH MANUFACTURING CA INC (CELLTECH MFG CA INC)
MEDEVA PHARMACEUTICALS MA INC (MEDEVA PHARMS MA)	CELLTECH MANUFACTURING INC (CELLTECH MFG)
NOVOPHARM LTD (NOVOPHARM)	TEVA PHARMACEUTICALS USA (TEVA)
NOVOPHARM PHARMACEUTICAL CO (NOVOPHARM PHARM)	TEVA PHARMACEUTICALS USA (TEVA)
NOVOPHARM NC INC (NOVOPHARM NC)	TEVA PHARMACEUTICALS USA (TEVA)

APPLICANT NAME CHANGES

<u>FORMER APPLICANT NAME (FORMER ABBREVIATED NAME)</u>	<u>NEW APPLICANT NAME (NEW ABBREVIATED NAME)</u>
OHMEDA PHARMACEUTICAL PRODUCTS DIV (OHMEDA)	BAXTER HEATHCARE CORPORATION ANESTHESIA & CRITICAL CARE (BAXTER HLTHCARE CORP)
ROBERTS LABORATORIES INC (ROBERTS LABS)	SHIRE PHARMACEUTICAL DEVELOPMENT INC (SHIRE PHARM)
ROBERTS PHARMACEUTICAL CORP (ROBERTS PHARM)	SHIRE PHARMACEUTICAL DEVELOPMENT INC (SHIRE PHARM)
ZENITH GOLDLINE (ZENITH GOLDLINE)	IVAX PHARMACEUTICALS INC (IVAX PHARMS)
ZENITH GOLDLINE PHARMACEUTICALS INC (ZENITH GOLDLINE)	IVAX PHARMACEUTICALS INC (IVAX PHARMS)

1.3 AVAILABILITY OF THE EDITION

The 21st Edition of the Orange Book and its monthly cumulative supplements are available by subscription from the Government Printing Office:

Superintendent of Documents
Government Printing Office
P.O. Box 371954
Pittsburgh, PA 15250-7954

The telephone number to charge your subscription is 202-512-1800. The cost is \$101.00 annually.

The Approved Drug Products with Therapeutic Equivalence Evaluation (Orange Book) and related drug information is also available on the Internet at the Food and Drug Administration, Center for Drug Evaluation and Research, Drug Info page.

There is an Electronic Orange Book Query (EOB) at <http://www.fda.gov/cder/ob>. The Query provides searching of the approved drug list by active ingredient, proprietary name, applicant holder or applicant 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. The data is updated concurrently with the publication of the annual edition or monthly cumulative supplements.

The Internet version of the hard copy Orange Book annual edition is at
<http://www.fda.gov/cder/orange/adp.htm>.

The Internet version of the hard copy monthly supplement is at
<http://www.fda.gov/cder/orange/supplement/cspreface.htm>. Changes to the annual edition are listed separately by month.

There are ASCII text files of the Orange Book drug product data at
<http://www.fda.gov/cder/orange/obreadme.htm>. The drug product text files are zipped into zipobtxt.exe. The files are updated concurrently with the publication of the annual edition or monthly cumulative supplements. Appendix A and Appendix B are updated quarterly.

The 21st annual edition of the 2000 Orange Book Patent and Exclusivity List is at
<http://www.fda.gov/cder/orange/21bookpub.pdf>.

The current year Patent and Exclusivity cumulative supplement list that denotes the current month additions is at <http://www.fda.gov/cder/orange/supplement/patents.pdf>.

The Patent Term Extension and new Patents, Docket Number *95S-0117, is at
<http://www.fda.gov/cder/orange/docket.pdf>. It is updated monthly as soon as available and as otherwise needed.

The Drug Price Competition and Patent Term Restoration Act requires that patent information be filed with all newly submitted Section 505 drug applications. To facilitate industry submission of the information, a patent submission sample format is available in HTML and PDF format at:

<http://www.fda.gov/cder/orange/patdecl.pdf>
<http://www.fda.gov/cder/orange/patdecl.html>

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

1.4 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 2000) 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 2000</u>	<u>MAR 2001</u>	<u>JUN 2001</u>	<u>SEP 2001</u>
DRUG PRODUCTS LISTED	10360	10372	10155	10094
SINGLE SOURCE	2682 (25.9%)	2696 (26.0%)	2665 (26.2%)	2643 (26.2%)
MULTISOURCE	7568 (73.1%)	7566 (72.9%)	7380 (72.7%)	7341 (72.7%)
THERAPEUTICALLY EQUIVALENT	7257 (70.0%)	7263 (70.0%)	7078 (69.7%)	7050 (69.8%)
NOT THERAPEUTICALLY	311 (3.0%)	303 (2.9%)	302 (3.0%)	291 (2.9%)
EQUIVALENT EXCEPTIONS ¹	110 (1.1%)	110 (1.1%)	110 (1.1%)	110 (1.1%)
NEW MOLECULAR ENTITIES APPROVED	2	6	3	4
NUMBER OF APPLICANTS	594	582	579	572

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

1.5 CUMULATIVE SUPPLEMENT LEGEND

The 21st Edition Orange book (OB) Cumulative Supplement (CS) layout has changed. The new format follows the Annual Edition and previous CS format. The List is sorted by Ingredient(s) and, within each grouping, by the Dosage Form;Route and then by trade name. The manner of displaying the individual product information has changed.

The individual product record follows the previous format layout for Therapeutic Equivalence Code, Reference Listed Drug symbol, applicant holder, strength(s), New Drug Approval number, product number, and approval date. Two new columns have been added to provide more information. 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 proceeded 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.
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.
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.
WDAG	Withdrawn. The applicant holder has notified the FDA in writing that the product is no longer being marketed resulting in the product approval being withdrawn by mutual agreement. The product will be listed in the Discontinued Section.
WDRP	Withdrawn. The application approval has been withdrawn for failure to provide Annual Reports. The product will be moved to the Discontinued Section in the next edition.

1.6 CHANGE OF A THERAPEUTIC EQUIVALENT CODE FOR A DRUG ENTITY

Metaxalone tablets were reviewed in the Drug Efficacy Study Implementation program. FDA published a Federal Register notice on August 15, 1974 (39 FR 29396) finding metaxalone tablets to be effective in the treatment of discomfort associated with acute, painful musculo-skeletal conditions. The Federal Register notice did not require the conduct of a bioavailability/bioequivalence study as a condition of marketing.

On March 6, 2001, URL Mutual Pharmaceutical Co. Inc submitted a citizen petition (Docket No. 01P-0117/CP1) asking FDA to reclassify the drug product metaxalone tablets from one not presenting bioequivalence problems to one that requires an in vivo demonstration of bioequivalence as a condition of approval for an ANDA. To support these assertions, the petition included results of two in vivo bioequivalence fasting studies and three separate in vitro dissolution tests. After a careful review of the data submitted by Mutual, the agency agreed that the firm has demonstrated a lack of correlation between in vitro dissolution and in vivo bioequivalence data of oral metaxalone tablets in two bioequivalence studies. The failure of both Mutual formulations to meet the 90% confidence intervals further supports the lack of in vitro/in vivo correlations.

Therefore, in accordance with our policy as enunciated in Section 1.9 of the Introduction to the 21st Edition of the Orange Book, we are providing a 60 day period in which interested parties may submit comments. The closing date for the comments will be November 30, 2001. The comments should be sent to the Director, Division of Bioequivalence, Office of Generic Drugs, Center for Drug Evaluation and Research, (MPN-2) HFD-650, 7500 Standish Place, Rockville, MD 20855. These comments should include scientific data either supporting or disagreeing with our proposal to change the therapeutic equivalence category for metaxalone tablets from a "non bioproblem" to a "bioproblem" drug.

PREScription DRUG PRODUCT LIST - 21ST EDITION
RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 10 - OCT 2001

1-1

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE

@ MIKART	150MG;180MG;15MG	N81095 001	OCT 26, 1990	MAY	DISC
@	150MG;180MG;60MG	N81097 001	OCT 26, 1990	MAY	DISC

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

TRIAPRIN

@ DUNHALL	325MG;50MG	N89268 001	JUL 02, 1987	FEB	WDRP
-----------	------------	------------	--------------	-----	------

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ANOQUAN

@ ROBERTS AND HAUCK	325MG;50MG;40MG	N87628 001	OCT 01, 1986	FEB	WDRP
---------------------	-----------------	------------	--------------	-----	------

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

AB ABLE	325MG;50MG;40MG	N40390 001	JUL 23, 2001	JUL	NEWA
AB	500MG;50MG;40MG	N40394 001	JUL 23, 2001	JUL	NEWA

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL; ACETAMINOPHEN; AND CAFFEINE WITH CODEINE PHOSPHATE

AB WEST WARD	325MG;50MG;40MG;30MG	N75618 001	MAR 23, 2001	MAR	NEWA
FIORICET W/ CODEINE					

AB + NOVARTIS	325MG;50MG;40MG;30MG	N20232 001	JUL 30, 1992	MAR	CFTG
PHRENILIN WITH CAFFEINE AND CODEINE					

AB AMARIN PHARMS	325MG;50MG;40MG;30MG	N74911 001	AUG 22, 2001	AUG	NEWA
------------------	----------------------	------------	--------------	-----	------

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

>D> AA MIKART	356.4MG;30MG;16MG	N40109 001	AUG 26, 1997	OCT	CRLD
>A> +	356.4MG;30MG;16MG	N40109 001	AUG 26, 1997	OCT	CRLD

>D> DHC PLUS

>D> AA + PURDUE FREDERICK	356.4MG;30MG;16MG	N88584 001	MAR 04, 1986	OCT	DISC
>A> @	356.4MG;30MG;16MG	N88584 001	MAR 04, 1986	OCT	DISC

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

+ MIKART	712.8MG;60MG;32MG	N40316 001	APR 28, 1999	JAN	CTNA
----------	-------------------	------------	--------------	-----	------

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA MALLINCKRODT	300MG;15MG	N40419 001	MAY 31, 2001	MAY	NEWA
AA	300MG;30MG	N40419 002	MAY 31, 2001	MAY	NEWA

AA	300MG;60MG	N40419 003	MAY 31, 2001	MAY	NEWA
----	------------	------------	--------------	-----	------

CAPITAL WITH CODEINE

@ CARNRICK	325MG;30MG	N83643 001	MAY 31, 1974	FEB	WDRP
------------	------------	------------	--------------	-----	------

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	MALLINCKRODT	500MG/15ML;7.5MG/15ML	N40418 001	JUN 27, 2001	JUN	NEWA
AA	+ MIKART	500MG/15ML;7.5MG/15ML	N81051 001	AUG 28, 1992	JUN	CDFR
	+ +	500MG/15ML;5MG/15ML	N81226 001	OCT 27, 1992	JUN	CDFR
AA	PHARM ASSOC	500MG/15ML;5MG/15ML	N89557 001	APR 29, 1992	JUN	CDFR
	TABLET; ORAL	500MG/15ML;7.5MG/15ML	N40182 001	MAR 13, 1998	JUN	CDFR
	+ ENDO PHARMS	400MG;5MG	N40288 001	NOV 27, 1998	SEP	CTEC
	+ +	400MG;7.5MG	N40288 002	NOV 27, 1998	SEP	CTEC
	+ +	400MG;10MG	N40288 003	NOV 27, 1998	SEP	CTEC
	+ WATSON LABS	325MG;7.5MG	N40248 001	APR 28, 2000	JUL	DISC
	@ @	325MG;7.5MG	N40248 001	APR 28, 2000	AUG	DISC
	+ +	750MG;10MG	N40094 004	MAR 22, 1999	APR	NEWA
	LORTAB					
AA	+ WATSON LABS	325MG;5MG	N40099 001	JUN 25, 1997	JAN	CAHN
	NORCO					
AA	WATSON LABS	325MG;7.5MG	N40148 003	SEP 12, 2000	APR	NEWA
AA	+ +	325MG;7.5MG	N40148 003	SEP 12, 2000	JUL	CRLD

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

AB	ABLE	650MG;100MG	N75838 001	JUL 11, 2001	JUL	NEWA
	@ HALSEY	325MG;50MG	N70115 001	JUN 12, 1985	MAY	DISC
	@ @	650MG;100MG	N70116 001	JUN 12, 1985	MAY	DISC
AB	MALLINCKRODT	650MG;100MG	N75738 001	FEB 02, 2001	FEB	NEWA
AB	VINTAGE PHARMS	325MG;50MG	N74843 002	FEB 15, 2001	FEB	NEWA

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

ULTRACET

+ +	JOHNSON RW	325MG;37.5MG
-----	------------	--------------

N21123 001 AUG 15, 2001 AUG NEWA

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

AP	GENSIA SICOR PHARMS	EQ 50MG BASE/ML	N75627 001	MAR 28, 2001	MAR	NEWA
	ACYCLOVIR SODIUM					
>D>	AP AESGEN	EQ 500MG BASE/VIAL	N75015 001	APR 30, 1998	OCT	CAHN

>A>	AP AM PHARM PARTNERS	EQ 500MG BASE/VIAL	N75015 001	APR 30, 1998	OCT	CAHN
-----	----------------------	--------------------	------------	--------------	-----	------

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

>D>	AB ALPHARMA	0.09MG/INH	N73045 001	AUG 19, 1997	OCT	CAHN
	AB ARMSTRONG PHARMS	0.09MG/INH	N72273 001	AUG 14, 1996	JUN	CAHN
>A>	AB GENPHARM	0.09MG/INH	N73045 001	AUG 19, 1997	OCT	CAHN

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION VENTOLIN HFA		
+ GLAXO	EQ 0.09MG BASE/INH	N20983 001 APR 19, 2001 APR NEWA
CAPSULE; INHALATION VENTOLIN ROTACAPS @ GLAXO WELLCOME	EQ 0.2MG BASE	N19489 001 MAY 04, 1988 JUL DISC
SOLUTION; INHALATION ACCUNEB		
+ DEY	EQ 0.021% BASE	N20949 002 APR 30, 2001 APR NEWA
+	EQ 0.042% BASE	N20949 001 APR 30, 2001 APR NEWA
ALBUTEROL SULFATE AN NEPHRON	EQ 0.5% BASE	N75664 001 JUN 26, 2001 JUN NEWA
AN ROXANE VENTOLIN @ GLAXO WELLCOME	EQ 0.083% BASE EQ 0.5% BASE	N75129 001 FEB 13, 2001 FEB NEWA N19773 001 APR 23, 1992 JUL DISC
@		N19269 002 JAN 16, 1987 JUL DISC
TABLET; ORAL @ GLAXO WELLCOME	EQ 2MG BASE	N19112 001 JUL 10, 1986 JUN DISC
@	EQ 4MG BASE	N19112 002 JUL 10, 1986 JUN DISC

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION DUONEB		
+ DEY	EQ 0.083% BASE;0.017%	N20950 001 MAR 21, 2001 MAR NEWA

ALLOPURINOL

TABLET; ORAL ZYLOPRIM		
AB PROMETHEUS LABS	100MG	N16084 001 AUG 19, 1966 MAY CAHN
AB +	300MG	N16084 002 JAN 14, 1974 MAY CAHN

ALMOTRIPTAN MALATE

TABLET; ORAL AXERT		
PHARMACIA AND UPJOHN	EQ 6.25MG BASE	N21001 001 MAY 07, 2001 MAY NEWA
+	EQ 12.5MG BASE	N21001 002 MAY 07, 2001 MAY NEWA

ALPRAZOLAM

SOLUTION; ORAL ALPRAZOLAM @ ROXANE	0.5MG/5ML	N74314 001 OCT 31, 1993 SEP DISC
--	-----------	----------------------------------

AMIKACIN SULFATE

INJECTABLE; INJECTION AMIKACIN SULFATE @ ABBOTT	EQ 250MG BASE/ML	N63265 001 NOV 30, 1994 APR DISC
@	EQ 250MG BASE/ML	N63266 001 OCT 31, 1994 APR DISC
@	EQ 250MG BASE/ML	N64099 001 JUN 20, 1995 MAY DISC
@ ELKINS SINK	EQ 250MG BASE/ML	N63275 001 MAY 18, 1992 APR DISC

AMINOCAPROIC ACID

TABLET; ORAL				
AMICAR				
AB + IMMUNEX	500MG		N15197 001	JUN 03, 1964
AMINOCAPROIC			MAY	CFTG
AB MIKART	500MG		N75602 001	MAY 24, 2001
			MAY	NEWA

AMIODARONE HYDROCHLORIDE

TABLET; ORAL				
AMIODARONE HCL				
AB BARR	200MG		N75389 001	JAN 25, 2001
AB TARO	200MG		N75424 001	MAR 30, 2001
			MAR	NEWA

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL				
AMITRIPTYLINE HCL				
@ TEVA	75MG		N85030 001	NOV 22, 1976
			JUL	DISC

AMLEXANOX

PASTE; DENTAL				
APHTHASOL				
+ GLAXOSMITHKLINE CONS	5%		N20511 001	DEC 17, 1996
			SEP	CAHN

AMOXICILLIN

CAPSULE; ORAL				
AMOXICILLIN				
@ LABS ATRAL	250MG		N62528 001	AUG 07, 1985
@	500MG		N62528 002	AUG 07, 1985
@ MYLAN	250MG		N62067 001	AUG 14, 1980
@	500MG		N62067 002	AUG 14, 1980
@ TEVA	250MG		N63030 001	FEB 28, 1989
@	500MG		N63031 001	FEB 28, 1989
TRIMOX				APR DISC
@ APOTHECON	250MG		N63099 001	MAR 20, 1992
@	500MG		N63099 002	MAR 20, 1992
WYMOX				APR DISC
@ WYETH AYERST	250MG		N62120 001	APR 28, 1978
@	500MG		N62120 002	APR 28, 1978
FOR SUSPENSION; ORAL				APR DISC
TRIMOX				
@ APOTHECON	50MG/ML		N61886 001	DEC 09, 1974
@	125MG/5ML		N61886 002	DEC 09, 1974
@	250MG/5ML		N61886 003	DEC 09, 1974
			MAY	DISC
			MAY	DISC
			MAY	DISC

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL				
AUGMENTIN ES-600				
+ GLAXOSMITHKLINE	600MG/5ML;EQ 42.9MG		N50755 001	JUN 22, 2001
	BASE/5ML		JUN	NEWA

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE;DEXTROAMPHETAMINE SULFATE

>A>	CAPSULE, EXTENDED RELEASE; ORAL						
>A>	ADDERALL XR 10						
>A>	SHIRE LABS	2.5MG;2.5MG;2.5MG;2.5MG	N21303	001	OCT 11, 2001	OCT	NEWA
>A>	ADDERALL XR 20						
>A>	SHIRE LABS	5MG;5MG;5MG;5MG	N21303	002	OCT 11, 2001	OCT	NEWA
>A>	ADDERALL XR 30						
>A>	+ SHIRE LABS	7.5MG;7.5MG;7.5MG;7.5MG	N21303	003	OCT 11, 2001	OCT	NEWA
	TABLET; ORAL						
	ADDERALL 7.5						
	SHIRE LABS	1.875MG;1.875MG;1.875MG;1. 875MG	N11522	011	AUG 31, 2000	APR	CRLD

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

⑧ ABBOTT	50MG/VIAL	N64141	001	DEC 23, 1996	MAY	DISC
INJECTABLE, LIPID COMPLEX; INJECTION						
AMPHOTEC						
+ INTERMUNE PHARMS	50MG/VIAL	N50729	001	NOV 22, 1996	FEB	CAHN
+	100MG/VIAL	N50729	002	NOV 22, 1996	FEB	CAHN

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

⑧ ELKINS SINK	EQ 125MG BASE/VIAL	N62692	001	JUN 24, 1986	MAY	DISC
⑧	EQ 250MG BASE/VIAL	N62692	002	JUN 24, 1986	MAY	DISC
⑧	EQ 500MG BASE/VIAL	N62692	003	JUN 24, 1986	MAY	DISC
⑧	EQ 1GM BASE/VIAL	N62692	004	JUN 24, 1986	MAY	DISC
⑧	EQ 2GM BASE/VIAL	N62692	005	JUN 24, 1986	MAY	DISC
⑧	EQ 10GM BASE/VIAL	N62692	006	JUN 24, 1986	MAY	DISC
⑧ HANFORD GC	EQ 125MG BASE/VIAL	N63143	001	APR 15, 1993	APR	DISC
⑧	EQ 250MG BASE/VIAL	N63145	001	APR 15, 1993	APR	DISC
⑧	EQ 500MG BASE/VIAL	N63146	001	APR 15, 1993	APR	DISC
⑧	EQ 500MG BASE/VIAL	N63147	001	APR 15, 1993	APR	DISC
⑧	EQ 1GM BASE/VIAL	N62772	001	APR 15, 1993	MAY	DISC
⑧	EQ 1GM BASE/VIAL	N63139	001	APR 15, 1993	APR	DISC
⑧	EQ 2GM BASE/VIAL	N63140	001	APR 15, 1993	APR	DISC
⑧	EQ 2GM BASE/VIAL	N63141	001	APR 15, 1993	APR	DISC
⑧	EQ 10GM BASE/VIAL	N63142	001	APR 15, 1993	APR	DISC
⑧ IBI	EQ 125MG BASE/VIAL	N62797	001	JUL 12, 1993	MAY	DISC
⑧	EQ 2GM BASE/VIAL	N62797	002	JUL 12, 1993	MAY	DISC
⑧ MARSAM	EQ 10GM BASE/VIAL	N62994	001	SEP 15, 1988	JUL	DISC
OMNIPEN-N						
⑧ WYETH AYERST	EQ 125MG BASE/VIAL	N62718	001	DEC 16, 1986	MAY	DISC
⑧	EQ 250MG BASE/VIAL	N62718	002	DEC 16, 1986	MAY	DISC

⑧	EQ 500MG BASE/VIAL	N62718 003 DEC 16, 1986 MAY DISC
⑧	EQ 1GM BASE/VIAL	N62718 004 DEC 16, 1986 MAY DISC
⑧	EQ 2GM BASE/VIAL	N62718 005 DEC 16, 1986 MAY DISC
TOTACILLIN-N		
SMITHKLINE BEECHAM	EQ 10GM BASE/VIAL	N60677 006 MAY 04, 1976 JUL CTEC

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

⑧ BIOCHEMIE

EQ 250MG BASE

N64082 001 AUG 29, 1995 MAY DISC

⑧

EQ 500MG BASE

N64082 002 AUG 29, 1995 MAY DISC

FOR SUSPENSION; ORAL

⑧ MYLAN

EQ 125MG BASE/5ML

N61829 002 JUL 29, 1974 MAY DISC

⑧

EQ 250MG BASE/5ML

N61829 001 JUL 29, 1974 MAY DISC

TOTACILLIN

⑧ SMITHKLINE BEECHAM

EQ 125MG BASE/5ML

N60666 001 MAY 07, 1970 FEB WDRP

⑧

EQ 250MG BASE/5ML

N60666 002 MAY 07, 1970 FEB WDRP

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

FOR SUSPENSION; ORAL

PROBAMPACIN

⑧ TEVA

EQ 3.5GM BASE/BOT;1GM/BOT

N61741 001 OCT 10, 1973 MAY DISC

ARBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

GENESA

⑧ GENSIA AUTOMEDICS

0.05MG/ML

N20420 001 SEP 12, 1997 MAR DISC

ARDEPARIN SODIUM

INJECTABLE; INJECTION

NORMIFLO

⑧ PHARMACIA AND UPJOHN

5,000 UNITS/0.5ML

N20227 002 MAY 23, 1997 JUL CAHN

⑧

10,000 UNITS/0.5ML

N20227 001 MAY 23, 1997 JUL CAHN

⑧ WYETH AYERST

5,000 UNITS/0.5ML

N20227 002 MAY 23, 1997 MAY DISC

⑧

10,000 UNITS/0.5ML

N20227 001 MAY 23, 1997 MAY DISC

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

INJECTABLE; IV (INFUSION)

INFUVITE PEDIATRIC

+ SABEX

80MG/VIAL;0.02MG/VIAL;400

N21265 001 FEB 21, 2001 FEB NEWA

IU/VIAL;0.001MG/VIAL;5MG/V

IAL;0.14MG/VIAL;17MG/VIAL;

1MG/VIAL;1.4MG/VIAL;1.2MG/

VIAL;7 IU/VIAL;2,300

IU/VIAL;0.2MG/VIAL

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

FOR SOLUTION; IV (INFUSION)

M.V.I. PEDIATRIC

+ ASTRazeneca

80MG/VIAL;0.02MG/VIAL;0.00

>D>

1MG/VIAL;5MG/VIAL;0.01MG/V
 IAL;0.14MG/VIAL;17MG/VIAL;
 0.2MG/VIAL;1MG/VIAL;1.4MG/
 VIAL;EQ 1.2MG
 BASE/VIAL;0.7MG/VIAL;7MG/V
 IAL N18920 001 SEP 21, 2000 OCT CAHN
 +
 80MG/VIAL;0.02MG/VIAL;0.00
 1MG/VIAL;5MG/VIAL;0.01MG/V
 IAL;0.14MG/VIAL;17MG/VIAL;
 0.2MG/VIAL;1MG/VIAL;1.4MG/
 VIAL;EQ 1.2MG
 BASE/VIAL;0.7MG/VIAL;7MG/V
 IAL N18920 001 SEP 21, 2000 FEB NEWA
 >A> + NEOSAN PHARMS
 80MG/VIAL;0.02MG/VIAL;0.00
 1MG/VIAL;5MG/VIAL;0.01MG/V
 IAL;0.14MG/VIAL;17MG/VIAL;
 0.2MG/VIAL;1MG/VIAL;1.4MG/
 VIAL;EQ 1.2MG
 BASE/VIAL;0.7MG/VIAL;7MG/V
 IAL N18920 001 SEP 21, 2000 OCT CAHN
ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID;
NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE
HYDROCHLORIDE; VITAMIN A; VITAMIN E
 INJECTABLE; INJECTION
 M.V.I.-12
 >D> AP + ASTRAZENECA
 10MG/ML;0.006MG/ML;0.5UGM/
 ML;1.5MG/ML;20
 IU/ML;0.04MG/ML;4MG/ML;0.4
 MG/ML;0.36MG/ML;0.3MG/ML;3
 30 UNITS/ML;1 IU/ML N08809 004 AUG 08, 1985 OCT CAHN
 >A> AP + NEOSAN PHARMS
 10MG/ML;0.006MG/ML;0.5UGM/
 ML;1.5MG/ML;20
 IU/ML;0.04MG/ML;4MG/ML;0.4
 MG/ML;0.36MG/ML;0.3MG/ML;3
 30 UNITS/ML;1 IU/ML N08809 004 AUG 08, 1985 OCT CAHN
 MVC PLUS
 @ STERIS
 10MG/ML;0.006MG/ML;0.5UGM/
 ML;1.5MG/ML;20
 IU/ML;0.04MG/ML;4MG/ML;0.4
 MG/ML;0.36MG/ML;0.3MG/ML;3
 30 UNITS/ML;1 IU/ML N18439 002 AUG 08, 1985 SEP DISC
ASPIRIN; BUTALBITAL; CAFFEINE
 TABLET; ORAL
 LANORINAL
 @ LANNETT 325MG;50MG;40MG N86986 002 OCT 18, 1985 JUL DISC
ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE
 TABLET; ORAL
 INVAGESIC
 AB GENEVA PHARMS TECH 385MG;30MG;25MG N74817 001 NOV 27, 1996 JAN CAHN
 INVAGESIC FORTE
 AB GENEVA PHARMS TECH 770MG;60MG;50MG N74817 002 NOV 27, 1996 JAN CAHN

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

PROPOXYPHENE COMPOUND 65

@ EON

389MG;32.4MG;65MG

N80044 002 SEP 16, 1983 MAY DISC

PROPOXYPHENE COMPOUND-65

@ GENEVA PHARMS

389MG;32.4MG;65MG

N83101 002 JUN 24, 1985 MAY DISC

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

>D> ROXIPRIN

>D> AA ROXANE

325MG;4.5MG;0.38MG

N87743 001 JUN 04, 1982 OCT DISC

>A> @

325MG;4.5MG;0.38MG

N87743 001 JUN 04, 1982 OCT DISC

ATENOLOL

TABLET; ORAL

ATENOLOL

@ GENPHARM

25MG

N74126 003 AUG 26, 1998 JUL DISC

@

50MG

N74126 001 MAR 23, 1994 JUL DISC

@

100MG

N74126 002 MAR 23, 1994 JUL DISC

ATORVASTATIN CALCIUM

TABLET; ORAL

LIPITOR

PFIZER

EQ 10MG BASE

N20702 001 DEC 17, 1996 MAR CAHN

EQ 20MG BASE

N20702 002 DEC 17, 1996 MAR CAHN

EQ 40MG BASE

N20702 003 DEC 17, 1996 MAR CAHN

+

EQ 80MG BASE

N20702 004 APR 07, 2000 MAR CAHN

ATROPINE SULFATE

INJECTABLE; IM-IV-SC

ATROPINE SULFATE ANSYR PLASTIC SYRINGE

ABBOTT

0.05MG/ML

N21146 002 JUL 09, 2001 JUL NEWA

+

0.1MG/ML

N21146 001 JUL 09, 2001 JUL NEWA

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HCL AND ATROPINE SULFATE

AA	INWOOD LABS	0.025MG;2.5MG	N85509 001	MAR 09, 1978	FEB	WDRP
AA	LANNETT	0.025MG;2.5MG	N85372 001	FEB 21, 1978	AUG	CMFD
AA	R AND S PHARMA	0.025MG;2.5MG	N85035 001	JUL 05, 1977	MAY	DISC
AA	WEST WARD	0.025MG;2.5MG	N87765 001	MAR 15, 1982	JUL	DISC
AA	EON	0.025MG;2.5MG	N86173 001	AUG 28, 1981	AUG	CMFD
	PVT FORM	0.025MG;2.5MG	N85766 001	DEC 22, 1978	MAY	DISC

AURANOFIN

CAPSULE; ORAL

RIDaura

+ PROMETHEUS LABS

3MG

N18689 001 MAY 24, 1985 MAY CAHN

AZATHIOPRINE

TABLET; ORAL						
IMURAN						
@ PROMETHEUS LABS	25MG		N16324 002	MAR 21, 1980	MAY	CAHN
AB +	50MG		N16324 001	MAR 20, 1968	MAY	CAHN

AZELASTINE HYDROCHLORIDE

SPRAY, METERED; NASAL						
ASTELIN						
>D> + WALLACE LABS	EQ 0.125MG BASE/SPRAY		N20114 001	NOV 01, 1996	OCT	CAHN
>A> + WALLACE PHARMS	EQ 0.125MG BASE/SPRAY		N20114 001	NOV 01, 1996	OCT	CAHN

AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATE

FOR SUSPENSION; TABLET; ORAL						
TROVAN/ZITHROMAX COMPLIANCE PAK						
@ PFIZER	EQ 1GM BASE;EQ 100MG BASE		N50762 001	DEC 18, 1998	MAY	DISC

BACITRACIN ZINC

POWDER; FOR RX COMPOUNDING						
ZIBA-RX						
@ PHARMA TEK	500,000 UNITS/BOT		N61737 001	APR 26, 1973	MAY	DISC

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC						
NEO-POLYCIN						
@ DOW PHARM	500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM		N60647 001	APR 19, 1954	FEB	WDRP

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL						
CORZIDE						
KING PHARMS	5MG;40MG		N18647 001	MAY 25, 1983	AUG	CAHN
+	5MG;80MG		N18647 002	MAY 25, 1983	AUG	CAHN

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION						
EMETE-CON						
+ PFIZER	EQ 50MG BASE/VIAL		N16820 001	MAR 20, 1974	MAY	CAHN

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL						
BETAMETHASONE DIPROPIONATE						
@ CLAY PARK	EQ 0.05% BASE		N74579 001	NOV 26, 1997	APR	DISC
DISC; TOPICAL						
DIPROSONE						
@ SCHERING	EQ 0.1% BASE		N17829 001	MAY 24, 1977	AUG	DISC

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL						
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE						
AB ALTANA	EQ 0.05% BASE;1%		N75502 001	JUN 05, 2001	JUN	NEWA
AB TARO	EQ 0.05% BASE;1%		N75673 001	MAY 29, 2001	MAY	NEWA

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL
LOTRISONE
AB + SCHERING EQ 0.05% BASE; 1% N18827 001 JUL 10, 1984 MAY CFTG

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
BETAXOLOL HCL
AT BAUSCH AND LOMB EQ 0.5% BASE N75630 001 APR 12, 2001 APR NEWA

BETHANECHOL CHLORIDE

INJECTABLE; INJECTION
URECHOLINE
@ SIDMAK LABS 5MG/ML N06536 001 OCT 12, 1948 AUG CAHN
TABLET; ORAL
DUVOID
@ WELLSPRING PHARM 10MG N86262 001 MAR 22, 1978 JUN CAHN
@ 25MG N86263 001 MAR 22, 1978 JUN CAHN
@ 50MG N85882 003 MAR 22, 1978 JUN CAHN
URECHOLINE
@ SIDMAK LABS 5MG N06536 003 FEB 03, 1949 AUG CAHN
@ 10MG N06536 002 OCT 12, 1948 AUG CAHN
@ 25MG N06536 004 OCT 12, 1948 AUG CAHN
@ 50MG N06536 005 JUN 24, 1980 AUG CAHN

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC
LUMIGAN
+ ALLERGAN 0.03% N21275 001 MAR 16, 2001 MAR NEWA

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL
HEЛИДАС

+ PROMTHEUS LABS 262.4MG;250MG;500MG N50719 001 AUG 15, 1996 MAY DISC

BISOPROLOL FUMARATE

TABLET; ORAL
BISOPROLOL FUMARATE

AB COPLEY PHARM 5MG N75644 001 JUN 26, 2001 JUN NEWA
AB 10MG N75644 002 JUN 26, 2001 JUN NEWA

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

@ APOTHECON	2.5MG;6.25MG	N75642 002 DEC 27, 2000 JUN DISC
@	5MG;6.25MG	N75642 001 DEC 27, 2000 JUN DISC
@	10MG;6.25MG	N75642 003 DEC 27, 2000 JUN DISC
AB TEVA	2.5MG;6.25MG	N75686 001 JAN 19, 2001 JAN NEWA
AB	5MG;6.25MG	N75686 002 JAN 19, 2001 JAN NEWA
AB	10MG;6.25MG	N75686 003 JAN 19, 2001 JAN NEWA

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN

>A>	AP	BEDFORD	EQ 15 UNITS BASE/VIAL	N65042 002	OCT 17, 2001	OCT	NEWA
>A>	AP		EQ 30 UNITS BASE/VIAL	N65042 001	OCT 17, 2001	OCT	NEWA

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN

@ ALLERGAN

0.5%

N20490 001 MAR 13, 1997 APR DISC

ALPHAGAN P

+ ALLERGAN

0.15%

N21262 001 MAR 16, 2001 MAR NEWA

BUDESONIDE

CAPSULE; ORAL

>A> ENTOCORT EC

>A> + ASTRAZENECA

3MG

N21324 001 OCT 02, 2001 OCT NEWA

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; SPINAL

MARCaine

AP + ABBOTT

0.75%

N18692 001 MAY 04, 1984 JUN CAHN

BUPRENORPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPRENEX

AP + RECKITT BENCKISER

EQ 0.3MG BASE/ML

N18401 001 DEC 29, 1981 JUL CAHN

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

WELLBUTRIN SR

GLAXO WELLCOME

50MG

N20358 001 OCT 04, 1996 APR CTEC

100MG

N20358 002 OCT 04, 1996 APR CTEC

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPAR

AB BRISTOL MYERS SQUIBB

5MG

N18731 001 SEP 29, 1986 MAR CFTG

AB

10MG

N18731 002 SEP 29, 1986 MAR CFTG

AB

15MG

N18731 003 APR 22, 1996 MAR NEWA

AB +

30MG

N18731 004 APR 22, 1996 JUN CFTG

BUSPIRONE HCL

AB DANBURY PHARMA

5MG

N74253 001 MAR 28, 2001 MAR NEWA

AB

10MG

N74253 002 MAR 28, 2001 MAR NEWA

AB MYLAN

15MG

N75272 003 MAR 28, 2001 MAR NEWA

AB MYLAN TECHNOLOGIES

30MG

N76008 001 JUN 28, 2001 JUN NEWA

AB PAR PHARM

7.5MG

N75467 002 MAR 28, 2001 MAR NEWA

BUTABARBITAL SODIUM

TABLET; ORAL

BUTISOL SODIUM

+ WALLACE LABS

15MG

N00793 002 JUN 05, 1939 MAY CTEC

BUTABARBITAL SODIUM

TABLET; ORAL
 SODIUM BUTABARBITAL
 @ LANNETT 15MG N85849 001 AUG 21, 1978 MAY DISC
 @ 30MG N85866 001 JUL 20, 1978 MAY DISC

BUTOCONAZOLE NITRATE

CREAM; VAGINAL
 GYNIAZOLE-1
 + KV PHARM 2% N19881 001 FEB 07, 1997 SEP CTNA

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION
 BUTORPHANOL TARTRATE
 >A> AP 2MG/ML N75697 001 OCT 23, 2001 OCT NEWA
 BUTORPHANOL TARTRATE PRESERVATIVE FREE
 >A> AP APOTEX 1MG/ML N75695 001 OCT 23, 2001 OCT NEWA
 >A> AP 2MG/ML N75695 002 OCT 23, 2001 OCT NEWA
 SPRAY, METERED; NASAL
 AB MYLAN 1MG/SPRAY N75759 001 AUG 08, 2001 AUG NEWA
 STADOL
 AB + BRISTOL MYERS SQUIBB 1MG/SPRAY N19890 001 DEC 12, 1991 AUG CFTG

CALCITONIN, SALMON

INJECTABLE; INJECTION
 CALCITONIN-SALMON
 @ ASTRAZENECA 200 IU/ML N73690 001 APR 14, 1995 JUN DISC

CALCITRIOL

CAPSULE; ORAL
 >A> CALCITRIOL
 >A> AB TEVA 0.25UGM N75765 001 OCT 12, 2001 OCT NEWA
 >A> AB 0.5UGM N75765 002 OCT 12, 2001 OCT NEWA
 ROCALTROL
 >D> ROCHE 0.25UGM N18044 001 AUG 17, 1978 OCT CFTG
 >A> AB 0.25UGM N18044 001 AUG 17, 1978 OCT CFTG
 >D> + 0.5UGM N18044 002 AUG 17, 1978 OCT CFTG
 >A> AB + 0.5UGM N18044 002 AUG 17, 1978 OCT CFTG

CALCIUM ACETATE

CAPSULE; ORAL
 PHOSLO
 BRAINTREE EQ 84.5MG CALCIUM N21160 001 APR 02, 2001 APR NEWA
 @ EQ 84.5MG CALCIUM N21160 001 APR 02, 2001 AUG DISC
 + EQ 169MG CALCIUM N21160 002 APR 02, 2001 APR NEWA
 + @ EQ 169MG CALCIUM N21160 002 APR 02, 2001 AUG DISC
 PHOSLO GELCAPS
 + BRAINTREE EQ 169MG CALCIUM N21160 003 APR 02, 2001 AUG NEWA

CAPTOPRIL

TABLET; ORAL
 CAPTOPRIL
 AB GENEVA PHARMS TECH 12.5MG N74481 001 FEB 13, 1996 JAN CAHN
 AB 25MG N74481 002 FEB 13, 1996 JAN CAHN

AB	50MG	N74481 003	FEB 13, 1996	JAN CAHN
AB	100MG	N74481 004	FEB 13, 1996	JAN CAHN

CARBACHOL

SOLUTION; INTRAOCULAR				
CARBASTAT				
AT NOVARTIS	0.01%	N73677 001	APR 28, 1995	FEB CAHN

CARBAMAZEPINE

TABLET, CHEWABLE; ORAL				
CARBAMAZEPINE				
AB CARACO	100MG	N75712 001	JUL 05, 2001	JUL NEWA
TABLET, EXTENDED RELEASE; ORAL				
TEGRETOL-XR				
NOVARTIS	100MG	N20234 001	MAR 25, 1996	JUL CRLD
	200MG	N20234 002	MAR 25, 1996	JUL CRLD

CARBIDOPA; LEVODOPA

TABLET; ORAL				
CARBIDOPA AND LEVODOPA				
@ SCS	10MG;100MG	N74080 001	MAR 25, 1994	FEB WDRP
@	25MG;100MG	N74080 002	MAR 25, 1994	FEB WDRP
@	25MG;250MG	N74080 003	MAR 25, 1994	FEB WDRP

CARISOPRODOL

TABLET; ORAL				
CARISOPRODOL				
AA ABLE	350MG	N40421 001	JUN 21, 2001	JUN NEWA

CASPOFUNGIN ACETATE

INJECTABLE; IV (INFUSION)				
CANCIDAS				
+ MERCK RES	50MG/VIAL	N21227 001	JAN 26, 2001	JAN NEWA
+	70MG/VIAL	N21227 002	JAN 26, 2001	JAN NEWA

CEFACLOR

CAPSULE; ORAL				
CECLR				
AB CEPH INTL	EQ 250MG BASE	N62205 001	JUL 28, 1979	JUN CAHN
AB	EQ 500MG BASE	N62205 002	JUL 28, 1979	JUN CAHN
FOR SUSPENSION; ORAL				
CEFACLOR				
@ ZENITH GOLDLINE	EQ 125MG BASE/5ML	N64087 001	APR 28, 1995	MAY DISC
@	EQ 187MG BASE/5ML	N64086 001	APR 28, 1995	MAY DISC
@	EQ 250MG BASE/5ML	N64085 001	APR 28, 1995	MAY DISC
TABLET, EXTENDED RELEASE; ORAL				
CECLR CD				
LILLY	EQ 375MG BASE	N50673 001	JUN 28, 1996	APR CTEC
AB +	EQ 500MG BASE	N50673 002	JUN 28, 1996	JAN CFTG
CEFACLOR				
AB ZENITH GOLDLINE	EQ 500MG BASE	N65057 001	JAN 05, 2001	JAN NEWA

CEFADROXIL/CEFADROXIL HEMIHYDRATE

TABLET; ORAL

CEFADROXIL

© ZENITH GOLDLINE

EQ 1GM BASE

N62774 001 APR 08, 1987 MAY DISC

CEFAMANDOLE NAFATE

INJECTABLE; INJECTION

MANDOL

© LILLY

©

EQ 1GM BASE/VIAL

EQ 2GM BASE/VIAL

N62560 001 SEP 10, 1985 MAY DISC
N62560 002 SEP 10, 1985 MAY DISCCEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

AP HIKMA

EQ 500MG BASE/VIAL

N65047 001 SEP 18, 2001 SEP NEWA

AP

EQ 1GM BASE/VIAL

N65047 002 SEP 18, 2001 SEP NEWA

© TEVA

EQ 250MG BASE/VIAL

N63016 001 MAR 14, 1989 APR DISC

©

EQ 500MG BASE/VIAL

N63016 002 MAR 14, 1989 APR DISC

© KEFZOL

EQ 1GM BASE/VIAL

N63016 003 MAR 14, 1989 APR DISC

© LILLY

EQ 500MG BASE/VIAL

N62557 001 SEP 10, 1985 MAY DISC

©

EQ 1GM BASE/VIAL

N62557 002 SEP 10, 1985 MAY DISC

CEFDITOREN PIVOXIL

TABLET; ORAL

SPECTRACEF

+ TAP PHARM

200MG

N21222 001 AUG 29, 2001 AUG NEWA

CEFONICID SODIUM

INJECTABLE; INJECTION

MONOCID

+ © SMITHKLINE BEECHAM

EQ 500MG BASE/VIAL

N50579 001 MAY 23, 1984 AUG DISC

+ ©

EQ 1GM BASE/VIAL

N50579 002 MAY 23, 1984 AUG DISC

©

EQ 1GM BASE/VIAL

N63295 001 JUL 26, 1993 APR DISC

©

EQ 10GM BASE/VIAL

N50579 004 MAY 23, 1984 AUG DISC

CEFOPERAZONE SODIUM

INJECTABLE; INJECTION

CEFOBID

© PFIZER

©

EQ 1GM BASE/VIAL

N63333 001 MAR 31, 1995 MAY DISC

EQ 2GM BASE/VIAL

N63333 002 MAR 31, 1995 MAY DISC

CEFORANIDE

INJECTABLE; INJECTION

PRECEF

© APOTHECON

500MG/VIAL

N62579 001 NOV 26, 1984 MAY DISC

©

1GM/VIAL

N62579 002 NOV 26, 1984 MAY DISC

©

2GM/VIAL

N62579 003 NOV 26, 1984 MAY DISC

©

10GM/VIAL

N62579 004 NOV 26, 1984 MAY DISC

©

20GM/VIAL

N62579 005 NOV 26, 1984 MAY DISC

CEFOXITIN SODIUM

INJECTABLE; INJECTION

MEFOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

© MERCK	EQ 20MG BASE/ML	N50581 003	SEP 20, 1984	JUL	DISC
©	EQ 40MG BASE/ML	N50581 004	SEP 20, 1984	JUL	DISC

CEFTAZIDIME

INJECTABLE; INJECTION

TAZICEF

AP ABBOTT	500MG/VIAL	N62662 001	MAR 06, 1986	JAN	CAHN
AP	1GM/VIAL	N62662 002	MAR 06, 1986	JAN	CAHN
AP	1GM/VIAL	N64032 001	OCT 31, 1993	JAN	CAHN
AP	2GM/VIAL	N62662 003	MAR 06, 1986	JAN	CAHN
AP	2GM/VIAL	N64032 002	OCT 31, 1993	JAN	CAHN
AP	6GM/VIAL	N62662 004	MAR 06, 1986	JAN	CAHN
TAZIDIME IN PLASTIC CONTAINER					
© LILLY	1GM/VIAL	N62739 001	JUL 10, 1986	MAY	DISC
©	2GM/VIAL	N62739 002	JUL 10, 1986	MAY	DISC

CEFUROXIME SODIUM

INJECTABLE; IM-IV

CEFUROXIME

AB AM PHARM PARTNERS	EQ 750MG BASE/VIAL	N65001 001	MAY 30, 2001	MAY	NEWA
AB TEVA	EQ 750MG BASE/VIAL	N64192 002	APR 16, 1998	MAY	CDFR
CEFUROXIME SODIUM					
AB HANFORD GC	EQ 750MG BASE/VIAL	N64125 001	MAY 30, 1997	MAY	CDFR
KEFUROX					
AB LILLY	EQ 750MG BASE/VIAL	N62591 001	JAN 10, 1986	MAY	CDFR
AB ZINACEF	EQ 750MG BASE/VIAL	N50558 002	OCT 19, 1983	MAY	CDFR
AB + GLAXO WELLCOME	EQ 750MG BASE/VIAL				
INJECTABLE; INJECTION					
CEFUROXIME					
AP AM PHARM PARTNERS	EQ 1.5GM BASE/VIAL	N65001 002	MAY 30, 2001	MAY	NEWA
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER					
+ B BRAUN	EQ 15MG BASE/ML	N50780 001	FEB 21, 2001	FEB	NEWA
+	EQ 30MG BASE/ML	N50780 002	FEB 21, 2001	FEB	NEWA
KEFUROX IN PLASTIC CONTAINER					
© LILLY	EQ 1.5GM BASE/VIAL	N62590 002	JAN 10, 1986	MAY	DISC
INJECTABLE; INTRAVENOUS					
© LILLY	EQ 750MG BASE/VIAL	N62590 001	JAN 10, 1986	MAY	DISC

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

© STEVENS J

EQ 500MG BASE

N62869 001 MAR 17, 1988 JUL DISC

© TEVA

EQ 500MG BASE

N62823 001 FEB 05, 1988 MAY DISC

KEFLEX

AB CEPH INTL

EQ 250MG BASE

N62118 001 MAR 27, 1978 JUN CAHN

AB

EQ 500MG BASE

N62118 002 MAR 27, 1978 JUN CAHN

FOR SUSPENSION; ORAL

CEPHALEXIN

© BARR

EQ 125MG BASE/5ML

N62778 001 AUG 06, 1987 MAY DISC

AB	RANBAXY	EQ 125MG BASE/5ML	N65081 001	JUL 27, 2001	JUL	NEWA
AB		EQ 250MG BASE/5ML	N65081 002	JUL 27, 2001	JUL	NEWA
	KEFLEX					
+ CEPH INTL		EQ 100MG BASE/ML	N62117 001	MAR 27, 1978	JUN	CAHN
AB		EQ 125MG BASE/5ML	N62117 002	MAR 27, 1978	JUN	CAHN
AB +		EQ 250MG BASE/5ML	N62117 003	MAR 27, 1978	JUN	CAHN
	TABLET; ORAL					
	KEFLET					
@ LILLY		EQ 250MG BASE	N62745 001	DEC 01, 1986	JUL	DISC
@		EQ 500MG BASE	N62745 002	DEC 01, 1986	JUL	DISC

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION						
KEFLIN IN PLASTIC CONTAINER						
@ LILLY	EQ 1GM BASE/VIAL	N62549 001	SEP 10, 1985	APR	DISC	
@	EQ 2GM BASE/VIAL	N62549 002	SEP 10, 1985	APR	DISC	

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL						
ZYRTEC-D 12 HOUR						
+ PFIZER	5MG;120MG	N21150 001	AUG 10, 2001	AUG	NEWA	

CHLORAMPHENICOL

CAPSULE; ORAL						
CHLORAMPHENICOL						
@ ZENITH GOLDLINE	250MG	N62247 001	APR 28, 1980	MAY	DISC	
CHLOROMYCETIN						
@ PARKADEALE	50MG	N60591 001	DEC 08, 1950	MAY	DISC	
@	100MG	N60591 003	DEC 08, 1950	MAY	DISC	
@	250MG	N60591 002	DEC 08, 1950	MAY	DISC	
MYCHEL						
+ ARMENPHARM	250MG	N60851 001	JUN 20, 1967	MAY	CRLD	
SOLUTION/DROPS; OPHTHALMIC						
CHLORAMPHENICOL						
@ AKORN	0.5%	N62042 001	AUG 31, 1981	FEB	WDRP	
@ ALCON	0.5%	N62628 001	SEP 25, 1985	MAY	DISC	
CHLOROPTIC						
+ ALLERGAN	0.5%	N50091 001	MAR 20, 1968	MAY	CTEC	

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL						
CHLORDIAZACHEL						
@ RACHELLE	5MG	N85086 001	MAY 11, 1976	FEB	WDRP	
@	10MG	N84639 001	MAY 11, 1976	FEB	WDRP	
@	25MG	N85087 001	MAY 11, 1976	FEB	WDRP	
CHLORDIAZEPOXIDE HCL						
@ FERRANTE	5MG	N85118 001	SEP 02, 1981	FEB	WDRP	
@	10MG	N85119 001	SEP 02, 1976	FEB	WDRP	
@	25MG	N85120 001	SEP 02, 1976	FEB	WDRP	
@ GENEVA PHARMS	5MG	N84678 001	JUN 15, 1976	JUL	DISC	
@	10MG	N84041 001	JUN 15, 1976	MAY	DISC	
@	25MG	N84679 002	SEP 07, 1976	MAY	DISC	
@ IMPAX LABS	5MG	N86213 001	JUL 10, 1979	JUL	DISC	

@	25MG	N86212 001	JUL 10, 1979	JUL	DISC
@ ROSEMONT	5MG	N84644 001	FEB 24, 1976	MAY	DISC
@	25MG	N84645 001	FEB 24, 1976	JUL	DISC

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE

@ TEVA EQ 150MG BASE

N87504 001 JAN 13, 1982 JUL DISC

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

@ ABC HOLDING	250MG	N85569 001	MAR 08, 1978	MAY	DISC
@ CHELSEA LABS	250MG	N86795 001	AUG 15, 1983	JUL	DISC
@ DANBURY PHARMA	250MG	N85173 001	NOV 04, 1977	MAY	DISC

CHLORPHENIRAMINE MALEATE

INJECTABLE; INJECTION

CHLORPHENIRAMINE MALEATE

@ STERIS	10MG/ML	N86096 001	OCT 09, 1979	JUL	DISC
TABLET; ORAL					
@ GENEVA PHARMS	4MG	N80961 001	DEC 20, 1972	MAY	DISC
AA + ICN	4MG	N80598 001	FEB 11, 1972	MAY	CRLD
@ PHARMAVITE	4MG	N85104 001	FEB 11, 1977	FEB	WDRP
@ WEST WARD	4MG	N83787 001	OCT 18, 1973	FEB	WDRP

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CHLORPHENIRAMINE MALEATE AND PHENYLPROPANOLAMINE HCL

@ GENEVA PHARMS	12MG;75MG	N88940 001	JAN 26, 1989	SEP	DISC
DRIZE					
@ ASCHER	12MG;75MG	N88359 001	FEB 13, 1986	SEP	DISC
ORNADE					
@ SMITHKLINE BEECHAM	12MG;75MG	N12152 004	JAN 06, 1981	SEP	DISC

CHLORPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLORPROMAZINE HCL

@ STERIS 25MG/ML N80365 001 FEB 13, 1974 MAY DISC

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

@ GENEVA PHARMS 25MG N87380 001 MAY 01, 1981 JUL DISC

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

@ DANBURY PHARMA 500MG N81019 001 JUL 29, 1991 MAY DISC

CIMETIDINE

TABLET; ORAL

CIMETIDINE

AB	GENEVA PHARMS TECH	200MG	N74506 001	JAN 24, 1996	JAN	CAHN
AB		300MG	N74506 002	JAN 24, 1996	JAN	CAHN
AB		400MG	N74506 003	JAN 24, 1996	JAN	CAHN
AB		800MG	N74506 004	JAN 24, 1996	JAN	CAHN

CINOXACIN

CAPSULE; ORAL

CINOBAC

LILLY

+		250MG	N18067 001	JUN 13, 1980	JUL	CTEC
	CINOXACIN	500MG	N18067 002	JUN 13, 1980	JUL	CTEC
@	TEVA	250MG	N73005 001	FEB 28, 1992	JUL	DISC
@		500MG	N73006 001	FEB 28, 1992	JUL	DISC

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN HCL

AB	+	PHARMACIA AND UPJOHN	EQ 300MG BASE	N50162 003	APR 14, 1988	FEB	CFTG
		CLINDAMYCIN HCL					
AB		RANBAXY	EQ 150MG BASE	N65061 001	FEB 02, 2001	FEB	NEWA
AB			EQ 300MG BASE	N65061 002	FEB 02, 2001	FEB	NEWA

CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE

BT	GALDERMA LABS LP	EQ 1% BASE	N50782 001	NOV 27, 2000	SEP	CAHN	
	INJECTABLE; INJECTION						
AP	+	PHARMACIA AND UPJOHN	EQ 18MG BASE/ML	N50639 003	APR 10, 1991	JUN	CFTG
	CLINDAMYCIN PHOSPHATE						
@	ABBOTT	EQ 150MG BASE/ML	N62943 001	SEP 29, 1988	MAY	DISC	
@	ELKINS SINK	EQ 150MG BASE/ML	N62806 001	OCT 15, 1987	MAY	DISC	
@		EQ 150MG BASE/ML	N62953 001	APR 21, 1988	MAY	DISC	
@	GENSIA SICOR PHARMS	EQ 150MG BASE/ML	N63041 001	DEC 29, 1989	APR	DISC	
@		EQ 150MG BASE/ML	N63282 001	MAY 29, 1992	APR	DISC	
@	LEDERLE	EQ 150MG BASE/ML	N63068 001	AUG 28, 1989	MAY	DISC	
AP	CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER						
AP	ABBOTT	EQ 6MG BASE/ML	N65027 001	JUN 29, 2001	JUN	NEWA	
AP		EQ 12MG BASE/ML	N65027 002	JUN 29, 2001	JUN	NEWA	
AP		EQ 18MG BASE/ML	N65027 003	JUN 29, 2001	JUN	NEWA	
	SOLUTION; TOPICAL						
	CLINDAMYCIN PHOSPHATE						
@	COPLEY PHARM	EQ 1% BASE	N62944 001	JAN 11, 1989	MAY	DISC	
@	TEVA	EQ 1% BASE	N62930 001	JUN 28, 1989	MAY	DISC	

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE

AB1	STIEFEL	0.05%
-----	---------	-------

N75338 001 FEB 09, 2001 FEB NEWA

AB2	0.05%	N75733 001 AUG 22, 2001 AUG NEWA
-----	-------	----------------------------------

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

ANAFRANIL

AB	TYCO HLTHCARE	25MG	N19906 001 DEC 29, 1989 JUN CAHN
AB	+	50MG	N19906 002 DEC 29, 1989 JUN CAHN
AB		75MG	N19906 003 DEC 29, 1989 JUN CAHN

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

AB	CARACO	0.5MG	N75423 001 APR 27, 2001 APR NEWA
AB		1MG	N75423 002 APR 27, 2001 APR NEWA
AB		2MG	N75423 003 APR 27, 2001 APR NEWA

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURACLON

+ ELAN PHARMS	0.1MG/ML	N20615 001 OCT 02, 1996 SEP CAHN
---------------	----------	----------------------------------

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM

@ ABLE	3.75MG	N71777 001 JUL 14, 1987 JAN DISC
@	7.5MG	N71778 001 JUL 14, 1987 JAN DISC
@	15MG	N71779 001 JUL 14, 1987 JAN DISC
TABLET; ORAL		
@ GENEVA PHARMS	3.75MG	N72512 001 MAY 11, 1990 JUL DISC

COLCHICINE; PROBENECID

TABLET; ORAL

PROBENECID AND COLCHICINE

>D> BP	IMPAK LABS	0.5MG;500MG	N83720 002 SEP 06, 1977 OCT DISC
>A>	@	0.5MG;500MG	N83720 002 SEP 06, 1977 OCT DISC

CORTICOTROPIN

INJECTABLE; INJECTION

H.P. ACTHAR GEL

BC	+	QUESTCOR PHARMS	40 UNITS/ML	N08372 006 FEB 06, 1956 AUG CAHN
BC	+		80 UNITS/ML	N08372 008 FEB 06, 1956 AUG CAHN

CORTISONE ACETATE

TABLET; ORAL

CORTISONE ACETATE

@ CHELSEA LABS	25MG	N85884 001 MAY 15, 1978 MAY DISC
----------------	------	----------------------------------

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION

INTAL

+ AVENTIS	0.8MG/INH	N18887 001 DEC 05, 1985 SEP CAHN
SOLUTION/DROPS; OPHTHALMIC		
CROMOLYN SODIUM		

<u>CROMOLYN SODIUM</u>			
SOLUTION/DROPS; OPHTHALMIC			
CROMOLYN SODIUM			
AT	NOVEX	4%	N75615 001 JAN 26, 2001 JAN NEWA
<u>CYCLACILLIN</u>			
TABLET; ORAL			
CYCLACILLIN			
② TEVA		250MG	N62895 001 AUG 04, 1988 MAY DISC
②		500MG	N62895 002 AUG 04, 1988 MAY DISC
<u>CYPHEPTADINE HYDROCHLORIDE</u>			
TABLET; ORAL			
CYPHEPTADINE HCL			
② GENEVA PHARMS		4MG	N86808 001 FEB 24, 1981 JUL DISC
<u>DACARBAZINE</u>			
INJECTABLE; INJECTION			
DACARBAZINE			
AP	BEDFORD	200MG/VIAL	
>A>	AP FAULDING	200MG/VIAL	N75812 001 JUN 15, 2001 JUN NEWA N75940 001 OCT 18, 2001 OCT NEWA
<u>DELAVIRDINE MESYLATE</u>			
TABLET; ORAL			
RESCRIPTOR			
AGOURON		100MG	
+		200MG	N20705 001 APR 04, 1997 AUG CRLD N20705 002 JUL 14, 1999 AUG NEWA
<u>DESERPIDINE; HYDROCHLOROTHIAZIDE</u>			
TABLET; ORAL			
ORETICYL 25			
② ABBOTT		0.125MG;25MG	N12148 001 DEC 14, 1959 MAR DISC
ORETICYL 50			
② ABBOTT		0.125MG;50MG	N12148 003 DEC 14, 1959 MAR DISC
ORETICYL FORTE			
② ABBOTT		0.25MG;25MG	N12148 002 DEC 14, 1959 MAR DISC
<u>DESONIDE</u>			
OINTMENT; TOPICAL			
DESONIDE			
AB	ALTANA	0.05%	N75751 001 MAR 12, 2001 MAR NEWA
<u>DEXAMETHASONE</u>			
TABLET; ORAL			
DEXAMETHASONE			
② DANBURY PHARMA		0.75MG	N80968 001 MAY 03, 1973 MAY DISC
<u>DEXAMETHASONE SODIUM PHOSPHATE</u>			
INJECTABLE; INJECTION			
DEXAMETHASONE SODIUM PHOSPHATE			
② DELL LABS		EQ 4MG PHOSPHATE/ML	N83161 001 JUN 06, 1978 FEB WDRP
② GENESIA SICOR PHARMS		EQ 4MG PHOSPHATE/ML	N81125 001 AUG 31, 1990 MAY DISC
OINTMENT; OPHTHALMIC			

DEXAMETHASONE SODIUM PHOSPHATE

OINTMENT; OPHTHALMIC

DECADRON

@ MERCK

EQ 0.05% PHOSPHATE

N11977 001 SEP 02, 1959 MAY DISC

MAXIDEX

+ ALCON

EQ 0.05% PHOSPHATE

N83342 001 OCT 23, 1973 MAY CTEC

SOLUTION/DROPS; OTIC

DEXAMETHASONE SODIUM PHOSPHATE

@ AKORN

EQ 0.1% PHOSPHATE

N84855 001 JUN 29, 1976 FEB WDRP

DEXAMETHASONE SODIUM PHOSPHATE; NEOMYCIN SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEODECADRON

+ MERCK

EQ 0.1% PHOSPHATE; EQ 3.5MG

BASE/ML

N50322 001 JUL 06, 1959 MAY CTEC

NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE

@ ALCON UNIVERSAL

EQ 0.1% PHOSPHATE; EQ 3.5MG

BASE/ML

N62714 001 JUL 21, 1986 MAY DISC

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

DEXACIDIN

AT NOVARTIS

0.1%;EQ 3.5MG

BASE/GM;10,000 UNITS/GM

N62566 001 FEB 22, 1985 FEB CAHN

@

0.1%;EQ 3.5MG

BASE/GM;10,000 UNITS/GM

N62566 001 FEB 22, 1985 MAY DISC

SUSPENSION/DROPS; OPHTHALMIC

AT NOVARTIS

0.1%;EQ 3.5MG

BASE/ML;10,000 UNITS/ML

N62544 001 OCT 29, 1984 FEB CAHN

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

AA BARR

5MG

N40361 001 JAN 31, 2001 JAN NEWA

AA

10MG

N40361 002 JAN 31, 2001 JAN NEWA

DEXTROSTAT

AA + SHIRE RICHWOOD

10MG

N84051 002 MAY 29, 1975 JAN CFTG

DIAZEPAM

GEL; RECTAL

DIASTAT

+ XCEL PHARMS

2.5MG/0.5ML

N20648 001 JUL 29, 1997 JUL CAHN

5MG/ML

N20648 002 JUL 29, 1997 JUL CAHN

10MG/2ML

N20648 003 JUL 29, 1997 JUL CAHN

15MG/3ML

N20648 004 JUL 29, 1997 JUL CAHN

+

20MG/4ML

N20648 005 JUL 29, 1997 JUL CAHN

DICLOFENAC POTASSIUM

TABLET; ORAL

DICLOFENAC POTASSIUM

AB EON

50MG

N75582 001 FEB 23, 2001 FEB NEWA

<u>DICLOFENAC SODIUM</u>			
GEL; TOPICAL			
SOLARAZE			
+ BIOLAN PHARMA PLC	3%		N21005 001 OCT 16, 2000 MAR CAHN
<u>DICLOXACILLIN SODIUM</u>			
CAPSULE; ORAL			
DYCILL			
@ SMITHKLINE BEECHAM	EQ 250MG BASE		N62238 001 DEC 31, 1979 APR DISC
@	EQ 500MG BASE		N62238 002 DEC 31, 1979 APR DISC
<u>DICYCLOMINE HYDROCHLORIDE</u>			
CAPSULE; ORAL			
DICYCLOMINE HCL			
@ HALSEY	10MG		N84505 001 OCT 21, 1986 MAY DISC
INJECTABLE; INJECTION			
@ STERIS	10MG/ML		N80614 001 FEB 11, 1986 JUL DISC
<u>DIETHYLPROPION HYDROCHLORIDE</u>			
TABLET, EXTENDED RELEASE; ORAL			
TENUATE DOSPAN			
+ AVENTIS PHARMS	75MG		N12546 001 NOV 07, 1960 JUL CTEC
TEPANIL TEN-TAB			
@ 3M	75MG		N17956 001 MAY 25, 1977 JUL DISC
<u>DILTIAZEM HYDROCHLORIDE</u>			
CAPSULE, EXTENDED RELEASE; ORAL			
DILTIAZEM HCL			
AB2 MYLAN	120MG		N75124 002 MAR 18, 1998 MAR CTEC
<u>DIPHENHYDRAMINE HYDROCHLORIDE</u>			
CAPSULE; ORAL			
DIPHENHYDRAMINE HCL			
@ CHELSEA LABS	50MG		N85083 001 JUN 29, 1976 MAY DISC
@ NEWTRON PHARMS	25MG		N86543 001 FEB 08, 1979 FEB WDRP
@	50MG		N86544 001 FEB 08, 1979 FEB WDRP
INJECTABLE; INJECTION			
DIPHENHYDRAMINE HCL PRESERVATIVE FREE			
@ AM PHARM PARTNERS	50MG/ML		N80586 002 JAN 10, 1973 JUL DISC
<u>DISULFIRAM</u>			
TABLET; ORAL			
ANTABUSE			
ODYSSEY PHARMS	250MG		N88482 001 DEC 08, 1983 JAN CAHN
+	500MG		N88483 001 DEC 08, 1983 JAN CAHN
@ SIDMAK LABS	250MG		N07883 003 NOV 03, 1970 MAR CAHN
@	500MG		N07883 002 JUN 01, 1953 MAR CAHN
<u>DOXAZOSIN MESYLATE</u>			
TABLET; ORAL			
DOXAZOSIN MESYLATE			
AB SIDMAK LABS	EQ 1MG BASE		N75750 001 JUN 08, 2001 JUN NEWA

AB		EQ 2MG BASE	N75750 002	JUN 08, 2001	JUN	NEWA
AB		EQ 4MG BASE	N75750 003	JUN 08, 2001	JUN	NEWA
AB		EQ 8MG BASE	N75750 004	JUN 08, 2001	JUN	NEWA
AB	TEVA	EQ 1MG BASE	N75353 001	JAN 12, 2001	JAN	NEWA
AB		EQ 2MG BASE	N75353 002	JAN 12, 2001	JAN	NEWA
AB		EQ 4MG BASE	N75353 003	JAN 12, 2001	JAN	NEWA
AB		EQ 8MG BASE	N75353 004	JAN 12, 2001	JAN	NEWA

DOXYCYCLINE

FOR SUSPENSION; ORAL

DOXYCHEL

@ RACHELLE

EQ 25MG BASE/5ML

N61720 001 JUN 18, 1973 FEB WDRP

VIBRAMYCIN

+ PFIZER

EQ 25MG BASE/5ML

N50006 001 DEC 06, 1967 FEB CTEC

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXY-LEMMON

@ TEVA

EQ 50MG BASE

N62497 001 AUG 23, 1984 APR DISC

@

EQ 100MG BASE

N62497 002 JUN 15, 1984 APR DISC

DOXYCYCLINE HYCLATE

@ CHELSEA LABS

EQ 50MG BASE

N62142 001 AUG 12, 1981 APR DISC

@

EQ 100MG BASE

N62142 002 AUG 12, 1981 APR DISC

AB HALSEY

EQ 50MG BASE

N61717 001 JUL 17, 1973 JUN CAHN

@

EQ 50MG BASE

N62418 001 JAN 28, 1983 APR DISC

AB

EQ 100MG BASE

N61717 002 JUL 17, 1973 JUN CAHN

@

EQ 100MG BASE

N62418 002 JAN 28, 1983 APR DISC

CAPSULE, COATED PELLETS; ORAL

@ SIDMAK LABS NJ

EQ 100MG BASE

N63187 001 JUN 30, 1992 MAY DISC

INJECTABLE; INJECTION

DOXYCHEL HYCLATE

@ RACHELLE

EQ 100MG BASE/VIAL

N61953 001 SEP 10, 1980 FEB WDRP

DOXYCYCLINE

@ BEDFORD

EQ 100MG BASE/VIAL

N62569 001 MAR 09, 1988 MAY DISC

@

EQ 200MG BASE/VIAL

N62569 002 MAR 09, 1988 MAY DISC

@ ELKINS SINK

EQ 100MG BASE/VIAL

N62450 001 OCT 27, 1983 APR DISC

@

EQ 200MG BASE/VIAL

N62450 002 OCT 27, 1983 APR DISC

DOXYCYCLINE HYCLATE

@ LEDERLE

EQ 100MG BASE/VIAL

N62992 001 FEB 16, 1989 MAY DISC

@

EQ 200MG BASE/VIAL

N62992 002 FEB 16, 1989 MAY DISC

TABLET; ORAL

DOXY-LEMMON

@ TEVA

EQ 100MG BASE

N62581 001 MAR 15, 1985 MAY DISC

DOXYCYCLINE HYCLATE

AB HALSEY

EQ 100MG BASE

N62269 001 SEP 03, 1980 JUN CAHN

AB

EQ 100MG BASE

N62269 002 NOV 08, 1982 JUN CAHN

DOXYCYCLINE HYCLATE

@ HALSEY

EQ 50MG BASE

N62269 003 SEP 03, 1980 JUN CAHN

PERIOSTAT

+ COLLAGENEX PHARMS

20MG

N50783 001 FEB 02, 2001 FEB NEWA

DROPERIDOL; FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE AND DROPERIDOL

© ASTRazeneca

2.5MG/ML;EQ 0.05MG BASE/ML

N72027 001 APR 13, 1989 JUL DISC

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL-28

YASMIN

+ BERLEX LABS

3MG;0.03MG

N21098 001 MAY 11, 2001 MAY NEWA

DYCLONINE HYDROCHLORIDE

SOLUTION; TOPICAL

DYCLONE

© ASTRazeneca

0.5%

N09925 002 JUN 13, 1974 AUG DISC

©

1%

N09925 001 AUG 03, 1955 AUG DISC

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

AB	TARO	2.5MG	
AB		5MG	N75657 002 JAN 23, 2001 JAN NEWA
AB		10MG	N75657 003 JAN 23, 2001 JAN NEWA
AB		20MG	N75657 004 JAN 23, 2001 JAN NEWA
AB	TORPHARM	2.5MG	N75178 002 MAR 23, 2001 MAR NEWA
AB		5MG	N75178 001 MAR 23, 2001 MAR NEWA
AB		10MG	N75178 003 MAR 23, 2001 MAR NEWA
AB		20MG	N75178 004 MAR 23, 2001 MAR NEWA

N75657 001 JAN 23, 2001 JAN NEWA

N75657 002 JAN 23, 2001 JAN NEWA

N75657 003 JAN 23, 2001 JAN NEWA

N75657 004 JAN 23, 2001 JAN NEWA

N75178 002 MAR 23, 2001 MAR NEWA

N75178 001 MAR 23, 2001 MAR NEWA

N75178 003 MAR 23, 2001 MAR NEWA

N75178 004 MAR 23, 2001 MAR NEWA

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

>A>	AB	DR REDDYS LABS LTD	5MG;12.5MG	N75909 001 OCT 15, 2001 OCT NEWA
>A>	AB		10MG;25MG	N75909 002 OCT 15, 2001 OCT NEWA
	AB	EON	5MG;12.5MG	N76116 001 SEP 19, 2001 SEP NEWA
	AB		10MG;25MG	N76116 002 SEP 19, 2001 SEP NEWA
	AB	MYLAN	5MG;12.5MG	N75624 001 SEP 18, 2001 SEP NEWA
	AB		10MG;25MG	N75624 002 SEP 18, 2001 SEP NEWA
	AB	TARO PHARM INDs	5MG;12.5MG	N75788 001 SEP 18, 2001 SEP NEWA
	AB		10MG;25MG	N75788 002 SEP 18, 2001 SEP NEWA
	AB	TEVA	5MG;12.5MG	N75727 001 SEP 18, 2001 SEP NEWA
	AB		10MG;25MG	N75727 002 SEP 18, 2001 SEP NEWA
		VASERETIC		
	AB	MERCK RES LABS	5MG;12.5MG	N19221 003 JUL 12, 1995 SEP CFTG
AB	+		10MG;25MG	N19221 001 OCT 31, 1986 SEP CFTG

N75909 002 OCT 15, 2001 OCT NEWA

N76116 001 SEP 19, 2001 SEP NEWA

N76116 002 SEP 19, 2001 SEP NEWA

N75624 001 SEP 18, 2001 SEP NEWA

N75624 002 SEP 18, 2001 SEP NEWA

N75788 001 SEP 18, 2001 SEP NEWA

N75788 002 SEP 18, 2001 SEP NEWA

N75727 001 SEP 18, 2001 SEP NEWA

N75727 002 SEP 18, 2001 SEP NEWA

ENFLURANE

LIQUID; INHALATION

ENFLURANE

AN MINRAD

99.9%

N74396 001 JUL 29, 1994 FEB CAHN

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

LOVENOX

+ AVENTIS	30MG/0.3ML	N20164 001	MAR 29, 1993	APR	CAHN
+	40MG/0.4ML	N20164 002	JAN 30, 1998	APR	CAHN
+	60MG/0.6ML	N20164 003	MAR 27, 1998	APR	CAHN
+	80MG/0.8ML	N20164 004	MAR 27, 1998	APR	CAHN
+	90MG/0.6ML	N20164 006	JUN 02, 2000	APR	CAHN
+	100MG/ML	N20164 005	MAR 27, 1998	APR	CAHN
+	120MG/0.8ML	N20164 007	JUN 02, 2000	APR	CAHN
+	150MG/ML	N20164 008	JUN 02, 2000	APR	CAHN

EPINEPHRINE BITARTRATE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

@ ASTRAZENECA	0.005MG/ML;1.5%	N17751 007	AUG 30, 1976	SEP	DISC
+ DENTSPLY PHARM	0.005MG/ML;1%	N17751 006	AUG 30, 1976	APR	CAHN
+	0.005MG/ML;1.5%	N17751 007	AUG 30, 1976	APR	CAHN
+	0.005MG/ML;1.5%	N21384 001	AUG 30, 1976	SEP	NEWA

EPINEPHRINE BITARTRATE; PRILOCaine HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE

@ ASTRAZENECA	0.005MG/ML;4%	N14763 008	JUN 29, 1970	SEP	DISC
+ DENTSPLY PHARM	0.005MG/ML;4%	N21383 001	JUN 29, 1970	SEP	NEWA

EPINEPHRINE; ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

@ DENTSPLY PHARM

0.005MG/ML;0.5%

N17751 004 AUG 30, 1976 APR CAHN

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HCL W/ EPINEPHRINE

@ INTL MEDICATION	0.01MG/ML;1%	N86402 001	FEB 04, 1980	JUL	DISC
@ STERIS	0.01MG/ML;1%	N80377 003	FEB 20, 1974	JUL	DISC
@	0.01MG/ML;2%	N80377 004	FEB 20, 1974	JUL	DISC
LIDOCATON					
@ PHARMATON	0.02MG/ML;2%	N84728 001	AUG 17, 1983	FEB	WDRP
XYLOCAINE W/ EPINEPHRINE					
@ ASTRAZENECA	0.01MG/ML;2%	N06488 003	NOV 19, 1948	SEP	DISC
+ DENTSPLY PHARM	0.01MG/ML;2%	N21381 001	NOV 19, 1948	SEP	NEWA
	0.02MG/ML;2%	N21381 002	NOV 19, 1948	SEP	NEWA

ERGOCALCIFEROL

CAPSULE; ORAL

VITAMIN D

@ IMPAX LABS

50,000 IU

N80951 001 JUL 13, 1973 FEB DISC

ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATES

@ DANBURY PHARMA	1MG	N87244 001 AUG 16, 1982 JUL DISC
TABLET; SUBLINGUAL		
@ DANBURY PHARMA	1MG	N87183 001 APR 16, 1981 JUL DISC

ERYTHROMYCIN

SOLUTION; TOPICAL

ERYTHROMYCIN

AT @ CLAY PARK	2%	N63038 001 JAN 11, 1991 APR DISC
	2%	N63038 001 JAN 11, 1991 MAY CMFD
TABLET, DELAYED RELEASE; ORAL		
E-BASE		
@ BARR	333MG	N63028 001 MAY 15, 1990 APR DISC
ILOTYCIN		
@ DISTA	250MG	N61910 001 FEB 27, 1975 MAY DISC

ERYTHROMYCIN ESTOLATE

CAPSULE; ORAL

ERYTHROMYCIN ESTOLATE

+ BARR	EQ 250MG BASE	N62162 002 JUN 15, 1981 MAY CTEC
@ DANBURY PHARMA	EQ 250MG BASE	N62087 001 JUN 14, 1979 APR DISC
ILOSONE		
@ LILLY	EQ 125MG BASE	N61897 001 JAN 06, 1975 MAY DISC
@	EQ 250MG BASE	N61897 002 JAN 06, 1975 MAY DISC
FOR SUSPENSION; ORAL		
@ DISTA	EQ 125MG BASE/5ML	N61893 001 JAN 06, 1975 MAY DISC
SUSPENSION/DROPS; ORAL		
@ LILLY	EQ 100MG BASE/ML	N61894 003 JAN 07, 1975 APR DISC
TABLET; ORAL		
@ LILLY	EQ 500MG BASE	N61896 001 JAN 03, 1975 APR DISC
TABLET, CHEWABLE; ORAL		
@ DISTA	EQ 125MG BASE	N61895 001 JAN 03, 1975 MAY DISC
@	EQ 250MG BASE	N61895 002 JAN 03, 1975 MAY DISC

ERYTHROMYCIN ETHYLSUCCINATE

TABLET; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

@ BARR	EQ 400MG BASE	N62256 001 APR 28, 1980 MAY DISC
--------	---------------	----------------------------------

ERYTHROMYCIN GLUCEPTATE

INJECTABLE; INJECTION

ILOTYCIN GLUCEPTATE

@ DISTA	EQ 250MG BASE/VIAL	N50370 001 JUN 23, 1964 JUL DISC
@	EQ 500MG BASE/VIAL	N50370 002 JUN 23, 1964 JUL DISC
@	EQ 1GM BASE/VIAL	N50370 003 JUN 23, 1964 JUL DISC

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROMYCIN STEARATE

@ BARR	EQ 500MG BASE	N63179 001 MAY 15, 1990 MAY DISC
--------	---------------	----------------------------------

© ZENITH GOLDLINE	EQ 250MG BASE	N61461 001	SEP 04, 1971	MAY	DISC
©	EQ 500MG BASE	N61461 002	APR 11, 1980	MAY	DISC
WYAMYCIN S					
© WYETH AYERST	EQ 250MG BASE	N61675 001	OCT 06, 1972	APR	DISC
©	EQ 500MG BASE	N61675 002	JUL 13, 1973	APR	DISC

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL					
NEXIUM					
+ ASTRAZENECA	EQ 20MG BASE	N21153 001	FEB 20, 2001	FEB	NEWA
+	EQ 40MG BASE	N21153 002	FEB 20, 2001	FEB	NEWA

ESTRADIOL VALERATE

INJECTABLE; INJECTION					
DELESTROGEN					
+ KING PHARMS	10MG/ML	N09402 002	AUG 18, 1962	AUG	CAHN
AO +	20MG/ML	N09402 004	JUN 23, 1961	AUG	CAHN
AO +	40MG/ML	N09402 003	FEB 24, 1961	AUG	CAHN

ESTRADIOL; NORETHINDRONE ACETATE

FILM, EXTENDED RELEASE; TRANSDERMAL					
COMBIPATCH					
NOVARTIS	0.05MG/24HR;0.14MG/24HR	N20870 001	AUG 07, 1998	MAR	CAHN
+	0.05MG/24HR;0.25MG/24HR	N20870 002	AUG 07, 1998	MAR	CAHN

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28					
PREMPRO					
+ WYETH AYERST	0.625MG;0.625MG;2.5MG;2.5M G	N20527 001	NOV 17, 1995	JAN	CTNA
+	0.625MG;0.625MG;5MG;5MG	N20527 003	JAN 09, 1998	JAN	CTNA
PREMPRO (PREMARIN;CYCRIN)					
+ WYETH AYERST	0.625MG;0.625MG;2.5MG;2.5M G	N20303 001	DEC 30, 1994	JAN	CTNA

ESTROGENS, ESTERIFIED

TABLET; ORAL					
ESTRATAB					
© SOLVAY	0.3MG	N86715 001	APR 08, 1981	JUL	DISC
©	0.625MG	N83209 001	JUN 17, 1977	JUL	DISC
MENEST					
+ MONARCH PHARMS	0.3MG 0.625MG	N84951 001	SEP 28, 1977	JUL	CTEC
		N84948 001	SEP 28, 1977	JUL	CTEC

ESTROPIPATE

TABLET; ORAL					
ORTHO-EST					
AB WOMEN FIRST HLTHCARE	0.75MG	N89567 001	FEB 27, 1991	JAN	CAHN
AB	1.5MG	N89582 001	JUL 17, 1991	JAN	CAHN

>A> ETHINYL ESTRADIOL; ETONOGESTREL
 >A> RING; VAGINAL
 >A> NUVARING
 >A> + ORGANON INC 0.015MG;0.12MG N21187 001 OCT 03, 2001 OCT NEWA

ETHINYL ESTRADIOL; LEVONORGESTREL
 TABLET; ORAL-21
 ALESSE
 AB + WYETH AYERST 0.02MG;0.1MG N20683 001 MAR 27, 1997 APR CTEC
 AVIANE-21
 AB DURAMED 0.02MG;0.1MG N75796 002 APR 30, 2001 APR NEWA
 ENPRESSE-21
 AB DURAMED 0.03MG,0.04MG;0.03MG;0.05MG N75809 001 JUL 16, 2001 JUL NEWA
 0.075MG;0.125MG
 TABLET; ORAL-28
 ALESSE
 AB WYETH AYERST 0.02MG;0.1MG N20683 002 MAR 27, 1997 APR CTEC
 AVIANE-28
 AB DURAMED 0.02MG;0.1MG N75796 001 APR 30, 2001 APR NEWA
 ENPRESSE-28
 AB DURAMED 0.03MG,0.04MG;0.03MG;0.05MG N75809 002 JUL 16, 2001 JUL NEWA
 0.075MG;0.125MG

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE
 TABLET; ORAL-28
 LOESTRIN FE 1.5/30
 AB + PARKE DAVIS 0.03MG;1.5MG N17355 001 APR 30, 1973 FEB CFTG
 LOESTRIN FE 1/20
 AB + PARKE DAVIS 0.02MG;1MG N17354 001 APR 30, 1973 FEB CFTG
 MICROGESTIN FE 1.5/30
 AB WATSON LABS 0.03MG;1.5MG N75548 001 FEB 05, 2001 FEB NEWA
 MICROGESTIN FE 1/20
 AB WATSON LABS 0.02MG;1MG N75647 001 FEB 05, 2001 FEB NEWA

ETHOSUXIMIDE
 SYRUP; ORAL
 ZARONTIN
 AA + PARKE DAVIS 250MG/5ML N80258 001 FEB 13, 1974 JAN CRLD

ETIDOCAINE HYDROCHLORIDE
 INJECTABLE; INJECTION
 DURANEST
 @ DENTSPLY PHARM 0.5% N17751 003 AUG 30, 1976 APR CAHN
 + 1% N17751 005 AUG 30, 1976 APR CAHN

ETODOLAC
 TABLET, EXTENDED RELEASE; ORAL
 ETODOLAC
 AB TEVA 400MG N75665 003 FEB 05, 2001 FEB NEWA

ETOPOSIDE

CAPSULE; ORAL ETOPOSIDE					
AB GENPHARM	50MG	N75635	001	SEP 19, 2001	SEP NEWA
VEPESID					
AB + BRISTOL INJECTABLE; INJECTION ETOPOSIDE	50MG	N19557	001	DEC 30, 1986	SEP CFTG
@ PIERRE FABRE	20MG/ML	N74813	001	JUL 09, 1997	SEP WDAG

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION ETOPOPHOS PRESERVATIVE FREE					
AB BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N20906	001	FEB 27, 1998	JUN DISC
@	EQ 1GM BASE/VIAL	N20906	002	FEB 27, 1998	JUN NEWA

FAMCICLOVIR

TABLET; ORAL FAMVIR					
NOVARTIS	125MG	N20363	003	DEC 11, 1995	JAN CAHN
	250MG	N20363	001	APR 26, 1996	JAN CAHN
+	500MG	N20363	002	JUN 29, 1994	JAN CAHN

FAMOTIDINE

INJECTABLE; INJECTION FAMOTIDINE					
AP AM PHARM PARTNERS	10MG/ML	N75709	001	APR 16, 2001	APR NEWA
AP APOTHECON	10MG/ML	N75707	001	APR 16, 2001	APR NEWA
@	10MG/ML	N75707	001	APR 16, 2001	MAY DISC
AP BEDFORD	10MG/ML	N75651	001	APR 16, 2001	APR NEWA
AP	10MG/ML	N75684	001	APR 16, 2001	APR NEWA
AP ESI LEDERLE	10MG/ML	N75488	001	APR 16, 2001	APR NEWA
AP FAULDING	10MG/ML	N75705	001	APR 16, 2001	APR NEWA
FAMOTIDINE PRESERVATIVE FREE					
AP AM PHARM PARTNERS	10MG/ML	N75813	001	APR 16, 2001	APR NEWA
AP APOTHECON	10MG/ML	N75708	001	APR 16, 2001	APR NEWA
@	10MG/ML	N75708	001	APR 16, 2001	MAY DISC
AP BEDFORD	10MG/ML	N75622	001	APR 16, 2001	APR NEWA
AP BEN VENUE	10MG/ML	N75825	001	APR 17, 2001	APR NEWA
AP ESI LEDERLE	10MG/ML	N75486	001	APR 16, 2001	APR NEWA
AP FAULDING	10MG/ML	N75669	001	APR 16, 2001	APR NEWA
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER					
AP BAXTER HLTHCARE	0.4MG/ML	N75591	001	MAY 10, 2001	MAY NEWA
PEPCID					
AP + MERCK	10MG/ML	N19510	001	NOV 04, 1986	APR CFTG
PEPCID PRESERVATIVE FREE					
AP + MERCK	10MG/ML	N19510	004	NOV 04, 1986	APR CFTG
PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER					
AP + MERCK	0.4MG/ML	N20249	001	FEB 18, 1994	MAY CFTG
TABLET; ORAL FAMOTIDINE					
AB CARLSBAD	20MG	N75805	001	APR 16, 2001	APR NEWA

AB		40MG	N75805 002	APR 16, 2001	APR	NEWA
AB	DANBURY PHARMA	20MG	N75062 002	APR 16, 2001	APR	NEWA
AB		40MG	N75062 001	APR 16, 2001	APR	NEWA
AB	DR REDDYS LABS LTD	20MG	N75718 001	APR 16, 2001	APR	NEWA
AB		40MG	N75718 002	APR 16, 2001	APR	NEWA
AB	EON	20MG	N75793 001	APR 16, 2001	APR	NEWA
AB		40MG	N75793 002	APR 16, 2001	APR	NEWA
AB	GENEVA PHARMS	20MG	N75302 001	APR 16, 2001	APR	NEWA
AB		40MG	N75302 002	APR 16, 2001	APR	NEWA
AB	GENPHARM	20MG	N75457 001	APR 18, 2001	APR	NEWA
AB		40MG	N75457 002	APR 18, 2001	APR	NEWA
AB	INVAMED	20MG	N75607 001	MAY 10, 2001	MAY	NEWA
AB		40MG	N75607 002	MAY 10, 2001	MAY	NEWA
AB	MYLAN	20MG	N75704 001	APR 16, 2001	APR	NEWA
AB		40MG	N75704 002	APR 16, 2001	APR	NEWA
AB	PUREPAC PHARM	20MG	N75650 001	SEP 14, 2001	SEP	NEWA
AB		40MG	N75650 002	SEP 14, 2001	SEP	NEWA
AB	TEVA	20MG	N75311 001	APR 16, 2001	APR	NEWA
AB		40MG	N75311 002	APR 16, 2001	APR	NEWA
AB	TORPHARM	20MG	N75611 001	JUL 23, 2001	JUL	NEWA
AB		40MG	N75611 002	JUL 23, 2001	JUL	NEWA
AB	WOCKHARDT	20MG	N75786 001	APR 16, 2001	APR	NEWA
AB		40MG	N75786 002	APR 16, 2001	APR	NEWA
AB	ZENITH GOLDLINE	20MG	N75511 001	APR 16, 2001	APR	NEWA
AB		40MG	N75511 002	APR 16, 2001	APR	NEWA
	PEPCID					
AB	MERCK	20MG	N19462 001	OCT 15, 1986	APR	CFTG
AB	+	40MG	N19462 002	OCT 15, 1986	APR	CFTG

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL
PLENDIL

>D>	+	ASTRAZENECA	2.5MG	N19834 004	SEP 22, 1994	OCT	CRLD
>A>			2.5MG	N19834 004	SEP 22, 1994	OCT	CRLD
>D>	+		5MG	N19834 001	JUL 25, 1991	OCT	CRLD
>A>			5MG	N19834 001	JUL 25, 1991	OCT	CRLD

FENOFLIBRATE

TABLET; ORAL
TRICOR

ABBOTT		54MG	N21203 001	SEP 04, 2001	AUG	NEWA
+		160MG	N21203 003	SEP 04, 2001	SEP	NEWA

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL
DURAGESIC

ALZA		1.2MG/24HR	N19813 003	AUG 07, 1990	MAY	CTEC
		1.8MG/24HR	N19813 002	AUG 07, 1990	MAY	CTEC
		2.4MG/24HR	N19813 001	AUG 07, 1990	MAY	CTEC

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE PRESERVATIVE FREE

@ MARSAM

EQ 0.05MG BASE/ML

N74917 001 FEB 03, 1998 JAN DISC

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

AB	ALPHAPHARM	50MG	N75442 001	JUL 31, 2001	JUL	NEWA
AB		100MG	N75442 002	JUL 31, 2001	JUL	NEWA
AB		150MG	N75442 003	JUL 31, 2001	JUL	NEWA
	TAMBOCOR					
AB	3M	50MG	N18830 004	AUG 23, 1988	JUL	CFTG
AB		100MG	N18830 001	OCT 31, 1985	JUL	CFTG
AB +		150MG	N18830 003	JUN 03, 1988	JUL	CFTG

FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINE

AP AM PHARM PARTNERS 500MG/VIAL N75837 001 FEB 22, 2001 FEB NEWA

FLUDEOXYGLUCOSE, F-18

INJECTABLE; INJECTION

FLUDEOXYGLUCOSE F 18

+ DOWNTSTATE CLINCL 4-90mCi/ML N20306 002 SEP 25, 2001 SEP NEWA

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

@ CLAY PARK 0.01% N86810 001 MAR 04, 1982 APR DISC
@ 0.025% N86811 001 MAR 04, 1982 APR DISCFLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-SYNALAR

@ MEDICIS 0.025%;EQ 3.5MG BASE/GM N60700 001 JUN 11, 1963 MAY DISC

FLUOROMETHOLONE

SUSPENSION/DROPS; OPHTHALMIC

FLUOR-OP

AB NOVARTIS 0.1% N70185 001 FEB 27, 1986 FEB CAHN

FLUOROURACIL

CREAM; TOPICAL

CARAC

+ DERMIC LABS 0.5% N20985 001 OCT 27, 2000 MAY CTNA

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE

AB BARR EQ 20MG BASE N74803 001 AUG 02, 2001 AUG NEWA
AB DR REDDYS LABS LTD EQ 40MG BASE N75465 003 AUG 02, 2001 AUG NEWA

AB	GENEVA PHARMS PROZAC	EQ 10MG BASE	N75049 001 AUG 02, 2001 AUG NEWA
AB	LILLY	EQ 10MG BASE	N18936 006 DEC 23, 1992 AUG CFTG
AB		EQ 20MG BASE	N18936 001 DEC 29, 1987 AUG CFTG
AB +		EQ 40MG BASE	N18936 003 JUN 15, 1999 AUG CFTG
	CAPSULE, DELAYED REL PELLETS; ORAL PROZAC WEEKLY		
+ LILLY	SOLUTION; ORAL FLUOXETINE	EQ 90MG BASE	N21235 001 FEB 26, 2001 FEB NEWA
AT	TEVA PROZAC	EQ 20MG BASE/5ML	N75506 001 AUG 02, 2001 AUG NEWA
AT + LILLY	TABLET; ORAL FLUOXETINE HCL	EQ 20MG BASE/5ML	N20101 001 APR 24, 1991 AUG CFTG
AB	ALPHAPHARM +	EQ 10MG BASE EQ 20MG BASE	N75755 001 AUG 02, 2001 AUG NEWA N75755 002 AUG 02, 2001 AUG NEWA
AB + LILLY		EQ 10MG BASE	N20974 001 MAR 09, 1999 AUG CFTG
<u>FLUPHENAZINE DECANOATE</u>			
	INJECTABLE; IM-SC FLUPHENAZINE DECANOATE		
AO	APOTEX	25MG/ML	N75918 001 AUG 17, 2001 AUG NEWA
<u>FLUPHENAZINE HYDROCHLORIDE</u>			
	TABLET; ORAL PERMITIL		
>D>			
>D> BP	SCHERING	10MG	N12034 006 JAN 07, 1964 OCT DISC
>A>	@	10MG	N12034 006 JAN 07, 1964 OCT DISC
<u>FLURAZEPAM HYDROCHLORIDE</u>			
	CAPSULE; ORAL FLURAZEPAM HCL		
	CH CHELSEA LABS	15MG	N72368 001 MAR 30, 1989 JUL DISC
	PUREPAC PHARM	15MG	N71927 001 SEP 09, 1987 JUL DISC
	@	30MG	N71551 001 SEP 09, 1987 JUL DISC
<u>FLURBIPROFEN</u>			
	TABLET; ORAL FLURBIPROFEN		
AB	CARACO	50MG	N75058 001 APR 27, 2001 APR NEWA
AB		100MG	N75058 002 APR 27, 2001 APR NEWA
<u>FLUTAMIDE</u>			
	CAPSULE; ORAL EULEXIN		
AB +	SCHERING FLUTAMIDE	125MG	N18554 001 JAN 27, 1989 SEP CFTG
AB	BARR	125MG	N75820 001 SEP 18, 2001 SEP NEWA
AB	EON	125MG	N75818 001 SEP 18, 2001 SEP NEWA
>A> AB	IVAX PHARMS	125MG	N75780 001 SEP 19, 2001 OCT NEWA
AB	TEVA	125MG	N75298 001 SEP 18, 2001 SEP NEWA

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

AB	BARR	25MG	N75897 001	JAN 25, 2001	JAN	NEWA
AB		50MG	N75897 002	JAN 25, 2001	JAN	NEWA
AB		100MG	N75897 003	JAN 25, 2001	JAN	NEWA
>A>	AB GENPHARM	50MG	N75950 001	OCT 15, 2001	OCT	NEWA
>A>	AB INVAMED	100MG	N75950 002	OCT 15, 2001	OCT	NEWA
AB		25MG	N75887 001	JAN 05, 2001	JAN	NEWA
AB		50MG	N75887 002	JAN 05, 2001	JAN	NEWA
AB		100MG	N75887 003	JAN 05, 2001	JAN	NEWA
AB	SYNTON PHARMS	25MG	N75899 001	JAN 17, 2001	JAN	NEWA
AB		50MG	N75899 002	JAN 17, 2001	JAN	NEWA
AB		100MG	N75899 003	JAN 17, 2001	JAN	NEWA
AB	TORPHARM	25MG	N75902 001	MAY 07, 2001	MAY	NEWA
AB		50MG	N75902 002	MAY 07, 2001	MAY	NEWA
AB		100MG	N75902 003	MAY 07, 2001	MAY	NEWA
AB	WATSON LABS	25MG	N75894 001	APR 18, 2001	APR	NEWA
AB		50MG	N75894 002	APR 18, 2001	APR	NEWA
AB		100MG	N75894 003	APR 18, 2001	APR	NEWA
AB	ZENITH GOLDLINE	25MG	N75898 001	MAR 12, 2001	MAR	NEWA
AB		50MG	N75898 002	MAR 12, 2001	MAR	NEWA
AB		100MG	N75898 003	MAR 12, 2001	MAR	NEWA

FOLIC ACID

TABLET; ORAL

FOLIC ACID

>D>	AA IMPAX LABS	1MG	N80686 001	JUL 20, 1973	OCT	DISC
>A>	@	1MG	N80686 001	JUL 20, 1973	OCT	DISC

FOLLITROPIN ALFA

INJECTABLE; INJECTION

GONAL-F

+ SERONO

1,200 IU/VIAL

N20378 004 FEB 28, 2001 AUG NEWA

FORMOTEROL FUMARATE

CAPSULE; INHALATION

FORADIL

+ NOVARTIS

0.012MG/INH

N20831 001 FEB 16, 2001 FEB NEWA

GABAPENTIN

CAPSULE; ORAL

NEURONTIN

PFIZER

100MG

N20235 001 DEC 30, 1993 MAR CAHN

300MG

N20235 002 DEC 30, 1993 MAR CAHN

+

N20235 003 DEC 30, 1993 MAR CAHN

SOLUTION; ORAL

@ PARKE DAVIS

250MG/5ML

N21129 001 MAR 02, 2000 OCT CMFD

>D> +

250MG/5ML

N21129 001 MAR 02, 2000 OCT CMFD

GALANTAMINE HYDROBROMIDE

SOLUTION; ORAL				
REMINYL				
+ JANSSEN	4MG/ML			N21224 001 JUN 22, 2001 JUN NEWA
TABLET; ORAL				
JANSSEN	EQ 4MG BASE			N21169 001 FEB 28, 2001 FEB NEWA
	EQ 8MG BASE			N21169 002 FEB 28, 2001 FEB NEWA
+	EQ 12MG BASE			N21169 003 FEB 28, 2001 FEB NEWA

GEMFIBROZIL

TABLET; ORAL				
GEMFIBROZIL				
AB GENEVA PHARMS TECH	600MG			N74615 001 SEP 29, 1995 JAN CAHN

GENTAMICIN SULFATE

CREAM; TOPICAL				
GENTAMICIN SULFATE				
@ BAUSCH AND LOMB	EQ 0.1% BASE			N64056 001 APR 29, 1994 MAY DISC
INJECTABLE; INJECTION				

@ GENSIA SICOR PHARMS	EQ 10MG BASE/ML			N63149 001 NOV 21, 1991 MAY DISC
@	EQ 40MG BASE/ML			N63106 002 NOV 21, 1991 APR DISC
@ STERIS	EQ 10MG BASE/ML			N62318 002 AUG 20, 1981 APR DISC
@	EQ 40MG BASE/ML			N62318 001 JUN 02, 1981 APR DISC

U-GENCIN				
@ PHARMACIA AND UPJOHN	EQ 10MG BASE/ML			N62248 001 MAY 02, 1980 FEB WDRP
@	EQ 40MG BASE/ML			N62248 002 MAY 02, 1980 FEB WDRP

INJECTABLE; INTRATHECAL				
GARAMYCIN				

@ SCHERING	EQ 2MG BASE/ML			N50505 001 OCT 01, 1979 APR DISC
OINTMENT; OPHTHALMIC				
GENTACIDIN				

AT NOVARTIS	EQ 0.3% BASE			N62501 001 JUL 26, 1984 FEB CAHN
@	EQ 0.3% BASE			N62501 001 JUL 26, 1984 MAY DISC

OINTMENT; TOPICAL				
-------------------	--	--	--	--

GARAMYCIN				
@ SCHERING	EQ 0.1% BASE			N60463 001 MAR 15, 1966 SEP DISC
GENTAMICIN				

AT + CLAY PARK	EQ 0.1% BASE			N62351 001 FEB 18, 1982 SEP CTEC
GENTAMICIN SULFATE				
@ BAUSCH AND LOMB	EQ 0.1% BASE			N64054 001 APR 29, 1994 MAY DISC
SOLUTION/DROPS; OPHTHALMIC				
GENTACIDIN				

AT NOVARTIS	EQ 0.3% BASE			N62480 001 MAR 30, 1984 FEB CAHN
>A> GENTAK				

>A> AT AKORN	EQ 0.3% BASE			N64163 001 OCT 12, 2001 OCT NEWA
GENTAMICIN SULFATE				
@ ALCON UNIVERSAL	EQ 0.3% BASE			N62523 001 NOV 25, 1985 APR DISC

GLIPIZIDE

TABLET; ORAL				
--------------	--	--	--	--

GLIPIZIDE				
-----------	--	--	--	--

AB GENEVA PHARMS TECH	5MG			N74542 001 JUN 20, 1995 JAN CAHN
-----------------------	-----	--	--	----------------------------------

AB		10MG	N74542 002	JUN 20, 1995	JAN	CAHN
AB	TORPHARM	5MG	N75795 001	JUN 13, 2001	JUN	NEWA
AB		10MG	N75795 002	JUN 13, 2001	JUN	NEWA

GLUTETHIMIDE

TABLET; ORAL						
GLUTETHIMIDE						
© CELLTECH PHARMS	500MG		N85171 001	DEC 22, 1976	SEP	DISC

GLYCOPYRROLATE

INJECTABLE; INJECTION						
GLYCOPYRROLATE						
© GENSIA SICOR PHARMS	0.2MG/ML		N81169 001	SEP 10, 1991	MAY	DISC
TABLET; ORAL						
ROBINUL						
+ FIRST HORIZON	1MG		N12827 001	AUG 11, 1961	AUG	CAHN
ROBINUL FORTE						
+ FIRST HORIZON	2MG		N12827 002	AUG 11, 1961	AUG	CAHN

GRANISETRON HYDROCHLORIDE

SOLUTION; ORAL						
KYTRIL						
+ ROCHE	EQ 2MG BASE/10ML		N21238 001	JUN 27, 2001	JUN	NEWA

GRISEOFULVIN, MICROCRYSTALLINE

SUSPENSION; ORAL						
GRIFULVIN V						
+ J AND J	125MG/5ML		N62483 001	JAN 26, 1984	MAR	CRLD
© JOHNSON AND JOHNSON	125MG/5ML		N50448 001	MAY 19, 1972	MAR	DISC

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL						
GRISACTIN ULTRA						
© WYETH AYERST	125MG		N62178 001	MAR 13, 1980	APR	DISC
©	250MG		N62178 002	MAR 13, 1980	APR	DISC
ULTRAGRIS-165						
© SIDMAK LABS NJ	165MG		N62645 001	JUN 30, 1992	MAY	DISC
ULTRAGRIS-330						
© SIDMAK LABS NJ	330MG		N62646 001	JUN 30, 1992	MAY	DISC

HALOPERIDOL

TABLET; ORAL						
HALOPERIDOL						
© DANBURY PHARMA	1MG		N70982 001	MAR 06, 1987	JUL	DISC
© ROXANE	0.5MG		N71128 001	FEB 17, 1987	AUG	DISC
©	1MG		N71129 001	FEB 17, 1987	AUG	DISC
©	2MG		N71130 001	FEB 17, 1987	AUG	DISC
©	5MG		N71131 001	FEB 17, 1987	AUG	DISC
©	20MG		N71133 001	MAY 12, 1987	AUG	DISC

HALOPERIDOL LACTATE

CONCENTRATE; ORAL				
HALOPERIDOL INTENSOL				
@ ROXANE	EQ 2MG BASE/ML	N72045 001	APR 12, 1988	AUG DISC
INJECTABLE; INJECTION				
HALOPERIDOL				
AP AM PHARM PARTNERS	EQ 5MG BASE/ML	N75689 001	MAR 09, 2001	JUN CTNA
AP BEDFORD	EQ 5MG BASE/ML	N75858 001	JUN 18, 2001	JUN NEWA
AP GENESIS SICOR PHARMS	EQ 5MG BASE/ML	N76035 001	AUG 29, 2001	AUG NEWA
HALOPERIDOL LACTATE				
AP AM PHARM PARTNERS	EQ 5MG BASE/ML	N75689 001	MAR 09, 2001	MAR NEWA

HALOTHANE

LIQUID; INHALATION				
HALOTHANE				
@ BH	99.99%	N84977 001	JUL 14, 1976	JAN DISC

HEPARIN SODIUM

INJECTABLE; INJECTION				
HEPARIN SODIUM				
@ ABBOTT	10,000 UNITS/ML	N40095 001	JUL 26, 1996	MAY DISC
HEPARIN SODIUM PRESERVATIVE FREE				
@ PHARMA SERVE NY	1,000 UNITS/ML	N86129 001	FEB 22, 1980	FEB WDRP

HOMATROPINE METHYLBROMIDE

TABLET; ORAL				
HOMAPIN-10				
@ MISSION PHARMA	10MG	N86308 001	APR 11, 1979	JUL DISC
HOMAPIN-5				
@ MISSION PHARMA	5MG	N86309 001	APR 11, 1979	JUL DISC

HYALURONIDASE

INJECTABLE; INJECTION				
WYDASE				
@ WYETH AYERST	150 UNITS/ML	N06343 002	MAR 22, 1950	JUL DISC
@	150 UNITS/VIAL	N06343 006	MAR 06, 1951	JUL DISC
@	1,500 UNITS/VIAL	N06343 005	MAR 06, 1951	JUL DISC

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION				
HYDRALAZINE HCL				
AP AM PHARM PARTNERS	20MG/ML	N40388 001	MAR 13, 2001	MAR NEWA

HYDRALAZINE HYDROCHLORIDE; HYDROCHLORTHIAZIDE; RESERPINE

TABLET; ORAL				
RESERPINE, HYDRALAZINE HCL AND HYDROCHLORTHIAZIDE				
@ DANBURY PHARMA	25MG;15MG;0.1MG	N85549 001	SEP 29, 1977	MAY DISC

HYDROCHLORTHIAZIDE

TABLET; ORAL				
HYDROCHLORTHIAZIDE				
@ DANBURY PHARMA	50MG	N83232 001	JAN 24, 1975	MAY DISC

© HALSEY	25MG	N83972 001	OCT 03, 1974	MAY	DISC
©	50MG	N83972 002	OCT 03, 1974	MAY	DISC
© IMPAX LABS	25MG	N84029 001	JUL 05, 1977	MAY	DISC
©	50MG	N83607 002	JUN 06, 1977	MAY	DISC
© PHARMERAL	25MG	N84325 001	JUN 24, 1976	MAY	DISC
©	50MG	N84324 001	JUN 24, 1976	MAY	DISC
© PVT FORM	50MG	N86597 001	OCT 11, 1978	JUL	DISC
© WEST WARD	50MG	N84878 001	JAN 31, 1977	MAY	DISC

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

HYDRO-RESERP

© ABC HOLDING	50MG;0.125MG	N84714 002	JUN 29, 1982	MAY	DISC
HYDROCHLOROTHIAZIDE W/ RESERPINE					
© DANBURY PHARMA	25MG;0.125MG	N84466 001	JAN 07, 1977	MAY	DISC
©	50MG;0.125MG	N84467 001	JAN 07, 1977	MAY	DISC
RESERPINE AND HYDROCHLOROTHIAZIDE-50					
© WEST WARD	50MG;0.125MG	N88189 001	MAY 10, 1984	FEB	WDRP

HYDROCORTISONE

CREAM; TOPICAL

DERMACORT

© MONARCH PHARMS	1%	N83011 002	APR 26, 1973	SEP	WDAG
HC (HYDROCORTISONE)					
© C AND M PHARMA	0.5%	N80482 003	MAR 20, 1973	FEB	WDRP
©	1%	N80482 004	MAR 20, 1973	FEB	WDRP
HYDROCORTISONE					
© TOPIDERM	1%	N89273 001	FEB 17, 1989	FEB	WDRP
NUTRACORT					
© HEALTHPOINT	1%	N80442 003	APR 04, 1972	JUL	DISC
PROCTOCORT					
© MONARCH PHARMS	1%	N83011 001	APR 26, 1973	FEB	DISC

LOTION; TOPICAL

ACTICORT

© BAKER NORTON

BETA-HC

© BETA DERMAC

GLYCORT

© HERAN

HYDROCORTISONE

© MERICON

©

OINTMENT; TOPICAL

HC (HYDROCORTISONE)

© C AND M PHARMA

POWDER; FOR RX COMPOUNDING

H-CORT

© TORCH

SOLUTION; TOPICAL

TEXACORT

AT + SIRIUS LABS

+

TABLET; ORAL

1%	N86535 001	FEB 04, 1981	JUL	DISC
1%	N89495 001	JAN 25, 1988	FEB	WDRP
1%	N87489 001	OCT 03, 1983	FEB	WDRP
0.5%	N85282 001	JUN 05, 1978	MAY	DISC
1%	N85282 002	FEB 26, 1987	MAY	DISC
1%	N80481 002	MAR 20, 1973	FEB	WDRP
100%	N87834 001	MAR 29, 1982	FEB	WDRP
1%	N80425 001	DEC 22, 1971	JUN	CAHN
2.5%	N81271 001	APR 17, 1992	MAY	CAHN

HYDROCORTISONE

TABLET; ORAL

HYDROCORTISONE

>D>	BP	IMPAK LABS	20MG	N80781 001	AUG 02, 1973	OCT	DISC
>A>	⑧		20MG	N80781 001	AUG 02, 1973	OCT	DISC
	⑧ LANNETT		20MG	N85070 001	MAY 07, 1976	MAY	DISC

HYDROCORTISONE ACETATE

CREAM; TOPICAL

MICORT-HC

FERNDALE LABS

2.5%

N40396 001 FEB 27, 2001 FEB NEWA

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

OINTMENT; TOPICAL

NEO-CORTEF

⑧ PHARMACIA AND UPJOHN 1%;EQ 3.5MG BASE/GM N60751 002 MAY 18, 1965 APR DISC

SUSPENSION/DROPS; OPHTHALMIC

COR-OTICIN

⑧ AKORN 1.5%;EQ 3.5MG BASE/ML N60188 001 OCT 26, 1968 FEB WDRP

HYDROCORTISONE VALERATE

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

AB ALTANA 0.2% N75085 001 JUL 31, 2001 JUL NEWA

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

NEO-OTOSOL-HC

⑧ ALCON 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML N62423 001 AUG 25, 1983 APR DISC

SUSPENSION/DROPS; OPHTHALMIC

CORTISPORIN

+ MONARCH PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML N50169 001 DEC 18, 1964 MAY CTEC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

⑧ ALCON UNIVERSAL 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML N62874 001 MAY 11, 1988 MAY DISC

SUSPENSION/DROPS; OTIC

⑧ ALCON UNIVERSAL 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML N62488 001 NOV 06, 1985 APR DISC

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HCL

⑧ ABBOTT 50MG/ML N86821 001 SEP 05, 1979 JUL DISC

⑧ AM PHARM PARTNERS 25MG/ML N88184 001 MAR 31, 1983 JUL DISC

⑧ 50MG/ML N88185 001 MAR 31, 1983 JUL DISC

⑧ STERIS 25MG/ML N85778 001 OCT 05, 1979 MAY DISC

TABLET; ORAL

⑧ PAR PHARM 10MG N87602 001 JAN 22, 1982 JUL DISC

⑧ 25MG N87603 001 JAN 22, 1982 JUL DISC

⑧ 50MG N87604 001 JAN 22, 1982 JUL DISC

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

© GENEVA PHARMS

EQ 50MG HCL

N81128 001 JUN 28, 1991 MAY DISC

©

EQ 100MG HCL

N81129 001 JUN 28, 1991 MAY DISC

© VANGARD

EQ 50MG HCL

N88393 001 SEP 19, 1983 FEB WDRP

IBUPROFEN

TABLET; ORAL

IBUPROFEN

>A>	AB	DR REDDYS LABS INC	400MG	N76112 001	OCT 31, 2001	OCT	NEWA
>A>	AB		600MG	N76112 002	OCT 31, 2001	OCT	NEWA
>A>	AB		800MG	N76112 003	OCT 31, 2001	OCT	NEWA
>D>	AB	LEDERLE	400MG	N70629 001	SEP 19, 1986	OCT	DISC
>A>		©	400MG	N70629 001	SEP 19, 1986	OCT	DISC
>D>	AB		600MG	N70630 001	SEP 19, 1986	OCT	DISC
>A>		©	600MG	N70630 001	SEP 19, 1986	OCT	DISC

IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

NOVARTIS

50MG

N21335 001 MAY 10, 2001 MAY NEWA

+

100MG

N21335 002 MAY 10, 2001 MAY NEWA

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HCL

© ROXANE

25MG

N83799 002 AUG 05, 1977 SEP DISC

©

50MG

N83799 003 AUG 05, 1977 SEP DISC

TOFRANIL

AB TYCO HLTHCARE

10MG

N87844 001 MAY 22, 1984 JUN CAHN

AB

25MG

N87845 001 MAY 22, 1984 JUN CAHN

AB +

50MG

N87846 001 MAY 22, 1984 JUN CAHN

IMIPRAMINE PAMOATE

CAPSULE; ORAL

TOFRANIL-PM

TYCO HLTHCARE

EQ 75MG HCL

N17090 001 MAR 15, 1973 JUN CAHN

EQ 100MG HCL

N17090 004 MAR 08, 1974 JUN CAHN

EQ 125MG HCL

N17090 003 MAR 08, 1974 JUN CAHN

+

EQ 150MG HCL

N17090 002 MAR 15, 1973 JUN CAHN

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

AB GENEVA PHARMS TECH

1.25MG

N74594 001 MAY 23, 1996 JAN CAHN

AB

2.5MG

N74594 002 MAY 23, 1996 JAN CAHN

INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

MERCK RES LABS

EQ 100MG BASE

N20685 006 APR 19, 2000 AUG NEWA

IPRATROPIUM BROMIDE

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

AN	ASLUNG PHARM	0.02%	N75693 001	JAN 26, 2001	JAN	NEWA
>A>	AN BAUSCH AND LOMB	0.02%	N75835 001	OCT 15, 2001	OCT	NEWA
AN	NEPHRON	0.02%	N75562 001	SEP 27, 2001	SEP	NEWA
AN	NOVEX	0.02%	N75441 001	MAR 28, 2001	MAR	NEWA
AN	WARRICK PHARMS	0.02%	N75507 001	JAN 19, 2001	JAN	NEWA

ISOFLURANE

LIQUID; INHALATION

ISOFLURANE

AN	MINRAD	99.9%	N74416 001	SEP 30, 1994	FEB	CAHN
----	--------	-------	------------	--------------	-----	------

ISONIAZID

SYRUP; ORAL

ISONIAZID

+ CAROLINA MEDCL	50MG/5ML	N88235 001	NOV 10, 1983	MAY	CTEC
@ MIKART	50MG/5ML	N81118 001	JUL 21, 1997	MAY	DISC
TABLET; ORAL		N80136 001	NOV 13, 1970	MAY	DISC

@ HALSEY

100MG

ISOSORBIDE MONONITRATE

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

AB	ZENITH GOLDLINE	30MG	N75448 002	AUG 07, 2001	AUG	NEWA
AB		120MG	N75448 003	AUG 07, 2001	AUG	NEWA

ISOTRETINOIN

CAPSULE; ORAL

ACCUVANE

+ HLR

20MG

N18662 004 MAR 28, 1983 APR CTEC

KANAMYCIN SULFATE

INJECTABLE; INJECTION

KANAMYCIN SULFATE

@ LOCH

EQ 75MG BASE/2ML

N63021 001 JUL 31, 1992 MAY DISC

@

EQ 500MG BASE/2ML

N63022 001 JUL 31, 1992 MAY DISC

@

EQ 1GM BASE/3ML

N63025 001 JUL 31, 1992 APR DISC

@ STERIS

EQ 1GM BASE/3ML

N62520 003 MAY 09, 1985 MAY DISC

KANTREX

+ APOTHECON

EQ 75MG BASE/2ML

N61901 003 MAR 06, 1975 MAY CTEC

+

EQ 500MG BASE/2ML

N61901 001 MAR 06, 1975 MAY CTEC

+

EQ 1GM BASE/3ML

N61901 002 MAR 06, 1975 MAY CTEC

KETOCONAZOLE

CREAM; TOPICAL

NIZORAL

AB + JANSSEN

2%

N19084 001 DEC 31, 1985 JUL CAHN

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

AP	APOTEX	15MG/ML	N75631 002	JUN 29, 2001	JUN	NEWA
AP		30MG/ML	N75626 001	JUL 24, 2001	JUL	NEWA
AP		30MG/ML	N75631 001	JUN 29, 2001	JUN	NEWA
	@ APOTHECON	15MG/ML	N75348 001	NOV 28, 2000	MAY	DISC
	@	30MG/ML	N75348 002	NOV 28, 2000	MAY	DISC
>D>	AP BEDFORD	15MG/ML	N75230 002	OCT 25, 1999	OCT	DISC
>A>	@	15MG/ML	N75230 002	OCT 25, 1999	OCT	DISC
>D>	AP	30MG/ML	N75230 001	OCT 25, 1999	OCT	DISC
>A>	@	30MG/ML	N75230 001	OCT 25, 1999	OCT	DISC

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HCL

@ APOTHECON

5MG/ML

N75355 001 NOV 29, 1999 MAY DISC

TRANDATE

AP + PROMETHEUS LABS

5MG/ML

N19425 001 DEC 31, 1985 MAY CAHN

LACTULOSE

SOLUTION; ORAL

LACTULOSE

AA VINTAGE PHARMS

10GM/15ML

N75993 001 JUL 26, 2001 JUL NEWA

SOLUTION; ORAL, RECTAL

@ ROXANE

10GM/15ML

N73590 001 MAY 29, 1992 SEP DISC

LAMOTRIGINE

TABLET, CHEWABLE; ORAL

LAMICTAL CD

GLAXO WELLCOME

2MG

N20764 004 SEP 08, 2000 MAR NEWA

LANSOPRAZOLE

FOR SUSPENSION, EXTENDED RELEASE; ORAL

PREVACID

TAP PHARM

15MG/PACKET

N21281 001 MAY 03, 2001 MAY NEWA

+

30MG/PACKET

N21281 002 MAY 03, 2001 MAY NEWA

LATANOPROST; TIMOLOL MALEATESOLUTION/DROPS; OPHTHALMICXALCOM+ PHARMACIA AND UPJOHN

0.005%;0.5%

N21219 001 OCT 02, 2001 OCT NEWA

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM PRESERVATIVE FREE

AP LUITPOLD

EQ 50MG BASE/VIAL

N40338 001 JAN 31, 2001 JAN NEWA

LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR

AP + SIGMA TAU

200MG/ML

N20182 001 DEC 16, 1992 MAR CFTG

LEVOCARNITINE

INJECTABLE; INJECTION

LEVOCARNITINE

AP	BEDFORD	200MG/ML	N75567 001	MAR 29, 2001	MAR	NEWA
AP	GENSIA SICOR PHARMS	200MG/ML	N75881 001	MAR 29, 2001	MAR	NEWA
AP	LUITPOLD	200MG/ML	N75861 001	JUN 22, 2001	JUN	NEWA

LEVODOPA

CAPSULE; ORAL

DOPAR

@ SHIRE LABS

250MG

N16913 001 JUN 04, 1970 MAY DISC

TABLET; ORAL

@ SHIRE LABS

250MG

N16913 004 JUL 06, 1972 JUN DISC

@

500MG

N16913 005 JUL 06, 1972 JUN DISC

LARODOPA

ROCHE

250MG

N16912 003 JUN 04, 1970 JUN CRLD

+

500MG

N16912 004 JUN 04, 1970 JUN CRLD

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

POLOCAINE W/ LEVONORDEFRIN

AP	DENTSPLY PHARM	0.05MG/ML;2%
----	----------------	--------------

N89517 001 APR 14, 1988 JUL CAHN

LEVOHYDROXYNE SODIUM

TABLET; ORAL

LEVOXYL

BX	+	JONES PHARMA	0.025MG	N21301 001	MAY 25, 2001	MAY	NEWA
BX			0.025MG	N21301 001	MAY 25, 2001	JUL	CRLD
BX			0.05MG	N21301 002	MAY 25, 2001	MAY	NEWA
BX			0.075MG	N21301 003	MAY 25, 2001	MAY	NEWA
BX			0.088MG	N21301 004	MAY 25, 2001	MAY	NEWA
BX			0.1MG	N21301 005	MAY 25, 2001	MAY	NEWA
BX			0.112MG	N21301 006	MAY 25, 2001	MAY	NEWA
BX			0.125MG	N21301 007	MAY 25, 2001	MAY	NEWA
BX			0.137MG	N21301 008	MAY 25, 2001	MAY	NEWA
BX			0.15MG	N21301 009	MAY 25, 2001	MAY	NEWA
BX			0.175MG	N21301 010	MAY 25, 2001	MAY	NEWA
BX			0.2MG	N21301 011	MAY 25, 2001	MAY	NEWA
BX	+		0.3MG	N21301 012	MAY 25, 2001	MAY	NEWA
		UNITHROID		N21301 012	MAY 25, 2001	JUL	CRLD
BX		STEVENS J	0.025MG	N21210 001	AUG 21, 2000	MAY	CTEC
BX			0.05MG	N21210 002	AUG 21, 2000	MAY	CTEC
BX			0.075MG	N21210 003	AUG 21, 2000	MAY	CTEC
BX			0.088MG	N21210 004	AUG 21, 2000	MAY	CTEC
BX			0.1MG	N21210 005	AUG 21, 2000	MAY	CTEC
BX			0.112MG	N21210 006	AUG 21, 2000	MAY	CTEC
BX			0.125MG	N21210 007	AUG 21, 2000	MAY	CTEC
BX			0.15MG	N21210 008	AUG 21, 2000	MAY	CTEC
BX			0.175MG	N21210 009	AUG 21, 2000	MAY	CTEC
BX	+		0.2MG	N21210 010	AUG 21, 2000	MAY	CTEC
			0.3MG	N21210 011	AUG 21, 2000	MAY	CTEC

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION					
LIDOCaine HCl					
@ STERIS	1%	N80377 001	FEB 20, 1974	JUL	DISC
@	2%	N80377 002	FEB 20, 1974	JUL	DISC
LIDOCATON					
@ PHARMATON	2%	N84727 001	AUG 17, 1983	FEB	WDRP
XYLOCAINE					
@ ASTRazeneca	2%	N06488 002	NOV 19, 1948	SEP	DISC
AP + DENTSPly PHARM	2%	N21380 001	NOV 19, 1948	SEP	NEWA

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION					
LINCOcIN					
+ PHARMACIA AND UPJOHN	EQ 300MG BASE/ML	N50317 001	DEC 29, 1964	MAY	CTEC
LINCOMYCIN HCl					
@ STERIS	EQ 300MG BASE/ML	N63180 001	APR 16, 1991	MAY	DISC

LISINOPRIL

TABLET; ORAL					
ZESTRIL					
AB ASTRazeneca	10MG	N19777 002	MAY 19, 1988	APR	CTEC

LITHIUM CARBONATE

CAPSULE; ORAL					
ESKALITH					
AB SMITHKLINE BEECHAM	300MG	N16860 001	APR 06, 1970	JUN	CRDL
LITHIUM CARBONATE					
AB ABLE	300MG	N76121 001	SEP 27, 2001	SEP	NEWA
+ ROXANE	600MG	N17812 003	JAN 28, 1987	JUN	CRDL

LOMEFLOXACIN HYDROCHLORIDE

TABLET; ORAL					
MAXAQIN					
+ UNIMED PHARMS	EQ 400MG BASE	N20013 001	FEB 21, 1992	SEP	CAHN

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL					
LOPERAMIDE HCl					
@ ROXANE	2MG	N73080 001	NOV 27, 1991	JUL	DISC

LORATADINE

TABLET; ORAL					
CLARITIN					
AB + SCHERING	10MG	N19658 001	APR 12, 1993	SEP	CFTG

LORAZEPAM

TABLET; ORAL					
LORAZEPAM					
AB RANBAXY	0.5MG	N76045 001	AUG 29, 2001	AUG	NEWA
AB	1MG	N76045 002	AUG 29, 2001	AUG	NEWA
AB	2MG	N76045 003	AUG 29, 2001	AUG	NEWA

@ WATSON LABS	0.5MG	N71086 001 MAR 23, 1987 JUL DISC
<u>LOSARTAN POTASSIUM</u>		
TABLET; ORAL		
COZAAR		
MERCK RES LABS	50MG	
+	100MG	N20386 002 APR 14, 1995 AUG CRLD N20386 003 OCT 13, 1998 AUG NEWA
<u>LOTEPREDNOL ETABONATE</u>		
SUSPENSION/DROPS; OPHTHALMIC		
ALREX		
>A> + BAUSCH AND LOMB	0.2%	N20803 001 MAR 09, 1998 OCT CAHN
>D> + PHARMOS	0.2%	N20803 001 MAR 09, 1998 OCT CAHN
LOTEMAX		
>A> + BAUSCH AND LOMB	0.5%	N20583 001 MAR 09, 1998 OCT CAHN
>D> + PHARMOS	0.5%	N20583 001 MAR 09, 1998 OCT CAHN
<u>MANNITOL</u>		
INJECTABLE; INJECTION		
MANNITOL 25%		
@ ASTRAZENECA	12.5GM/50ML	N89240 001 MAY 06, 1987 SEP DISC
<u>MECLIZINE HYDROCHLORIDE</u>		
TABLET; ORAL		
MECLIZINE HCL		
@ CHELSEA LABS	12.5MG	N85269 001 NOV 11, 1976 MAY DISC
<u>MEGESTROL ACETATE</u>		
SUSPENSION; ORAL		
MEGACE		
AB + BRISTOL MYERS SQUIBB	40MG/ML	N20264 001 SEP 10, 1993 JUL CFTG
MEGESTROL ACETATE		
AB PAR PHARM	40MG/ML	N75671 001 JUL 25, 2001 JUL NEWA
<u>MELOXICAM</u>		
TABLET; ORAL		
MOBIC		
BOEHRINGER INGELHEIM	7.5MG	N20938 001 APR 13, 2000 JUL CRLD
+	15MG	N20938 002 AUG 23, 2000 JUL NEWA
<u>MENOTROPINS (FSH;LH)</u>		
INJECTABLE; INJECTION		
>D> HUMEGON		
>D> AB ORGANON	75 IU/VIAL;75 IU/VIAL	N20328 001 SEP 01, 1994 OCT DISC
>A> @	75 IU/VIAL;75 IU/VIAL	N20328 001 SEP 01, 1994 OCT DISC
>D> AB	150 IU/VIAL;150 IU/VIAL	N20328 002 SEP 01, 1994 OCT DISC
>A> @	150 IU/VIAL;150 IU/VIAL	N20328 002 SEP 01, 1994 OCT DISC
PERGONAL		
>D> AB + SERONO	75 IU/AMP;75 IU/AMP	N17646 001 AUG 22, 1975 OCT CTEC
>A> BX +	75 IU/AMP;75 IU/AMP	N17646 001 AUG 22, 1975 OCT CTEC
>D> AB +	150 IU/AMP;150 IU/AMP	N17646 002 MAY 20, 1985 OCT CTEC
>A> BX +	150 IU/AMP;150 IU/AMP	N17646 002 MAY 20, 1985 OCT CTEC

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HCL

a ASTRAZENECA

50MG/ML

N89784 001 MAR 31, 1989 JUN DISC

a

100MG/ML

N89788 001 MAR 31, 1989 JUN DISC

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPIVACAINE HCL

a INT'L MEDICATION

1%

N87509 001 OCT 05, 1982 JUL DISC

POLOCAINE

AP DENTSPLY PHARM

3%

N88653 001 AUG 21, 1984 JUL CAHN

MEPROBAMATE

TABLET; ORAL

AMOSENE

a FERNDALE LABS

400MG

N84030 001 MAY 10, 1974 FEB WDRP

MEPROBAMATE

a HALSEY

400MG

N80699 002 OCT 16, 1972 MAY DISC

a IMPAX LABS

200MG

N14322 002 JUL 23, 1973 AUG DISC

a

400MG

N14322 001 JUL 15, 1963 AUG DISC

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL

SOLAGE

+ WESTWOOD SQUIBB

2%;0.01%

N20922 001 DEC 10, 1999 JUN CAHN

MESALAMINE

SUPPOSITORY; RECTAL

CANASA

+ AXCAN SCANDIPHARM

500MG

N21252 001 JAN 05, 2001 JAN NEWA

MESNA

INJECTABLE; INTRAVENOUS

MESNA

AP AM PHARM PARTNERS

100MG/ML

N75811 001 APR 26, 2001 APR NEWA

AP GENESIA SICOR PHARMS

100MG/ML

N75764 001 APR 27, 2001 APR NEWA

MESNEX

AP + ASTA

100MG/ML

N19884 001 DEC 30, 1988 APR CFTG

METAPROTERENOL SULFATE

SOLUTION; INHALATION

METAPROTERENOL SULFATE

AN NEPHRON

0.4%

N71855 001 JUL 14, 1988 AUG CAHN

AN NOVEX

0.6%

N71726 001 JUL 14, 1988 AUG CAHN

AN NOVEX

0.4%

N75402 001 FEB 28, 2001 FEB NEWA

AN NOVEX

0.6%

N75403 001 FEB 28, 2001 FEB NEWA

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DESOXYN

+ ABBOTT

5MG

N05378 002 DEC 31, 1943 JUL CTEC

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

METHAMPHETAMINE HCL

@ REXAR

5MG

@

10MG

N84931 001 JAN 19, 1976 JUL DISC
N84931 002 AUG 22, 1977 JUL DISCMETHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

@ APPLIED ANAL

25MG

@

50MG

N40011 001 JUL 17, 1997 MAY DISC
N40011 002 JUL 17, 1997 MAY DISCMETHIMAZOLE

TABLET; ORAL

METHIMAZOLE

AB EON

5MG

AB

10MG

+ GENPHARM

20MG

N40411 001 MAR 27, 2001 MAR NEWA
N40411 002 MAR 27, 2001 MAR NEWA
N40350 003 JUN 07, 2001 JUN NEWAMETHOTREXATE SODIUM

TABLET; ORAL

TREXALL

BARR

EQ 5MG BASE

N40385 001 MAR 21, 2001 MAR NEWA
N40385 002 MAR 21, 2001 MAR NEWA
N40385 003 MAR 21, 2001 MAR NEWA
N40385 004 MAR 21, 2001 MAR NEWA

+

EQ 7.5MG BASE

EQ 10MG BASE

EQ 15MG BASE

N40385 001 MAR 21, 2001 MAR NEWA
N40385 002 MAR 21, 2001 MAR NEWA
N40385 003 MAR 21, 2001 MAR NEWA
N40385 004 MAR 21, 2001 MAR NEWAMETHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

VASOXYL

@ GLAXO WELLCOME

20MG/ML

N06772 001 MAR 28, 1949 SEP WDAG

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

@ PVT FORM

2.5MG

PAMINE

N80970 001 OCT 18, 1976 MAY DISC

+ BRADLEY PHARMS

2.5MG

N08848 001 APR 09, 1953 MAY CTEC

METHYLCLOTHIAZIDE

TABLET; ORAL

METHYLCLOTHIAZIDE

@ PAR PHARM

2.5MG

@

5MG

N89135 001 FEB 12, 1986 JUL DISC
N89136 001 FEB 12, 1986 JUL DISCMETHYLDOPA

TABLET; ORAL

METHYLDOPA

@ LEDERLE

125MG

N70070 003 OCT 15, 1985 MAY DISC

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL					
METADATE CD					
+ CELLTECH PHARMS	20MG	N21259	001	APR 03, 2001	APR NEWA
TABLET; ORAL					
METHYLPHENIDATE HCL					
AB ABLE	5MG	N40404	001	MAR 29, 2001	MAR NEWA
AB	10MG	N40404	002	MAR 29, 2001	MAR NEWA
AB	20MG	N40404	003	MAR 29, 2001	MAR NEWA
TABLET, EXTENDED RELEASE; ORAL					
METADATE ER					
AB CELLTECH PHARMS	10MG	N40306	001	OCT 20, 1999	APR CTEC
METHYLPHENIDATE HCL					
AB ABLE	20MG	N76032	001	MAY 09, 2001	MAY NEWA
AB DANBURY PHARMA	20MG	N40410	001	FEB 09, 2001	FEB NEWA

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION					
DEPO-MEDROL					
PHARMACIA AND UPJOHN	40MG/ML	N11757	001	APR 27, 1959	MAY CTEC
METHYLPREDNISOLONE ACETATE					
@ STERIS	40MG/ML	N85600	001	MAR 14, 1979	MAY DISC

METHYLPREDNISOLONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL					
NEO-MEDROL ACETATE					
Ø PHARMACIA AND UPJOHN	0.25%;EQ 3.5MG BASE/GM	N60611	002	DEC 07, 1964	MAY DISC
Ø	1%;EQ 3.5MG BASE/GM	N60611	001	DEC 07, 1964	MAY DISC

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION					
METHYLPREDNISOLONE SODIUM SUCCINATE					
Ø GENESIA SICOR PHARMS	EQ 500MG BASE/VIAL	N81267	001	NOV 30, 1992	MAY DISC
Ø	EQ 1GM BASE/VIAL	N81268	001	NOV 30, 1992	MAY DISC

METHYLTESTOSTERONE

TABLET; Buccal					
ORETON					
Ø SCHERING	10MG	N80281	001	AUG 03, 1979	FEB DISC
TABLET; Buccal/Sublingual					
METHYLTESTOSTERONE					
Ø IMPAX LABS	10MG	N84287	001	JUL 16, 1974	JUL DISC
Ø LILLY	10MG	N80256	001	DEC 22, 1971	JUL DISC
TABLET; ORAL					
Ø LILLY	25MG	N80256	002	DEC 22, 1971	JUL DISC

METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC					
METIPRANOLOL					
AT FALCON PHARMS	0.3%	N75720	001	AUG 06, 2001	AUG NEWA
OPTIPRANOLOL					
AT + BAUSCH AND LOMB	0.3%	N19907	001	DEC 29, 1989	AUG CFTG

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION					
METOCLOPRAMIDE HCL					
@ ABBOTT	EQ 5MG BASE/ML	N70506 001	JUN 22, 1989	MAY	DISC
SOLUTION; INJECTION					
METOCLOPRAMIDE					
AA UDL	EQ 5MG BASE/5ML	N75051 001	JAN 26, 2001	JAN	NEWA
SOLUTION; ORAL					
AA UDL	EQ 5MG BASE/5ML	N75051 001	JAN 26, 2001	MAY	CDFR
TABLET; ORAL					
METOCLOPRAMIDE HCL					
AB GENEVA PHARMS TECH	EQ 5MG BASE	N74478 001	OCT 05, 1995	JAN	CAHN
AB	EQ 10MG BASE	N74478 002	OCT 05, 1995	JAN	CAHN
@ MUTUAL PHARM	EQ 5MG BASE	N71536 002	JAN 16, 1997	JUL	DISC
@	EQ 10MG BASE	N71536 001	APR 28, 1993	JUL	DISC

METOCURINE IODIDE

INJECTABLE; INJECTION					
METUBINE IODIDE					
@ LILLY	2MG/ML	N06632 003	FEB 15, 1952	SEP	WDAG

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL					
TOPROL-XL					
+ ASTRAZENECA	EQ 25MG TARTRATE	N19962 004	FEB 05, 2001	FEB	NEWA
	EQ 25MG TARTRATE	N19962 004	FEB 05, 2001	JUL	CRLD
	EQ 100MG TARTRATE	N19962 002	JAN 10, 1992	JUL	CRLD

METRONIDAZOLE

INJECTABLE; INJECTION					
METRO I.V.					
@ B BRAUN	500MG/100ML	N18674 001	AUG 31, 1982	MAY	DISC
METRONIDAZOLE					
@ ABBOTT	500MG/100ML	N18889 001	NOV 18, 1983	MAY	DISC
@ ELKINS SINK	500MG/100ML	N18907 001	MAR 30, 1984	MAY	DISC
TABLET; ORAL					
PROTOSTAT					
@ JOHNSON RW	250MG	N18871 001	MAR 02, 1983	MAR	DISC
@	500MG	N18871 002	MAR 02, 1983	MAR	DISC

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION					
MEZLIN					
@ BAYER	EQ 1GM BASE/VIAL	N62372 005	JAN 13, 1983	MAY	DISC
@	EQ 2GM BASE/VIAL	N62372 001	MAY 13, 1982	MAY	DISC
@	EQ 3GM BASE/VIAL	N62372 002	MAY 13, 1982	MAY	DISC
@	EQ 4GM BASE/VIAL	N62372 003	MAY 13, 1982	MAY	DISC
@	EQ 20GM BASE/VIAL	N62372 004	MAR 02, 1988	MAY	DISC

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HCL

>D>	AP	AESGEN	EQ 1MG BASE/ML	N75154 002	JUN 20, 2000	OCT	CAHN
>D>	AP		EQ 5MG BASE/ML	N75154 001	JUN 20, 2000	OCT	CAHN
>A>	AP	AM PHARM PARTNERS	EQ 1MG BASE/ML	N75154 002	JUN 20, 2000	OCT	CAHN
>A>	AP		EQ 5MG BASE/ML	N75154 001	JUN 20, 2000	OCT	CAHN
	@	APOTHECON	EQ 1MG BASE/ML	N75620 001	NOV 01, 2000	MAY	DISC
	@		EQ 5MG BASE/ML	N75620 002	NOV 01, 2000	MAY	DISC
	@		EQ 5MG BASE/ML	N75641 001	OCT 19, 2000	MAY	DISC
	@	ASTRAZENECA	EQ 5MG BASE/ML	N75263 001	JUN 26, 2000	MAY	DISC
	@	BEDFORD	EQ 5MG BASE/ML	N75249 001	JUN 23, 2000	SEP	DISC
	@	BEN VENUE	EQ 5MG BASE/ML	N75455 001	JUN 20, 2000	SEP	DISC

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

MINOCIN

AB	LEDERLE	EQ 75MG BASE	N50649 003	FEB 12, 2001	MAR	NEWA
AB	+	EQ 100MG BASE	N50649 002	MAY 31, 1990	MAR	CRLD
	MINOCYCLINE HCL					
AB	DANBURY PHARMA	EQ 100MG BASE	N63065 001	DEC 30, 1991	MAR	CRLD
AB	IMPAK LABS	EQ 75MG BASE	N65005 003	APR 18, 2001	APR	NEWA
	VECTRIN					
	@ MEDICIS	EQ 75MG BASE	N63067 002	SEP 15, 1999	MAY	DISC
	@	EQ 100MG BASE	N63067 001	JUL 31, 1990	MAY	DISC
	POWDER, EXTENDED RELEASE; DENTAL					
	ARESTIN					
+ +	ORAPHARMA	EQ 1MG BASE	N50781 001	FEB 16, 2001	FEB	NEWA

MIRTAZAPINE

TABLET, ORALLY DISINTEGRATING; ORAL

REMERON SOLTAB

+ +	ORGANON INC	15MG	N21208 001	JAN 12, 2001	JAN	NEWA
		30MG	N21208 002	JAN 12, 2001	JAN	NEWA
		45MG	N21208 003	JAN 12, 2001	JAN	NEWA

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

KADIAN

+ +	FAULDING PHARMS	20MG	N20616 001	JUL 03, 1996	JUN	CAHN
+ +		30MG	N20616 004	MAR 09, 2001	JUN	CAHN
+ +		50MG	N20616 002	JUL 03, 1996	JUN	CAHN
+ +		60MG	N20616 005	MAR 09, 2001	JUN	CAHN
+ +		100MG	N20616 003	JUL 03, 1996	JUN	CAHN

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

AB	WATSON LABS	100MG	N75656 001	JAN 30, 2001	JAN	NEWA
	ORAMORPH SR					
BC	ELAN PHARMS	15MG	N19977 004	NOV 23, 1994	SEP	CAHN
BC		30MG	N19977 001	AUG 15, 1991	SEP	CAHN
BC		60MG	N19977 002	AUG 15, 1991	SEP	CAHN
BC		100MG	N19977 003	AUG 15, 1991	SEP	CAHN

NABUMETONE

TABLET; ORAL

NABUMETONE

AB TEVA 750MG N75189 002 SEP 24, 2001 SEP NEWA

NADOLOL

TABLET; ORAL

CORGARD

AB APOTHECON 40MG N18063 001 DEC 10, 1979 AUG CRLD
NADOLOL
AB GENEVA PHARMS TECH 20MG N74501 001 NOV 09, 1995 JAN CAHN
AB 40MG N74501 002 NOV 09, 1995 JAN CAHN
AB 80MG N74501 003 NOV 09, 1995 JAN CAHNNAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

② APOTHECON

+ EQ 500MG BASE/VIAL N61984 001 APR 29, 1976 MAY DISC
② EQ 500MG BASE/VIAL N62527 001 AUG 02, 1984 MAY CRLD
② EQ 1GM BASE/VIAL N61984 002 APR 29, 1976 MAY DISC
② EQ 2GM BASE/VIAL N61984 003 APR 29, 1976 MAY DISC
② EQ 4GM BASE/VIAL N61984 005 APR 29, 1976 MAY DISC
② MARSAM EQ 500MG BASE/VIAL N62844 001 OCT 26, 1988 MAY DISC
② EQ 1GM BASE/VIAL N62844 002 OCT 26, 1988 MAY DISN
② EQ 1.5GM BASE/VIAL N62844 003 OCT 26, 1988 MAY DISC
② EQ 2GM BASE/VIAL N62844 004 OCT 26, 1988 MAY DISC
② EQ 4GM BASE/VIAL N62844 005 OCT 26, 1988 MAY DISC
② EQ 10GM BASE/VIAL N63008 001 SEP 29, 1988 MAY DISC

NALLPEN

② SMITHKLINE BEECHAM

② EQ 500MG BASE/VIAL N61999 001 JUL 10, 1978 MAY DISC
② EQ 1GM BASE/VIAL N61999 002 JUL 10, 1978 MAY DISC
② EQ 2GM BASE/VIAL N61999 003 JUL 10, 1978 MAY DISC
② EQ 10GM BASE/VIAL N61999 004 JUL 17, 1978 MAY DISC

UNIPEN

② WYETH AYERST

② EQ 500MG BASE/VIAL N50320 001 JUN 23, 1970 MAY DISC
② EQ 500MG BASE/VIAL N62717 001 DEC 16, 1986 MAY DISC
② EQ 1GM BASE/VIAL N62717 002 DEC 16, 1986 MAY DISC
② EQ 2GM BASE/VIAL N50320 003 JUN 23, 1970 MAY DISC
② EQ 2GM BASE/VIAL N62717 004 DEC 16, 1986 MAY DISC
② EQ 4GM BASE/VIAL N50320 004 JUN 23, 1970 MAY DISC
② EQ 10GM BASE/VIAL N50320 005 DEC 21, 1978 MAY DISC

UNIPEN IN PLASTIC CONTAINER

② WYETH AYERST

EQ 1GM BASE/VIAL N50320 002 JUN 23, 1970 MAY DISC

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HCL

② ASTRazeneca

10MG/ML N72070 001 APR 10, 1989 JUN DISC
② 20MG/ML N72073 001 APR 10, 1989 JUN DISC

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

@ WYETH AYERST	0.02MG/ML	N70188 001	SEP 24, 1986	JAN	DISC
@	0.02MG/ML	N70189 001	SEP 24, 1986	JAN	DISC
@	0.4MG/ML	N70190 001	SEP 24, 1986	JAN	DISC
@	0.4MG/ML	N70191 001	SEP 24, 1986	JAN	DISC

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

NALOXONE HCL AND PENTAZOCINE

AB AMIDE PHARM	EQ 0.5MG BASE;EQ 50MG BASE	N75735 001	JUL 11, 2001	JUL	NEWA
----------------	----------------------------	------------	--------------	-----	------

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALBALON

AT ALLERGAN	0.1%	N80248 001	MAR 24, 1972	JUL	CRLD
AT +	0.1%	N80248 001	MAR 24, 1972	AUG	CRLD
NAPHCON FORTE		N80229 001	MAR 06, 1974	JUL	DISC
@ ALCON	0.1%				
OPCON					
@ BAUSCH AND LOMB	0.1%	N87506 001	DEC 01, 1981	JUL	DISC
VASOCON					
AT NOVARTIS	0.1%	N80235 002	MAR 24, 1983	FEB	CAHN

NAPROXEN

TABLET, EXTENDED RELEASE; ORAL

NAPROXEN

AB + ALPHAPHARM	375MG	N75390 001	APR 19, 2001	APR	NEWA
AB +	500MG	N75390 002	APR 19, 2001	APR	NEWA

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

AB GENEVA PHARMS TECH	EQ 250MG BASE	N74495 001	DEC 05, 1994	JAN	CAHN
AB	EQ 500MG BASE	N74495 002	DEC 05, 1994	JAN	CAHN

NEDOCROMIL SODIUM

AEROSOL, METERED; INHALATION

TILADE

+ AVENTIS

1.75MG/INH

N19660 001 DEC 30, 1992 SEP CAHN

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

SERZONE

BRISTOL MYERS SQUIBB

50MG

N20152 001 DEC 22, 1994 APR CTEC

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATES

@ STERIS

EQ 40MG BASE/ML;200,000

UNITS/ML

N62664 001 APR 08, 1986 MAY DISC

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOSPORIN G.U. IRRIGANT

MONARCH PHARMS

EQ 40MG BASE/ML;200,000
UNITS/ML

N60707 001 JUN 28, 1966 MAY CTEC

SOLUTION/DROPS; OPHTHALMIC

STATROL

@ ALCON

EQ 3.5MG BASE/ML;16,250
UNITS/ML

N62339 001 NOV 30, 1984 JUL DISC

NESIRITIDE

FOR SOLUTION; INTRAVENOUS

NATRECOR

+ SCIOS

1.5MG/VIAL

N20920 001 AUG 10, 2001 AUG NEWA

NETILMICIN SULFATE

INJECTABLE; INJECTION

NETROMYCIN

@ SCHERING

EQ 100MG BASE/ML

N50544 003 FEB 28, 1983 MAY DISC

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

@ CHASE LABS NJ

10MG

N72409 001 JUL 04, 1990 FEB WDRP

@

20MG

N73421 001 JUN 19, 1991 FEB WDRP

TABLET, EXTENDED RELEASE; ORAL

ADALAT CC

AB1

BAYER

30MG

N20198 001 APR 21, 1993 APR CTEC

NIFEDIPINE

AB2

BIOVAIL

30MG

N75289 002 FEB 06, 2001 FEB NEWA

>A > AB1 ELAN PHARM

60MG

N75659 001 OCT 26, 2001 OCT NEWA

PROCARDIA XL

AB2 +

PFIZER

30MG

N19684 001 SEP 06, 1989 FEB CTEC

NITROFURAZONE

OINTMENT; TOPICAL

NITROFURAZONE

@ CLAY PARK

0.2%

N84968 001 JAN 25, 1978 MAY DISC

POWDER; TOPICAL

FURACIN

@ ROBERTS LABS

0.2%

N83791 001 OCT 17, 1975 FEB WDRP

SOLUTION; TOPICAL

NITROFURAZONE

@ CLAY PARK

0.2%

N85130 001 NOV 02, 1978 MAY DISC

+ WENDT

0.2%

N87081 001 JUL 22, 1981 MAY CTEC

NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

@ POHL BOSKAMP

0.4MG/SPRAY

N18705 001 OCT 31, 1985 APR DISC

NOREPINEPHRINE BITARTRATE; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAIN AND NOVOCAIN W/ LEVOPHED

@ EASTMAN KODAK

EQ 0.033MG BASE/ML;2%;0.4%

N08592 003 MAR 11, 1955 SEP WDAG

NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE

AB

BARR

5MG

N75951 001 MAY 25, 2001 MAY NEWA

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

PAMELOR

AB

TYCO HLTHCARE

EQ 10MG BASE

N18013 001 AUG 01, 1977 JUN CAHN

AB

EQ 25MG BASE

N18013 002 AUG 01, 1977 JUN CAHN

AB

EQ 50MG BASE

N18013 004 JUN 14, 1979 JUN CAHN

AB +

EQ 75MG BASE

N18013 003 JUN 14, 1979 JUN CAHN

SOLUTION; ORAL

AA

TYCO HLTHCARE

EQ 10MG BASE/5ML

N18012 001 AUG 01, 1977 JUN CAHN

NYSTATIN

CREAM; TOPICAL

NILSTAT

AB

LEDERLE

100,000 UNITS/GM

N61445 001 APR 02, 1971 MAY DISC

NYSTATIN

AB

TEVA

100,000 UNITS/GM

N61966 001 MAY 25, 1976 MAY DISC

OINTMENT; TOPICAL

NILSTAT

AB

LEDERLE

100,000 UNITS/GM

N61444 001 MAR 29, 1971 MAY DISC

NYSTATIN

AT +

ALTANA

100,000 UNITS/GM

N62124 002 SEP 23, 1982 MAY CTEC

SUSPENSION; ORAL

AB ROXANE

100,000 UNITS/ML

N62832 001 DEC 27, 1991 MAY DISC

AB TEVA

100,000 UNITS/ML

N62670 001 JUN 18, 1987 MAY DISC

AB

100,000 UNITS/ML

N62776 001 DEC 17, 1987 MAY DISC

AB THAMES

100,000 UNITS/ML

N62876 001 FEB 29, 1988 JUL DISC

TABLET; ORAL

AB EON

500,000 UNITS

N62065 001 JUL 22, 1977 MAY DISC

AB ROSEMONT

500,000 UNITS

N62524 001 NOV 26, 1985 MAY DISC

TABLET; VAGINAL

KOROSTATIN

AB

HOLLAND RANTOS

100,000 UNITS

N61718 001 SEP 30, 1974 FEB WDRP

NYSTATIN: TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

MYCO-TRIACET II

AB

TEVA

100,000 UNITS/GM;0.1%

N62045 002 NOV 26, 1985 MAY DISC

NYSTATIN AND TRIAMCINOLONE ACETONIDE

AB

CLAY PARK

100,000 UNITS/GM;0.1%

N62280 002 OCT 10, 1985 MAY DISC

OLANZAPINE

TABLET; ORAL

ZYPREXA

LILLY

+

15MG

20MG

N20592 005 SEP 09, 1997 JUN CRLD
N20592 006 SEP 09, 1997 JUN CMFDOXACILLIN SODIUM

INJECTABLE; INJECTION

BACTOCILL

@ SMITHKLINE BEECHAM

@

OXACILLIN SODIUM

EQ 1GM BASE/VIAL

EQ 2GM BASE/VIAL

N62736 001 DEC 19, 1986 FEB DISC
N62736 002 DEC 19, 1986 FEB DISC

AP + APOTHECON

EQ 1GM BASE/VIAL

N61490 003 APR 08, 1971 FEB CRLD

AP +

EQ 2GM BASE/VIAL

N62737 002 DEC 23, 1986 FEB CRLD

@ IBI

EQ 125MG BASE/VIAL

N62798 003 DEC 11, 1995 MAY DISC

@

EQ 250MG BASE/VIAL

N62798 004 DEC 11, 1995 MAY DISC

@

EQ 500MG BASE/VIAL

N62798 005 DEC 11, 1995 MAY DISC

@

EQ 1GM BASE/VIAL

N62798 001 DEC 11, 1995 MAY DISC

@

EQ 2GM BASE/VIAL

N62798 002 DEC 11, 1995 MAY DISC

OXAPROZIN

TABLET; ORAL

DAYPRO

AB + SEARLE

600MG

N18841 004 OCT 29, 1992 JAN CFTG

OXAPROZIN

AB DR REDDYS LABS LTD

600MG

N75855 001 JAN 31, 2001 JAN NEWA

AB EON

600MG

N75845 001 JAN 31, 2001 JAN NEWA

AB GENEVA PHARMS

600MG

N75850 001 APR 27, 2001 APR NEWA

AB GENPHARM

600MG

N75847 001 FEB 28, 2001 FEB NEWA

AB INVAMED

600MG

N75842 001 APR 12, 2001 APR NEWA

AB MYLAN

600MG

N75851 001 AUG 17, 2001 AUG NEWA

>A> AB PUREPAC PHARM

600MG

N75843 001 OCT 03, 2001 OCT NEWA

AB WATSON LABS

600MG

N75848 001 FEB 09, 2001 FEB NEWA

OXAZEPAM

CAPSULE; ORAL

SERAX

AB FAULDING PHARMS

10MG

N15539 002 SEP 29, 1966 JUN CAHN

AB

15MG

N15539 004 SEP 29, 1966 JUN CAHN

AB +

30MG

N15539 006 SEP 29, 1966 JUN CAHN

TABLET; ORAL

+ FAULDING PHARMS

15MG

N15539 008 NOV 16, 1967 JUN CAHN

OXCARBAZEPINE

SUSPENSION; ORAL

TRILEPTAL

+ NOVARTIS

300MG/5ML

N21285 001 MAY 25, 2001 MAY NEWA

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

ROXICODONE

ELAN PHARMS

15MG

N21011 001 AUG 31, 2000 SEP CAHN

+	30MG	N21011 002	AUG 31, 2000	SEP CAHN
TABLET, EXTENDED RELEASE; ORAL				
OXYCONTIN				
© PURDUE PHARMA LP	160MG	N20553 005	MAR 15, 2000	JUN DISC

OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL					
OXYTETRACYCLINE HCL					
© IMPAX LABS	EQ 250MG BASE	N60760 001	AUG 09, 1967	FEB DISC	
© PROTER	EQ 250MG BASE	N60869 001	JAN 29, 1964	FEB WDRP	
© WEST WARD	EQ 250MG BASE	N60770 001	SEP 29, 1967	MAY DISC	
TERRAMYCIN					
+	PFIZER	EQ 250MG BASE	N50286 002	SEP 08, 1964	MAY CTEC

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL

AP	BEDFORD	6MG/ML	N75190 001	JUL 27, 2001	JUL NEWA
AP	MYLAN	6MG/ML	N75278 001	JUL 23, 2001	JUL NEWA
AP	ZENITH GOLDLINE	6MG/ML	N75297 001	MAR 27, 2001	MAR NEWA

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

AREDIA

AP	+	NOVARTIS	30MG/VIAL	N20036 001	OCT 31, 1991	APR CFTG
AP	+		90MG/VIAL	N20036 004	MAY 06, 1993	APR CFTG
PAMIDRONATE DISODIUM						
AP		BEDFORD	30MG/VIAL	N75290 001	APR 30, 2001	APR NEWA
AP			90MG/VIAL	N75290 003	APR 30, 2001	APR NEWA

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE

©	ASTRAZENECA	1MG/ML	N72210 001	MAR 31, 1988	JUL DISC
©		2MG/ML	N72211 001	MAR 31, 1988	JUL DISC
©		2MG/ML	N72213 001	MAR 31, 1988	JUL DISC

PANTOPRAZOLE SODIUM

INJECTABLE; IV (INFUSION)

PROTONIX IV

+	WYETH AYERST	EQ 40MG BASE/VIAL	N20988 001	MAR 22, 2001	MAR NEWA
TABLET, DELAYED RELEASE; ORAL					
PROTONIX					
+	WYETH AYERST	EQ 20MG BASE	N20987 002	JUN 12, 2001	JUN NEWA

PEMOLINE

TABLET; ORAL

PEMOLINE

AB	MALLINCKRODT	18.75MG	N75726 003	MAR 30, 2001	MAR NEWA
AB		37.5MG	N75726 002	MAR 30, 2001	MAR NEWA
AB		75MG	N75726 001	MAR 30, 2001	MAR NEWA
AB	WATSON LABS	18.75MG	N75287 001	JUN 13, 2001	JUN NEWA

PENCICLOVIR SODIUM

CREAM; TOPICAL

DENAVIR

+ NOVARTIS

1%

N20629 001 SEP 24, 1996 JUL CAHN

PENICILLIN G POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN

@ TEVA

200,000 UNITS/5ML

N60307 002 MAY 27, 1964 JUL DISC

@

400,000 UNITS/5ML

N60307 004 MAY 27, 1964 JUL DISC

PENICILLIN-2

@ TEVA

250,000 UNITS/5ML

N60307 003 MAY 27, 1964 JUL DISC

TABLET; ORAL

PENICILLIN G POTASSIUM

@ TEVA

200,000 UNITS

N60306 001 JUN 01, 1964 MAY DISC

@

250,000 UNITS

N60306 002 JUN 01, 1964 MAY DISC

@

400,000 UNITS

N60306 003 JUN 01, 1964 MAY DISC

@

500,000 UNITS

N60306 004 JUN 26, 1979 MAY DISC

PENICILLIN G PROCAINE

INJECTABLE; INJECTION

PENICILLIN G PROCAINE

@ PFIZER

300,000 UNITS/VIAL

N60099 001 NOV 10, 1948 MAY DISC

@

1,500,000 UNITS/VIAL

N60099 002 NOV 10, 1948 MAY DISC

PFIZERPEN-AS

@ PFIZER

300,000 UNITS/ML

N60286 001 NOV 01, 1950 MAY DISC

@

600,000 UNITS/ML

N60286 002 NOV 01, 1950 MAY DISC

WYCILLIN

+ KING PHARMS

300,000 UNITS/ML

N60101 002 APR 26, 1948 MAY CTEC

+

600,000 UNITS/ML

N60101 001 APR 26, 1948 MAY CTEC

PENICILLIN G SODIUM

INJECTABLE; IM-IV

PENICILLIN G SODIUM

+ BIOCHEMIE

5,000,000 UNITS/VIAL

N65068 001 FEB 26, 2001 FEB NEWA

@ MARSAM

5,000,000 UNITS/VIAL

N63014 001 SEP 13, 1988 FEB DISC

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN V POTASSIUM

@ MYLAN

EQ 125MG BASE/5ML

N61624 002 AUG 07, 1972 MAY DISC

@

EQ 250MG BASE/5ML

N61624 001 JUN 05, 1972 MAY DISC

V-CILLIN K

@ LILLY

EQ 125MG BASE/5ML

N60004 001 AUG 21, 1958 MAY DISC

@

EQ 250MG BASE/5ML

N60004 002 APR 07, 1967 MAY DISC

TABLET; ORAL

PEN-VEE K

AB + WYETH AYERST

EQ 500MG BASE

N60006 003 JAN 13, 1958 MAY CRLD

PENICILLIN V POTASSIUM

AB + BIOCHEMIE

EQ 500MG BASE

N64071 002 NOV 30, 1995 MAY CTEC

@ MYLAN

EQ 250MG BASE

N61530 001 NOV 18, 1971 MAY DISC

@

EQ 500MG BASE

N61530 002 MAR 20, 1972 MAY DISC

PENICILLIN V POTASSIUM

TABLET; ORAL				
V-CILLIN K				
@ LILLY	EQ 125MG BASE	N60003 001	SEP 17, 1957	MAY DISC
@	EQ 250MG BASE	N60003 002	SEP 17, 1957	MAY DISC
@	EQ 500MG BASE	N60003 003	SEP 17, 1957	MAY DISC

PENTOBARBITAL

ELIXIR; ORAL				
NEMBUTAL				
@ ABBOTT	18.2MG/5ML	N83244 001	JAN 08, 1975	JUL DISC

PENTOBARBITAL SODIUM

CAPSULE; ORAL				
NEMBUTAL SODIUM				
@ ABBOTT	50MG	N84093 001	JAN 14, 1975	JUL DISC
SUPPOSITORY; RECTAL				
NEMBUTAL				
@ ABBOTT	30MG	N83247 001	JAN 25, 1982	JUL DISC
@	60MG	N83247 002	JAN 25, 1982	JUL DISC
@	120MG	N83247 003	JAN 25, 1982	JUL DISC
@	200MG	N83247 004	JAN 25, 1982	JUL DISC

PERFLUTREN

INJECTABLE; INTRAVENOUS				
DEFINITY				
+ DUPONT PHARMS	6.52MG/ML	N21064 001	JUL 31, 2001	JUL NEWA

PERPHENAZINE

CONCENTRATE; ORAL				
PERPHENAZINE				
+ PHARM ASSOC	16MG/5ML	N40360 001	MAY 25, 2001	MAY NEWA
TRILAFON				
@ SCHERING	16MG/5ML	N11557 001	DEC 12, 1958	MAR DISC

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL				
SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PENAZOPYRIDINE HCL				
+ ABLE	200MG;800MG;160MG	N21105 001	JUN 26, 2001	JUN NEWA

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL				
PHENDIMETRAZINE TARTRATE				
@ EON	35MG	N85633 001	JUL 13, 1978	JUL DISC
@	35MG	N85694 001	JUN 05, 1978	MAY DISC
AA +	35MG	N85695 001	JUN 05, 1978	JUL CRLD
@	35MG	N85702 001	JUN 07, 1978	JUL DISC
CAPSULE, EXTENDED RELEASE; ORAL				
BC + EON	105MG	N18074 001	APR 16, 1979	JUL CRLD
@ GENEVA PHARMS	105MG	N87378 001	NOV 03, 1981	JUL DISC
TABLET; ORAL				
PHENAZINE-35				
@ ABC HOLDING	35MG	N85512 001	MAY 06, 1977	MAY DISC

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

PHENDIMETRAZINE TARTRATE

© EON	35MG	N85402 001	MAY 19, 1978	MAY	DISC
©	35MG	N85497 001	AUG 19, 1977	MAY	DISC
© MIKART	35MG	N89452 001	OCT 30, 1991	JUL	DISC
© ROSEMONT	35MG	N84399 001	MAY 28, 1981	MAY	DISC
STATOBEK					
© TEVA	35MG	N86013 001	DEC 16, 1977	JUL	DISC
X-TROZINE					
© SHIRE RICHWOOD	35MG	N86550 001	SEP 16, 1981	JUL	DISC
©	35MG	N86551 001	SEP 16, 1981	JUL	DISC
©	35MG	N86552 001	SEP 16, 1981	JUL	DISC

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

OBY-TRIM

© SHIRE RICHWOOD

30MG

N87764 001 MAR 18, 1982 JUL DISC

PHENTERMINE HCL

© ABC HOLDING

30MG

N85411 001 SEP 10, 1980 MAY DISC

AA ABLE

30MG

N40403 001 AUG 30, 2001 AUG NEWA

AA

30MG

N40427 001 AUG 30, 2001 AUG NEWA

© ROSEMONT

30MG

N84487 001 APR 09, 1982 MAY DISC

TABLET; ORAL

AA ABLE

37.5MG

N40402 001 AUG 30, 2001 AUG NEWA

+ EON

30MG

N88605 001 SEP 28, 1987 MAY CMFD

PHENYTOIN

SUSPENSION; ORAL

PHENYTOIN

AB UDL

125MG/5ML

N40342 001 JAN 31, 2001 JAN NEWA

PIPERAZINE CITRATE

SYRUP; ORAL

PIPERAZINE CITRATE

© LANNETT

EQ 500MG BASE/5ML

N80963 001 JUL 25, 1974 MAY DISC

TABLET; ORAL

© IMPAX LABS

EQ 250MG BASE

N80874 001 JUL 19, 1973 MAY DISC

PIPOBROMAN

TABLET; ORAL

VERCYTE

© ABBOTT

25MG

N16245 002 JUL 01, 1966 JUL DISC

PORACTANT ALFA

SUSPENSION; INTRATRACHEAL

CUROSURF

+ DEY

80MG/ML

N20744 001 NOV 18, 1999 SEP CAIN

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE; ORAL

K-DUR 10

AB KEY PHARMS

10MEQ

N19439 002 JUN 13, 1986 APR CTEC

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HCL

@ PUREPAC PHARM

EQ 1MG BASE

N72991 001 MAY 16, 1989 JUL DISC

@

EQ 2MG BASE

N72921 001 MAY 16, 1989 JUL DISC

@

EQ 5MG BASE

N72992 001 MAY 16, 1989 JUL DISC

PREDNICARBATE

OINTMENT; TOPICAL

DERMATOP

+ AVENTIS PHARMS

0.1%

N19568 001 SEP 23, 1991 MAR CMFD

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

>A>	AA	KV PHARM	5MG/5ML	N40423 001	OCT 22, 2001	OCT	NEWA
		PRELONE					
>D>		+ MURO	5MG/5ML	N89654 001	JAN 17, 1989	OCT	CFTG
>A>	AA	+	5MG/5ML	N89654 001	JAN 17, 1989	OCT	CFTG

TABLET; ORAL

PREDNISOLONE

@ CHELSEA LABS

5MG

N85085 002 FEB 23, 1977 MAY DISC

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

VASOCIDIN

AT		NOVARTIS	0.5%;10%	N88791 001	OCT 05, 1984	FEB	CAHN
		SUSPENSION/DROPS; OPHTHALMIC					
		METIMYD					
	+	SCHERING	0.5%;10%	N10210 001	FEB 24, 1956	FEB	CTEC
		PREDAMIDE					
	@	AKORN	0.5%;10%	N88059 001	JUL 29, 1983	FEB	WDRP
		SULPHRIN					
	@	BAUSCH AND LOMB	0.5%;10%	N88089 001	DEC 28, 1982	FEB	WDRP

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION/DROPS; OPHTHALMIC

INFLAMASE FORTE

AT	+	NOVARTIS	EQ 0.9% PHOSPHATE	N80751 002	DEC 19, 1973	FEB	CAHN
		INFLAMASE MILD					
AT	+	NOVARTIS	EQ 0.11% PHOSPHATE	N80751 001	DEC 19, 1973	FEB	CAHN
		PREDNISOLONE SODIUM PHOSPHATE					
	@	AKORN	EQ 0.11% PHOSPHATE	N83358 001	AUG 21, 1974	FEB	WDRP
	@		EQ 0.9% PHOSPHATE	N83358 002	AUG 21, 1974	FEB	WDRP
	@	ALCON UNIVERSAL	EQ 0.11% PHOSPHATE	N81043 001	OCT 24, 1991	MAY	DISC
	@		EQ 0.9% PHOSPHATE	N81044 001	OCT 24, 1991	MAY	DISC

PREDNISONE

TABLET; ORAL

PREDNISONE

@ CHELSEA LABS

5MG

N85084 002 DEC 15, 1981 MAY DISC

@ GENEVA PHARMS

5MG

N80336 002 JUL 29, 1976 MAY DISC

	@ HALSEY	10MG	N86595 001	APR 10, 1979	JUL	DISC
	@ LANNETT	20MG	N84275 001	JUN 27, 1974	MAY	DISC
AB	TRIGEN	5MG	N40362 002	AUG 29, 2001	AUG	NEWA
AB		10MG	N40362 001	AUG 29, 2001	AUG	NEWA
PRILOCAINE HYDROCHLORIDE						
INJECTABLE; INJECTION						
CITANESE PLAIN						
	@ ASTRAZENECA	4%	N14763 007	NOV 18, 1965	SEP	DISC
+ DENTSPLY PHARM		4%	N21382 001	NOV 18, 1965	SEP	NEWA
PRIMIDONE						
SUSPENSION; ORAL						
MYSOLINE						
+ XCEL PHARMS		250MG/5ML	N10401 001	JUL 05, 1956	JUL	CAHN
TABLET; ORAL						
AB	ELAN PHARMA	50MG	N09170 003	MAR 08, 1954	MAY	CFTG
AB	XCEL PHARMS	50MG	N09170 003	MAR 08, 1954	JUL	CAHN
AB +		250MG	N09170 002	MAR 08, 1954	JUL	CAHN
PRIMIDONE						
AB	LANNETT	50MG	N84903 002	MAY 24, 2001	MAY	NEWA
PROCHLORPERAZINE						
SUPPOSITORY; RECTAL						
COMPAZINE						
AB	SMITHKLINE BEECHAM	2.5MG	N11127 003	FEB 09, 1959	JUL	CFTG
AB		5MG	N11127 001	SEP 27, 1957	JUL	CFTG
PROCHLORPERAZINE						
AB	ABLE	2.5MG	N40407 001	JUL 11, 2001	JUL	NEWA
AB		5MG	N40407 002	JUL 11, 2001	JUL	NEWA
AB		25MG	N40407 003	JUL 11, 2001	JUL	NEWA
PROCHLORPERAZINE EDISYLATE						
INJECTABLE; INJECTION						
PROCHLORPERAZINE EDISYLATE						
@ WYETH AYERST		EQ 5MG BASE/ML	N86348 001	JUL 05, 1979	JUL	DISC
PROCHLORPERAZINE MALEATE						
TABLET; ORAL						
PROCHLORPERAZINE MALEATE						
AB	GENEVA PHARMS TECH	EQ 5MG BASE	N40101 001	JUL 19, 1996	JAN	CAHN
AB		EQ 10MG BASE	N40101 002	JUL 19, 1996	JAN	CAHN
AB		EQ 25MG BASE	N40101 003	JUL 19, 1996	JAN	CAHN
PROGESTERONE						
INJECTABLE; INJECTION						
PROGESTERONE						
AO	AM PHARM PARTNERS	50MG/ML	N75906 001	APR 25, 2001	APR	NEWA
>A>	AO + SCHEIN	50MG/ML	N17362 002	MAY 08, 1978	OCT	CAHN
>D>	AO + STERIS	50MG/ML	N17362 002	MAY 08, 1978	OCT	CAHN
	AO +	50MG/ML	N17362 002	MAY 08, 1978	APR	CFTG

PROMETHAZINE HYDROCHLORIDE

SUPPOSITORY; RECTAL

PROMETHACON	50MG	N84902 001	OCT 05, 1981	MAY	DISC
@ POLYMEDICA					
TABLET; ORAL					
PHENERGAN					
WYETH AYERST	12.5MG	N07935 002	MAR 29, 1951	MAY	CTEC
PROMETHAZINE HCL					
@ LANNETT	12.5MG	N80949 001	JUL 28, 1976	MAY	DISC
@	25MG	N80949 002	JUN 28, 1976	MAY	DISC
@	50MG	N80949 003	JUN 28, 1976	MAY	DISC
@ PVT FORM	25MG	N83658 001	OCT 01, 1976	MAY	DISC

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

PROPOXYPHENE HCL					
@ GENEVA PHARMS	65MG	N83125 002	APR 14, 1976	MAY	DISC
@ IMPAX LABS	65MG	N83317 001	OCT 23, 1973	MAY	DISC

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERAL LA

WYETH AYERST LABS	60MG	N18553 004	MAR 18, 1987	AUG	CRLD
	80MG	N18553 002	APR 19, 1983	AUG	CRLD
	120MG	N18553 003	APR 19, 1983	AUG	CRLD

TABLET; ORAL

PROPRANOLOL HCL

@ LEDERLE	10MG	N70125 001	JUL 30, 1985	MAY	DISC
>D> AB	20MG	N70126 001	JUL 30, 1985	OCT	DISC
>A>	20MG	N70126 001	JUL 30, 1985	OCT	DISC
@	40MG	N70127 001	JUL 30, 1985	SEP	DISC
@ WATSON LABS	20MG	N70549 001	APR 11, 1986	MAY	DISC

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

PROTRIPTYLINE HCL

AB	ODYSSEY PHARMS	5MG	N73644 001	AUG 24, 1995	JAN	CAHN
AB		10MG	N73645 001	AUG 24, 1995	JAN	CAHN
	VIVACTIL					
AB	ODYSSEY PHARMS	5MG	N73644 001	AUG 24, 1995	MAR	CTNA
AB	+	10MG	N73645 001	AUG 24, 1995	MAR	CTNA
@ SIDMAK LABS		5MG	N16012 001	SEP 27, 1967	MAR	DISC
@		10MG	N16012 002	SEP 27, 1967	MAR	DISC

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

TRILITRON

@ NEWTRON PHARMS	30MG/5ML;1.25MG/5ML	N88474 001	FEB 12, 1985	FEB	WDRP
------------------	---------------------	------------	--------------	-----	------

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE; ORAL				
QUINAGLUTE				
BX + BERLEX LABS	324MG		N16647 001	DEC 08, 1969 MAR CTEC
QUINIDINE GLUCONATE				
BX DANBURY PHARMA	324MG		N87810 001	SEP 29, 1982 MAR CTEC
@ GENEVA PHARMS	324MG		N89894 001	DEC 15, 1988 MAR DISC
BX MUTUAL PHARM	324MG		N89338 001	FEB 11, 1987 MAR CTEC

QUINIDINE SULFATE

TABLET; ORAL				
QUINIDINE SULFATE				
@ IMPAX LABS	200MG		N83347 001	DEC 08, 1976 FEB DISC
@ MUTUAL PHARM	300MG		N81031 001	APR 14, 1989 MAY DISC
@ WEST WARD	200MG		N83862 001	SEP 02, 1976 MAY DISC

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL				
ZANTAC 150				
@ GLAXO WELLCOME	EQ 150MG BASE		N20095 001	MAR 08, 1994 AUG DISC
TABLET; ORAL				
RANITIDINE HCL				
@ BOEHRINGER INGELHEIM	EQ 150MG BASE		N74662 001	AUG 29, 1997 SEP DISC
@	EQ 300MG BASE		N74662 002	AUG 29, 1997 SEP DISC

RIBAVIRIN

CAPSULE; ORAL				
REBETOL				
+ SCHERING PLOUGH RES	200MG		N20903 002	JUL 25, 2001 AUG NEWA

RIFAMPIN

CAPSULE; ORAL				
RIFAMPIN				
AB VERSAPHARM	150MG		N65028 001	MAR 14, 2001 MAR NEWA
AB	300MG		N65028 002	MAR 14, 2001 MAR NEWA

RISPERIDONE

TABLET; ORAL				
RISPERDAL				
JANSSEN	0.5MG		N20272 007	JAN 27, 1999 APR CRLD
+	1MG		N20272 001	DEC 29, 1993 APR CRLD
	4MG		N20272 004	DEC 29, 1993 APR CRLD

SECOBARBITAL SODIUM

CAPSULE; ORAL				
SECOBARBITAL SODIUM				
@ ICN	100MG		N85477 001	DEC 10, 1981 FEB WDRP

SECRETIN

INJECTABLE; INJECTION				
SECRETIN-FERRING				
@ FERRING	75CU/VIAL		N18290 001	MAY 29, 1981 JUN DISC

SILVER SULFADIAZINE

DRESSING; TOPICAL

SILDAFLO

@ QUESTCOR PHARMS

1%

N19608 001 NOV 30, 1989 MAY CTNA

SIMVASTATIN

TABLET; ORAL

ZOCOR

MERCK

5MG

N19766 001 DEC 23, 1991 APR CTEC

SODIUM IODIDE, I-131

CAPSULE; ORAL

SODIUM IODIDE I 131

>D> + CIS 100uCi
 >A> @ 100uCi

N17316 002 NOV 10, 1976 OCT DISC
 N17316 002 NOV 10, 1976 OCT DISC

SODIUM POLYSTYRENE SULFONATE

SUSPENSION; ORAL, RECTAL

SPS

AA + CAROLINA MEDCL 15GM/60ML

N87859 001 DEC 08, 1982 MAY CRLD

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL

@ ELKINS SINK

1%

N05970 004 AUG 13, 1946 JUL DISC

@

3%

N05970 005 AUG 13, 1946 JUL DISC

SOMATROPIN RECOMBINANT

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

SAIZEN

+ SERONO

8.8MG/VIAL

N19764 003 AUG 29, 2000 AUG NEWA

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SORINE

AB UPSHER SMITH 80MG

N75500 001 APR 27, 2001 APR NEWA

AB 120MG

N75500 004 APR 27, 2001 APR NEWA

AB 160MG

N75500 002 APR 27, 2001 APR NEWA

AB 240MG

N75500 003 APR 27, 2001 APR NEWA

SOTALOL HCL

>A> AB MUTUAL PHARM 80MG

N75515 001 OCT 15, 2001 OCT NEWA

>A> AB 120MG

N75515 004 OCT 15, 2001 OCT NEWA

>A> AB 160MG

N75515 002 OCT 15, 2001 OCT NEWA

>A> AB 240MG

N75515 003 OCT 15, 2001 OCT NEWA

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

AB MYLAN 25MG

N40424 001 AUG 20, 2001 AUG NEWA

AB 50MG

N40424 002 AUG 20, 2001 AUG NEWA

AB 100MG

N40424 003 AUG 20, 2001 AUG NEWA

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

@ PFIZER

EQ 1GM BASE/VIAL

N60076 001 FEB 18, 1946 MAY DISC

@

EQ 5GM BASE/VIAL

N60076 002 FEB 18, 1946 MAY DISC

+ PHARMA TEK

EQ 1GM BASE/VIAL

N64210 001 JUN 30, 1998 MAY CTEC

SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPH-10

@ ALLERGAN

10%

N84015 001 JAN 07, 1975 MAY DISC

CETAMIDE

AT + ALCON

10%

N80021 001 SEP 27, 1972 MAY CTEC

SODIUM SULAMYD

@ SCHERING

10%

N05963 002 NOV 26, 1947 MAY DISC

SOLUTION/DROPS; OPHTHALMIC

BLEPH-10

AT + ALLERGAN

10%

N80028 001 MAY 25, 1971 MAY CRLD

BLEPH-30

AT + ALLERGAN

30%

N80028 002 MAY 25, 1971 MAY CRLD

SODIUM SULAMYD

@ SCHERING

10%

N05963 001 AUG 01, 1946 MAY DISC

@

30%

N05963 003 NOV 26, 1947 MAY DISC

SULF-10

@ NOVARTIS

10%

N80025 001 JUN 03, 1971 FEB CAHN

AT

10%

N80025 001 JUN 03, 1971 SEP CMFD

SULF-15

AT NOVARTIS

15%

N89047 001 OCT 31, 1995 FEB CAHN

SULTEN-10

@ BAUSCH AND LOMB

10%

N87818 001 FEB 03, 1983 FEB WDRP

SULFAMETHOXAZOLE

TABLET; ORAL

GANTANOL

+ ROCHE

500MG

N12715 002 NOV 17, 1961 MAY CTEC

SULFAMETHOXAZOLE

@ GENEVA PHARMS

500MG

N85844 001 MAR 23, 1978 MAY DISC

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

BACTRIM

>D> AP + ROCHE

80MG/ML;16MG/ML

N18374 001 JUN 23, 1981 OCT CAHN

>A> AP + WOMEN FIRST HLTHCARE

80MG/ML;16MG/ML

N18374 001 JUN 23, 1981 OCT CAHN

SUSPENSION; ORAL

>D> @ ROCHE

200MG/5ML;40MG/5ML

N17560 001 APR 16, 1975 OCT CAHN

>A> @ WOMEN FIRST HLTHCARE

200MG/5ML;40MG/5ML

N17560 001 APR 16, 1975 OCT CAHN

BACTRIM PEDIATRIC

>D> AB + ROCHE

200MG/5ML;40MG/5ML

N17560 002 DEC 10, 1979 OCT CAHN

>A> AB + WOMEN FIRST HLTHCARE

200MG/5ML;40MG/5ML

N17560 002 DEC 10, 1979 OCT CAHN

TRIMETH/SULFA

@ NASKA

200MG/5ML;40MG/5ML

N72399 001 MAY 23, 1988 FEB WDRP

TABLET; ORAL

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL			
BACTRIM			
>D>	AB	ROCHE	400MG;80MG
>A>	AB	WOMEN FIRST HLTHCARE	400MG;80MG
BACTRIM DS			
>D>	AB	+ ROCHE	800MG;160MG
>A>	AB	+ WOMEN FIRST HLTHCARE	800MG;160MG
SULFAMETHOXAZOLE AND TRIMETHOPRIM			
>D>	AB	ROXANE	400MG;80MG
>A>	@		400MG;80MG
	@ TEVA		400MG;80MG
	@		800MG;160MG
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH			
>D>	AB	PLANTEX	800MG;160MG
	@ ROXANE		800MG;160MG
>A>	AB	TEVA	800MG;160MG

N17377 001	JUL 30, 1973	OCT	CAHN
N17377 001	JUL 30, 1973	OCT	CAHN

N17377 002	MAR 01, 1978	OCT	CAHN
N17377 002	MAR 01, 1978	OCT	CAHN

N72768 001	AUG 30, 1991	OCT	DISC
N72768 001	AUG 30, 1991	OCT	DISC

N18242 001	MAY 19, 1981	MAY	DISC
N18242 002	MAY 19, 1981	MAY	DISC

N70037 001	JUN 02, 1987	OCT	CAHN
N72769 001	AUG 30, 1991	SEP	DISC

N70037 001	JUN 02, 1987	OCT	CAHN
------------	--------------	-----	------

SULFANILAMIDE

CREAM; VAGINAL			
AVC			
AT	+	NOVAVAX	15%
SUPPOSITORY; VAGINAL			
+ NOVAVAX 1.05GM			

N06530 003	JAN 27, 1987	JAN	CAHN
------------	--------------	-----	------

N06530 004	JAN 27, 1987	JAN	CAHN
------------	--------------	-----	------

SULFISOXAZOLE

TABLET; ORAL			
SULFISOXAZOLE			
@ GENEVA PHARMS 500MG			

N85628 001	JUN 13, 1977	JUL	DISC
------------	--------------	-----	------

TECHNETIUM TC-99M APCITIDE

INJECTABLE; INJECTION			
ACUTECT			
BERLEX LABS N/A			
DIATIDE RES LABS N/A			
N/A N/A			

N20887 001	SEP 14, 1998	MAY	CAHN
------------	--------------	-----	------

N20887 001	SEP 14, 1998	APR	CAHN
------------	--------------	-----	------

N20887 001	SEP 14, 1998	JUL	CAHN
------------	--------------	-----	------

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION			
MPI DTPA KIT - CHELATE			
@ NYCOMED AMERSHAM N/A			

N17255 001	APR 15, 1976	SEP	WDAG
------------	--------------	-----	------

TEMAZEPAM

CAPSULE; ORAL			
RESTORIL			
TYCO HLTHCARE 7.5MG			
AB		15MG	
AB	+	30MG	

N18163 003	OCT 25, 1991	JUN	CAHN
------------	--------------	-----	------

N18163 001	FEB 27, 1981	JUN	CAHN
------------	--------------	-----	------

N18163 002	FEB 27, 1981	JUN	CAHN
------------	--------------	-----	------

>A> TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL			
VIREAD			
>A>	+	GILEAD	300MG

N21356 001	OCT 26, 2001	OCT	NEWA
------------	--------------	-----	------

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL					
TERAZOSIN HCL					
AB TORPHARM	EQ 1MG BASE	N75498 001	APR 12, 2001	APR	NEWA
AB	EQ 2MG BASE	N75498 002	APR 12, 2001	APR	NEWA
AB	EQ 5MG BASE	N75498 003	APR 12, 2001	APR	NEWA
AB	EQ 10MG BASE	N75498 004	APR 12, 2001	APR	NEWA
AB ZENITH GOLDLINE	EQ 1MG BASE	N75614 002	JAN 30, 2001	JAN	NEWA
AB	EQ 2MG BASE	N75614 001	JAN 30, 2001	JAN	NEWA
AB	EQ 5MG BASE	N75614 003	JAN 30, 2001	JAN	NEWA
AB	EQ 10MG BASE	N75614 004	JAN 30, 2001	JAN	NEWA

TERBUTALINE SULFATE

TABLET; ORAL					
BRETHINE					
AB NOVARTIS	2.5MG	N17849 001	MAY 17, 1976	JUN	CFTG
AB +	5MG	N17849 002	MAY 17, 1976	JUN	CFTG
TERBUTALINE SULFATE					
AB IMPAX LABS	2.5MG	N75877 001	JUN 26, 2001	JUN	NEWA
AB	5MG	N75877 002	JUN 26, 2001	JUN	NEWA

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL					
PANMYCIN					
⑧ PHARMACIA AND UPJOHN	250MG	N60347 001	SEP 28, 1954	MAY	DISC
ROBITET					
⑧ WYETH AYERST	250MG	N61734 001	JUN 06, 1973	MAY	DISC
⑧	500MG	N61734 002	JUN 06, 1973	MAY	DISC
TETRACYCLINE HCL					
⑧ DANBURY PHARMA	250MG	N62343 001	OCT 02, 1981	MAY	DISC
⑧	500MG	N62343 002	OCT 02, 1981	MAY	DISC
⑧ EON	250MG	N61471 001	OCT 28, 1971	MAY	DISC
⑧ WEST WARD	250MG	N60768 001	AUG 24, 1964	MAY	DISC
⑧	500MG	N60768 002	NOV 07, 1977	MAY	DISC
⑧ WYETH AYERST	250MG	N61685 001	DEC 11, 1972	JUL	DISC
⑧	500MG	N61685 002	DEC 11, 1972	JUL	DISC

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL					
THIORIDAZINE HCL					
⑧ CHELSEA LABS	10MG	N88561 001	MAY 11, 1984	JUL	DISC
⑧ TEVA	10MG	N88493 001	MAY 17, 1985	JUL	DISC
⑧ ZENITH GOLDLINE	50MG	N88194 001	APR 14, 1983	JUL	DISC

THIOTEP A

INJECTABLE; INJECTION					
THIOPLEX					
AP + IMMUNEX	15MG/VIAL	N20058 001	DEC 22, 1994	APR	CFTG
THIOTEP A					
AP AAIPHARMA	15MG/VIAL	N75698 001	SEP 20, 2001	SEP	NEWA
AP BEDFORD	15MG/VIAL	N75547 001	APR 02, 2001	APR	NEWA
AP GENESIA SICOR PHARMS	15MG/VIAL	N75730 001	APR 20, 2001	APR	NEWA

+ @ IMMUNEX	30MG/VIAL 15MG/VIAL	N75730 002 APR 20, 2001 APR NEWA N11683 001 FEB 19, 1959 APR DISC
----------------	------------------------	--

THYROGLOBULIN

TABLET; ORAL THYROGLOBULIN @ IMPAX LABS	64.8MG	N80151 001 AUG 07, 1973 FEB DISC
---	--------	----------------------------------

TICARCILLIN DISODIUM

INJECTABLE; INJECTION TICAR @ SMITHKLINE BEECHAM	EQ 3GM BASE/VIAL	N62690 001 DEC 19, 1986 MAY DISC
--	------------------	----------------------------------

TOBRAMYCIN

SOLUTION; INHALATION TOBI + CHIRON	300MG/5ML	N50753 001 DEC 22, 1997 SEP CAHN
SOLUTION/DROPS; OPHTHALMIC TOBRAMYCIN @ ALCON UNIVERSAL	0.3%	N63176 001 MAY 25, 1994 MAY DISC
AT ALTANA	0.3%	N65026 001 SEP 11, 2001 SEP NEWA

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION NEBCIN AP + LILLY	EQ 1.2GM BASE/VIAL	N50519 001 JUN 11, 1979 AUG CFTG
TOBRAMYCIN AP PHARMA TEK	EQ 1.2GM BASE/VIAL	N65013 001 AUG 17, 2001 AUG NEWA
TOBRAMYCIN SULFATE @ ASTRazeneca	EQ 10MG BASE/ML	N63119 001 OCT 31, 1994 AUG DISC
@ ELKINS SINK	EQ 40MG BASE/ML	N63121 001 OCT 31, 1994 MAY DISC
@ LEDERLE	EQ 10MG BASE/ML	N63128 001 NOV 27, 1991 MAY DISC
	EQ 40MG BASE/ML	N63127 001 NOV 27, 1991 MAY DISC
	EQ 10MG BASE/ML	N63113 001 APR 26, 1991 MAY DISC

TOLMETIN SODIUM

CAPSULE; ORAL TOLMETIN SODIUM @ GENEVA PHARMS	EQ 400MG BASE	N73462 001 APR 30, 1992 JUL DISC
---	---------------	----------------------------------

TOPIRAMATE

TABLET; ORAL TOPAMAX + JOHNSON RW	25MG 200MG	N20505 004 DEC 24, 1996 MAR CRLD N20505 002 DEC 24, 1996 MAR CRLD
---	---------------	--

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC TRAVATAN + ALCON UNIVERSAL	0.004%	N21257 001 MAR 16, 2001 MAR NEWA
---	--------	----------------------------------

TRIAMCINOLONE

TABLET; ORAL

TRIAMCINOLONE

@ IMPAX LABS

4MG

N84340 001 APR 22, 1975 FEB DISC

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; INHALATION

AZMACORT

+ AVENTIS

0.1MG/INH

N18117 001 APR 23, 1982 SEP CAHN

AEROSOL, METERED; NASAL

NASACORT

+ AVENTIS

0.055MG/INH

N19798 001 JUL 11, 1991 SEP CAHN

CREAM; TOPICAL

TRIAMCINOLONE ACETONIDE

@ TARO

0.025%

N40038 001 OCT 26, 1994 MAY DISC

@ TOPIDERM

0.025%

N89274 001 FEB 21, 1989 FEB WDRP

@

0.1%

N89275 001 FEB 21, 1989 FEB WDRP

@

0.5%

N89276 001 FEB 21, 1989 FEB WDRP

OINTMENT; TOPICAL

ARISTOCORT

@ FUJISAWA HLTHCARE

0.5%

N80745 002 MAY 28, 1974 JUL DISC

ARISTOCORT A

@ FUJISAWA HLTHCARE

0.5%

N80745 003 SEP 23, 1975 JUL DISC

TRIAMCINOLONE ACETONIDE

@ G AND W LABS

0.025%

N89795 001 DEC 23, 1988 JUL DISC

@

0.1%

N89796 001 DEC 23, 1988 JUL DISC

AT THAMES

0.025%

N40374 001 JUN 05, 2001 JUN NEWA

AT

0.5%

N40386 001 JUN 05, 2001 JUN NEWA

SPRAY; TOPICAL

KENALOG

+ APOTHECON

0.147MG/GM

N12104 001 DEC 24, 1959 JUL CDFR

SPRAY, METERED; NASAL

NASACORT AQ

+ AVENTIS

0.055MG/SPRAY

N20468 001 MAY 20, 1996 SEP CAHN

TRICHLORMETHIAZIDE

TABLET; ORAL

TRICLOREX

@ LANNETT

4MG

N83436 001 AUG 11, 1980 MAY DISC

@

4MG

N85630 001 MAY 16, 1977 FEB WDRP

>D> BP IMPAX LABS

4MG

N83967 001 JAN 17, 1978 OCT DISC

>A>

4MG

N83967 001 JAN 17, 1978 OCT DISC

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

TRIFLUOPERAZINE HCL

@ GENEVA PHARMS

EQ 10MG BASE/ML

N85787 001 APR 15, 1982 MAY DISC

TABLET; ORAL

AB GENEVA PHARMS TECH

EQ 1MG BASE

N40153 001 OCT 25, 1996 JAN CAHN

AB

EQ 2MG BASE

N40153 002 OCT 25, 1996 JAN CAHN

AB

EQ 5MG BASE

N40153 003 OCT 25, 1996 JAN CAHN

AB EQ 10MG BASE N40153 004 OCT 25, 1996 JAN CAHN

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HCL

@ STERIS

100MG/ML

N86577 001 OCT 19, 1982 JUL DISC

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

@ ASCENT PEDS

EQ 25MG BASE/5ML

N74374 001 JUN 23, 1995 JUN DISC

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

SURMONTIL

SIDMAK LABS

EQ 25MG BASE

N16792 001 JUN 12, 1979 AUG CAHN

EQ 50MG BASE

N16792 002 JUN 12, 1979 AUG CAHN

+

EQ 100MG BASE

N16792 003 SEP 15, 1982 AUG CAHN

TRIPELENNAmine HYDROCHLORIDE

TABLET; ORAL

TRIPELENNAmine HCL

>D> AA IMPAX LABS

50MG

N80785 001 AUG 07, 1973 OCT DISC

>A>

@

50MG

N80785 001 AUG 07, 1973 OCT DISC

TRIPLE SULFA (SULFABENZAMIDE; SULFACETAMIDE; SULFATHIAZOLE)

CREAM; VAGINAL

TRIPLE SULFA

@ FOUGERA

3.7%;2.86%;3.42%

N86424 001 MAY 31, 1979 JUN DISC

TRIPTORELIN PAMOATE

INJECTABLE; INTRAMUSCULAR

TRELSTAR

+ DEBIO RECHERCHE

11.25MG/VIAL

N21288 001 JUN 29, 2001 JUN NEWA

+ PHARMACIA AND UPJOHN

11.25MG/VIAL

N21288 001 JUN 29, 2001 SEP CAHN

TRELSTAR DEPOT

+ DEBIO RECHERCHE

EQ 3.75MG BASE/VIAL

N20715 001 JUN 15, 2000 JUN CDFR

+ PHARMACIA AND UPJOHN

EQ 3.75MG BASE/VIAL

N20715 001 JUN 15, 2000 SEP CAHN

TUBOCURARINE CHLORIDE

INJECTABLE; INJECTION

TUBOCURARINE CHLORIDE

@ LILLY

3MG/ML

N06325 001 NOV 28, 1947 SEP WDAG

URACIL MUSTARD

CAPSULE; ORAL

URACIL MUSTARD

@ SHIRE PHARM

1MG

N12892 001 SEP 13, 1962 JUN DISC

UREA, C-13

FOR SOLUTION; ORAL

BREATHTEK UBT FOR H-PYLORI

+ MERETEK

EQ 75MG /POUCHE

N20586 002 MAY 10, 2001 AUG NEWA

UREA, C-13

FOR SOLUTION; ORAL
 MERETEK UBT KIT (W/ PRANACTIN)
 @ MERETEK 125MG/VIAL

N20586 001 SEP 17, 1996 AUG DISC

VALGANCICLOVIR HYDROCHLORIDE

TABLET; ORAL
 VALCYTE
 + SYNTEX (USA) INC LLC EQ 450MG BASE

N21304 001 MAR 29, 2001 MAR NEWA

VALPROIC ACID

CAPSULE; ORAL
 VALPROIC ACID
 @ PAR PHARM 250MG
 @ SCHERER RP 250MG

N70431 001 FEB 28, 1986 MAY DISC
N70195 001 JUL 02, 1987 JUL DISCVALSARTAN

TABLET; ORAL
 DIOVAN
 NOVARTIS 80MG
 + 160MG
 320MG

N21283 001 JUL 18, 2001 JUL NEWA
N21283 002 JUL 18, 2001 JUL NEWA
N21283 003 JUL 18, 2001 JUL NEWAVANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION
 VANCOMYCIN HCL
 @ ELKINS SINK EQ 500MG BASE/VIAL
 @ EQ 1GM BASE/VIAL

N62879 001 AUG 02, 1988 MAY DISC
N62879 002 AUG 02, 1988 MAY DISCVECURONIUM BROMIDE

INJECTABLE; INJECTION
 VECURONIUM BROMIDE
 + ABBOTT 4MG/VIAL

N75558 001 SEP 11, 2001 SEP NEWA

VINBLASTINE SULFATE

INJECTABLE; INJECTION
 VELBAN
 @ LILLY 10MG/VIAL
 VINBLASTINE SULFATE
 AP + BEDFORD 10MG/VIAL

N12665 001 MAR 06, 1961 MAY DISC
N89395 001 APR 09, 1987 MAY CRLDVITAMIN A PALMITATE

CAPSULE; ORAL
 VITAMIN A
 @ WEST WARD EQ 50,000 UNITS BASE
 INJECTABLE; INJECTION
 AQUASOL A
 + NEOSAN PHARMS EQ 50,000 UNITS BASE/ML

N80967 001 MAY 04, 1973 FEB WDRP
N06823 001 MAY 18, 1949 AUG CAHN

WARFARIN SODIUM

TABLET; ORAL						
COUMADIN						
AB DUPONT MERCK	2.5MG	N09218 018	NOV 29, 1961	JUL	CRLD	
AB +	5MG	N09218 007	FEB 17, 1964	JUL	CRLD	
AB	5MG	N09218 007	FEB 17, 1964	AUG	CRLD	

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL						
GEODON						
PFIZER	20MG	N20825 001	FEB 05, 2001	FEB	NEWA	
	40MG	N20825 002	FEB 05, 2001	FEB	NEWA	
	60MG	N20825 003	FEB 05, 2001	FEB	NEWA	
+	80MG	N20825 004	FEB 05, 2001	FEB	NEWA	

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)						
ZOMETA						
+ NOVARTIS	EQ 4MG BASE/VIAL	N21223 001	AUG 20, 2001	SEP	CPOT	
+	4.264MG/VIAL	N21223 001	AUG 20, 2001	AUG	NEWA	

ZOLMITRIPTAN

TABLET, ORALLY DISINTEGRATING; ORAL						
ZOMIG-ZMT						
ASTRAZENECA	2.5MG	N21231 001	FEB 13, 2001	FEB	NEWA	

PREScription DRUG PRODUCT LIST - 21ST EDITION
OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 10 - OCT 2001

2-1

ACETAMINOPHEN

SUPPOSITORy; RECTAL

ACETAMINOPHEN

@ ABLE	120MG	N73106 001	FEB 27, 1995	SEP	WDAG
@	325MG	N73107 001	FEB 27, 1995	SEP	WDAG
@	650MG	N73108 001	FEB 27, 1995	SEP	WDAG
ALPHARMA US PHARM	120MG	N18337 003	SEP 12, 1983	MAR	CAHN
	325MG	N18337 002	AUG 21, 1981	MAR	CAHN
+	650MG	N18337 001	APR 22, 1980	MAR	CAHN
INFANTS' FEVERALL					
ALPHARMA US PHARM	80MG	N18337 004	AUG 26, 1992	MAR	CAHN

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

TAVIST ALLERGY/SINUS/HEADACHE

+ NOVARTIS	500MG;EQ 0.25MG BASE;30MG	N21082 001	MAR 01, 2001	MAR	NEWA
------------	---------------------------	------------	--------------	-----	------

ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

AVAGARD

+ 3M	61%;1%	N21074 001	JUN 07, 2001	JUN	NEWA
------	--------	------------	--------------	-----	------

>A> ASPIRIN

>A> TABLET; ORAL

>A> BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

>A> + BAYER	500MG	N21317 001	OCT 18, 2001	OCT	NEWA
-------------	-------	------------	--------------	-----	------

BROMPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BROMATAPP

@ COBLEY PHARM	12MG;75MG	N71099 001	JUL 02, 1987	JUL	DISC
----------------	-----------	------------	--------------	-----	------

CLEMASTINE FUMARATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TAVIST-D

@ NOVARTIS	1.34MG;75MG	N18298 002	AUG 21, 1992	JAN	DISC
------------	-------------	------------	--------------	-----	------

@	1.34MG;75MG	N20640 001	AUG 09, 1996	JAN	DISC
---	-------------	------------	--------------	-----	------

CLOTRIMAZOLE

CREAM; VAGINAL

TRIVAGIZOLE 3

TARO	2%	N21143 001	APR 12, 2000	JUL	CRLD
------	----	------------	--------------	-----	------

CROMOLYN SODIUM

SPRAY, METERED; NASAL

CROMOLYN SODIUM

ALPHARMA	5.2MG/INH	N74800 001	JUL 26, 2001	JUL	NEWA
----------	-----------	------------	--------------	-----	------

BAUSCH AND LOMB	5.2MG/SPRAY	N75702 001	JUL 03, 2001	JUL	NEWA
-----------------	-------------	------------	--------------	-----	------

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

DR REDDYS LABS LTD

10MG

N75758 001 AUG 17, 2001 AUG NEWA

TEVA

10MG

N75312 001 MAY 31, 2001 MAY NEWA

ZENITH GOLDLINE

10MG

N75512 001 JUL 26, 2001 JUL NEWA

IBUPROFEN

TABLET; ORAL

>D> ACHE-S-N-PAIN

LEDERLE

200MG

N71065 001 MAY 28, 1987 OCT DISC

>D>

@

200MG

N71065 001 MAY 28, 1987 OCT DISC

>A>

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

IBUPROHM COLD AND SINUS

OHM LABS

200MG;30MG

N74567 001 APR 17, 2001 APR NEWA

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

REGULAR PURIFIED PORK INSULIN

+ NOVO NORDISK

100 UNITS/ML

N18381 001 MAR 17, 1980 MAY CTEC

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

NOVOLIN R

+ NOVO NORDISK

100 UNITS/ML

N19938 001 JUN 25, 1991 MAY CTEC

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 MAY CTEC

NOVOLIN 70/30

+ NOVO NORDISK

30 UNITS/ML;70 UNITS/ML

N19991 001 JUN 25, 1991 MAY CTEC

INSULIN RECOMBINANT PURIFIED HUMAN

INJECTABLE; INJECTION

VELOSULIN BR HUMAN

@ NOVO NORDISK

100 UNITS/ML

N19450 001 MAY 30, 1986 SEP WDAG

MICONAZOLE NITRATE

CREAM; TOPICAL

MONISTAT 3 COMBINATION PACK

+ PERSONAL PRODS

2%;4%

N21261 001 FEB 02, 2001 FEB NEWA

CREAM; TOPICAL, VAGINAL

+ PERSONAL PRODS

2%;4%

N21261 001 FEB 02, 2001 MAY CDFR

CREAM; VAGINAL

MINOXIDIL

SOLUTION; TOPICAL

MINOXIDIL EXTRA STRENGTH (FOR MEN)

PERRIGO

5%

N75598 001 JUN 13, 2001 JUN NEWA

MINOXIDIL

SOLUTION; TOPICAL

MINOXIDIL EXTRA STRENGTH FOR MEN

>A> NOVEX 5% N75839 001 OCT 01, 2001 OCT NEWA

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

>A> NICORETTE (ORANGE)

>A> + SMITHKLINE BEECHAM EQ 4MG BASE N20066 004 SEP 25, 2000 OCT NEWA

>A> NICORETTE (ORANGE)

>A> + SMITHKLINE BEECHAM EQ 2MG BASE N18612 004 SEP 25, 2000 OCT NEWA

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

CUMULATIVE SUPPLEMENT NUMBER 10 OCTOBER '01

NO OCTOBER 2001 APPROVALS

This data is provided to the Division of Data Management and Services from the Office of Orphan Products Development and it is not edited prior to publication.

Orphan Products Designations and Approvals List
October 2001

Name:	Sponsor & Address
Generic Name	DD=Date Designated
<u>TN</u> =Trade Name	MA=Marketing Approval
Indication Designated:	

(R)-N-[2-(6-chloro-5-methyl-1H-indol-3-yl)propyl]acetamide Treatment of circadian rhythm sleep disorders in blind people with no light perception Phase 2 Discovery, Inc.
TN= 3130 Highland Avenue, Third Floor
Cincinnati OH 45219-2374
DD= 10/3/01 MA=

2-chloroethyl-3-sarcosinamide-1-nitrosourea Treatment for malignant gliomas Lawrence Panasci, MD
TN= Professor of Medicine, McGill
3755 Cote Ste Catherine
Montreal, Quebec H3T 1E2
DD= 8/3/01 MA=

2-methoxyestradiol Treatment of multiple myeloma EntreMed, Inc.
TN=Panzem 9640 Medical Center Drive
Rockville MD 20850
DD= 7/10/01 MA=

3-(4'aminoisoindoline-1'-one)-1-piperidine-2,6-dione Treatment for multiple myeloma Celegene Corporation
TN=Revimid (proposed) 7 Powder Horn Drive
Warren NJ 07059
DD= 9/20/01 MA=

9-nitro-20-(S)-camptothecin Treatment of pediatric HIV infection/AIDS NovoMed Pharmaceuticals, Inc.
TN=Camvirex P.O. Box 900
Germantown MD 20875-0900
DD= 5/15/01 MA=

acetylcysteine For the intravenous treatment of moderate to severe acetaminophen overdose Cumberland Pharmaceuticals Inc.
TN=Acetadote 209 10th Street South
Suite 332
Nashville TN 37203
DD= 10/19/01 MA=

of

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
Adeno-associated viral vector containing the gene for human coagulation factor IX TN=Coagulin-B	Intrahepatic treatment of patients with moderate to severe hemophilia	Avigen, Inc. 1301 Harbor Bay Parkway Alameda CA 94502 DD= 6/13/01 MA=
Adeno-associated viral vector containing the gene for human coagulation factor IX TN=Coagulin-B	Intramuscular treatment of patients with moderate to severe hemophilia	Avigen, Inc. 1301 Harbor Bay Parkway Alameda CA 94502 DD= 6/13/01 MA=
adenovirus-mediated herpes simplex virus-thymidine kinase TN=	Use with gancyclovir in the treatment of malignant glioma	Ark Therapeutics Ltd 6 Warren Mews London W1T 6AR UK DD= 7/31/01 MA=
Alendronate disodium TN=Fosamax	Treatment of the bone manifestations of Gaucher disease	Richard J. Wenstrup, M.D. Division of Human Genetics Children's Hospital Research Cincinnati OH 45229-3039 DD= 2/13/01 MA=
Angiotensin 1-7 TN=MARstem	Treatment of myelodysplastic syndrome	Maret Pharmaceutical Corporation 4041 MacArthur Boulevard, Suite Newport Beach CA 92660 DD= 8/3/01 MA=
arsenic trioxide TN=Trisenox	Treatment of chronic myeloid leukemia	Cell Therapeutics, Inc. 201 Elliott Avenue West Suite 400 Seattle WA 98119 DD= 10/18/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
augmerosan <u>TN=GenaSense</u>	Treatment of multiple myeloma	Genta Incorporated Two Oak Way Berkeley Heights NJ 07922 DD= 8/28/01 MA=
augmerosan <u>TN=GenaSense</u>	Treatment of chronic lymphocytic leukemia	Genta Incorporated Two Oak Way Berkeley Heights NJ 07922 DD= 8/28/01 MA=
augmerosan <u>TN=GenaSense</u>	Treatment of acute myelocytic leukemia	Genta Incorporated Two Oak Way Berkeley Heights NJ 07922 DD= 8/28/01 MA=
B Lymphocyte Stimulator <u>TN=BLYS</u>	Treatment of common variable immunodeficiency (CVID)	Human Genome Sciences, Inc. 9410 Key West Avenue Rockville MD 20850 DD= 2/21/01 MA=
Beclomethasone 17,21-dipropionate <u>TN=</u>	Prevention of gastrointestinal graft-versus-host disease	Enteron Pharmaceuticals, Inc. 1680 Michigan Ave. Suite 700 Miami FL 33139 DD= 8/28/01 MA=
Benzophenone-3, octylmethoxycinnamate, avobenzone, titanium <u>TN= Total Block VL</u> SPF 75	For the prevention of visible light induced skin photosensitivity as a result of porfimer sodium dioxide, zinc oxide photodynamic therapy	Fallien Cosmeceuticals Ltd. 677 W. Dekalb Pike King of Prussia PA 19406 DD= 8/13/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
Busulfan <u>TN=Spartajet-Busulfan</u>	Intrathecal therapy for neoplastic meningitis	SuperGen, Inc. 4140 Dublin Boulevard Dublin CA 94568 DD= 3/5/01 MA=
Coenzyme Q10 <u>TN=</u>	For the treatment of Huntington's disease	Vitaline Corporation 385 Williamson Way Ashland OR 97520 DD= 3/5/01 MA=
DHA-paclitaxel <u>TN=Taxoprexin</u>	Treatment of pancreatic cancer	Protarga, Inc. 2200 Renaissance Blvd. Suite 450 King of Prussia PA 19406 DD= 9/25/01 MA=
digitoxin <u>TN=</u>	Treatment of soft tissue sarcomas	PrimeCyte, Inc. 130 Fifth Ave., N. Seattle WA 98109-4933 DD= 10/18/01 MA=
docosahexanoic acid-paclitaxel <u>TN=Taxoprexin</u>	Treatment of hormone-refractory prostate cancer.	Protarga, Inc. 1100 East Hector Street Suite 450 Conshohocken PA 19428-2377 DD= 3/5/01 MA=
Glatiramer acetate for Injection <u>TN=Copaxone</u>	Treatment of primary-progressive multiple sclerosis	TEVA Pharmaceuticals, USA 1090 Horsham Road North Wales PA 19454 DD= 6/5/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
h5G1.1mAb TN=	Idiopathic membranous glomerular nephropathy	Alexion Pharmaceuticals, Inc. 352 Knotter Drive Cheshire CT 06410 DD= 3/5/01 MA=
Hsp E7 TN=	Treatment of recurrent respiratory papillomatosis (RRP)	StressGen Biotechnologies, Inc. 409 2nd Avenue Suite 201 Collegeville PA 19426-2655 DD= 3/19/01 MA=
human gammaglobulin TN=	Treatment for juvenile rheumatoid arthritis	Protein Therapeutics, Inc 9040 S. Rita Rd., Suite 1100 Tucson AZ 84747 DD= 5/25/01 MA=
humanized monoclonal antibody against Shiga-like toxin II TN=	To prevent the development of or to decrease the incidence and severity of hemolytic uremic syndrome and associated sequelae of Shiga-like toxin-producing <i>E. coli</i> .	Teijin America, Inc. 600 Alexander Park Suite 304 Princeton NJ 08540 DD= 9/12/01 MA=
Imatinib TN=Gleevec	Treatment of chronic myelogenous leukemia	Novartis Pharmaceuticals 59 Route 10 East Hanover NJ 07936-1080 DD= 1/31/01 MA= 5/10/01
Imexon TN= n/a	Treatment of metastatic malignant melanoma	AmpliMed Corporation 2321 Camino La Zorrela Tucson AZ 85718 DD= 8/3/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
INH-A00021 TN=	Reduction (prevention) of nosocomial bacteremia caused by staphylococci in very low birth weight infants.	Inhibitex, Inc. 8995 Westside Parkway Suite 150 Alpharetta GA 30004 DD= 6/13/01 MA=
C. Interferon-alfa-1b TN=	Treatment of multiple myeloma	Ernest C.Borden Center for Cancer Drug Discovery 9500 Euclid Avenue Cleveland OH 44195 DD= 4/17/01 MA=
Intraoral fluoride releasing system TN=IFRS	Prevention of dental caries due to radiation-induced xerostomia in patients with head and neck cancer	Digestive Care, Inc. 1120 Win Drive Bethlehem PA 18017 DD= 7/31/01 MA=
L-glutamine TN=	Treatment of sickle cell disease	Orphan Drugs International, LLC PO Box 0401 Montrose CA 91021-0401 DD= 8/1/01 MA=
Latrodectus immune F(ab)2 TN=Aracmyn	Treatment of black widow spider envenomations	Rare Disease Therapeutics, Inc. 1101 Kermit Drive, Suite 608 Nashville TN 37217 DD= 6/18/01 MA=
Medroxyprogesterone acetate TN=Hematrol	Treatment of immune thrombocytopenic purpura.	InKine Pharmaceutical Company, 1787 Sentry Parkway West Building 18, Suite 440 Blue Bell PA 19422 DD= 2/22/01 MA=
metreleptin TN=	Treatment of metabolic disorders secondary to lipodystrophy	Amgen, Inc., One Amgen Center Drive Thousand Oaks CA 91320-1799 DD= 8/22/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
metreleptin TN=	Treatment of leptin deficiency secondary to generalized lipodystrophy and partial familial lipodystrophy	Amgen, Inc. One Amgen Center Drive Thousand Oaks CA 91320-1799 DD= 8/22/01 MA=
MTC-DOX for Injection TN=	Treatment of hepatocellular carcinoma	FeRx Incorporated 4330 La Jolla Village Drive Suite #250 San Diego CA 92122 DD= 1/3/01 MA=
muramyltripeptide, phosphatidyl-ethanolamin e encased in multi-lamellar liposomes TN=	Treatment of children and adolescent osteosarcoma	Jenner Biotherapies, Inc. 541 Kenosa Street Walworth WI 53184 DD= 6/5/01 MA=
nitazoxanide TN=Cryptaz	Treatment for intestinal amebiasis	Romark Laboratories, L.C. 6200 Courtney Campbell Causeway Suite 880 Tampa FL 33607 DD= 10/23/01 MA=
Nitisinone TN=Orfadin	Treatment of alkaptonuria	Swedish Orphan AB Kungsgatan 37, 7th Floor SE-111 56 Stockholm, Sweden DD= 10/19/01 MA=
Nitroprusside TN=	Treatment and prevention of cerebral vasospasm following subarachnoid hemorrhage.	Thomas, MD, Jeffrey Evan Thomas Jefferson University 834 Walnut Street, Suite 650 Philadelphia PA 19107-5102 DD= 2/21/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name TN=Trade Name	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
Nolatrexed TN=THYMITAQ	Treatment of hepatocellular carcinoma	Zarix, Inc. 1055 Westlakes Drive Suite 200 Berwyn PA 19312 DD= 10/18/01 MA=
Novel Acting Thrombolytic (NAT) TN=	Treatment of peripheral arterial occlusion (PAO)	Amgen, Inc. One Amgen Center Drive Thousand Oaks CA 91320-1799 DD= 1/26/01 MA=
NZ-1002 TN=	Enzyme replacement therapy in patients with all subtypes of Mucopolysaccharidosis I.	Novazyme Pharmaceuticals, Inc. 800 Research Parkway Suite 200 Oklahoma City OK 73104 DD= 4/11/01 MA=
oglufanide disodium TN=	Treatment of ovarian cancer	Cytran, Inc. 10230 NE Points Dr., NE Suite 530 Kirkland WA 98033-7869 DD= 9/24/01 MA=
p1-(uridine 5'--)p4-(2'-deoxycytidin e 5'--) tetraphosphate, tetrasodium salt TN=	For the treatment of cystic fibrosis	Inspire Pharmaceuticals, Inc. 4222 Emperor Blvd. Suite 470 Durham NC 27703 DD= 3/7/01 MA=
pemetrexed disodium TN=Alimta	Treatment of malignant pleural mesothelioma	Eli Lilly and Company Lilly Corporate Center Indianapolis IN 46285 DD= 8/28/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
Perflubron <u>TN=LiquiVent</u>	Treatment of acute respiratory distress disease (ARDS) in adults	Alliance Pharmaceutical Corp. 3040 Science Park Road San Diego CA 92191 DD= 4/26/01 MA=
Polyethylene glycol (PEG)-uricase <u>TN=</u>	To control the clinical consequences of hyperuricemia in patients with severe gout in whom conventional therapy is contraindicated or has been ineffective.	Bio-Technology General Corporation 70 Wood Avenue South Iselin NJ 08830 DD= 2/21/01 MA=
porfimer <u>TN= Photofrin</u>	For the ablation of High-Grade Dysplasia in Barrett's Esophagus in patients who are not considered to be candidates for esophagectomy	Axcan Scandipharm Inc. 22 Inverness Parkway Suite 310 Birmingham AL 35242 DD= 10/19/01 MA=
Pyruvate <u>TN=</u>	Treatment of interstitial lung disease.	Cellular Sciences, Inc 84 park Avenue P.O. Box 968 Flemington NJ 08822 DD= 2/21/01 MA=
recombinant human alpha-1 antitrypsin (rAAT) <u>TN=</u>	To delay progression of chronic obstructive pulmonary disease resulting from AAT deficiency-mediated emphysema and bronchiectasis	Baxter Healthcare Corporation 550 N. Brand Blvd. Glendale CA 91203 DD= 8/28/01 MA=
Recombinant Human Alpha-Fetoprotein <u>TN=</u>	Treatment of myasthenia gravis	Atlantic Biopharmaceuticals, Inc. 50 Church Street 5th floor Cambridge MA 02138 DD= 2/22/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
recombinant human endostatin protein <u>TN=</u>	Treatment of neuroendocrine tumors.	EntreMed, Inc. 9640 Medical Center Drive Rockville MD 20850 DD= 8/13/01 MA=
Reviparin sodium <u>TN=Clivarine</u>	Treatment of deep vein thrombosis which may lead to pulmonary embolism in pediatric patients	Knoll AG Ludwigshafen, Germany DD= 6/18/01 MA=
Reviparin sodium <u>TN=Clivarine</u>	Long-term treatment of acute deep vein thrombosis with or without pulmonary embolism in pregnant patients	Knoll AG Ludwigshafen, Germany DD= 6/18/01 MA=
squalamine lactate <u>TN=</u>	Treatment of ovarian cancer refractory or resistant to standard chemotherapy	Genaera Corporation 5110 Campus Drive Plymouth Meeting PA 19462 DD= 5/11/01 MA=
Synthetic Human Parathyroid Hormone 1-34 <u>TN=</u>	Treatment of hypoparathyroidism	Orphan Pharmaceuticals, U.S., Inc. 1101 Kermit Drive, Suite 608 Nashville TN 37217 DD= 1/26/01 MA=
Thyrotropin alfa <u>TN=Thyrogen</u>	Treatment of well-differentiated papillary, follicular or combined papillary/follicular carcinomas of the thyroid	Genzyme Corporation One Kendall Square Cambridge MA 02139-1562 DD= 8/3/01 MA=
Unconjugated Chimeric (human-murine) G250 IgG monoclonal antibody <u>TN=</u>	Treatment of renal cell carcinoma.	Wilex Biotechnology GmbH Grillparzerstrasse 10B 81675 Munich Germany DE DD= 3/22/01 MA=

Orphan Products Designations and Approvals List
October 2001

Name: Generic Name <u>TN=Trade Name</u>	Indication Designated:	Sponsor & Address DD=Date Designated MA=Marketing Approval
Vasoactive intestinal peptide <u>TN=</u>	Treatment of Acute Respiratory Distress Syndrome.	Sami I. Said, M.D. State University of New York at Health Sciences Center T17, 040 Stony Brook NY 11794-8172 DD= 3/9/01 MA=
Virulizin <u>TN=Virulizin</u>	Treatment of pancreatic cancer.	Lorus Therapeutics Inc. 7100 Woodbine Avenue, Suite 215 Markham, ON L3R 5J2 Canada DD= 2/1/01 MA=

NO OCT

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

NO OCTOBER 2001 ADDITIONS

k at
040

215

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019408 002 021056 001	BETAMETHASONE DIPROPIONATE; DIFROLENE BEXAROTENE; TARGRETIN	4489070 5780576 5962273 5466861 1712251 5712251 5688819	MAY 13, 2003 JUL 14, 2016 OCT 05, 2016 NOV 14, 2012 SEP 18, 2001 SEP 18, 2001 SEP 21, 2012	ODE	DEC 29, 2006		
020498 001	BICALUTAMIDE; CASODEX						
021275 001 021290 001	BIMATOPROST; LUMIGAN BOSENTAN; TRACLEER						
021290 002	BOSENTAN; TRACLEER						
020490 001	BRIMONIDINE TARTRATE; ALPHAGAN	6194415 6248741 6194415*PED 6248741*PED	JUN 28, JUN 28, DEC 28, DEC 28,	2015 2015 2015 2015	U-394 U-394 U-394 U-394	NCE NCE NCE NCE	MAR 16, 2006 NOV 20, 2008 NOV 20, 2006 NOV 20, 2008
020613 001	BRIMONIDINE TARTRATE; ALPHAGAN P	6194415*PED 5736165 5736165*PED 6248741	DEC 28, APR 07, OCT 07, JUN 28,	2015 2015 2015 2015	U-395 U-399 U-399 U-395	NCE NCE NCE NCE	SEP 06, 2001 SEP 06, 2002 SEP 06, 2001 MAR 06, 2002
021262 001	BRIMONIDINE TARTRATE; ALPHAGAN P	6194415 6248741*PED	JUN 28, DEC 28,	2015 2015	U-395 U-395	NCE NCE	MAR 16, 2004 SEP 16, 2004
020816 001 021324 001 020746 001	BRINZOLAMIDE; AZOPT BUDESONIDE; ENTOCORT EC BUDESONIDE; RHINOCORT	5424078 5378703 5643602 6291445	JUN 13, APR 01, JUL 01, APR 29,	2012 2012 2012 2017	U-224	NP	OCT 02, 2004
020358 001 020358 002 020358 003 018731 001 018731 002 018731 003 018731 004 074253 001 074253 002 075272 003 075467 002 076008 001 020524 001 018874 001	BUPROPTION HYDROCHLORIDE; WELLBUTRIN SR BUPROPTION HYDROCHLORIDE; WELLBUTRIN SR BUPROPTION HYDROCHLORIDE; WELLBUTRIN SR BUPTRONE HYDROCHLORIDE; BUSPAR BUPTRONE HYDROCHLORIDE; BUSPAR BUPTRONE HYDROCHLORIDE; BUSPAR BUPTRONE HYDROCHLORIDE; BUSPAR BUPTRONE HYDROCHLORIDE; BUSPIRONE HCL BUPTRONE HYDROCHLORIDE; BUSPIRONE HCL BUPTRONE HYDROCHLORIDE; BUSPIRONE HCL BUPTRONE HYDROCHLORIDE; BUSPIRONE HCL BUPTRONE HYDROCHLORIDE; BUSPIRONE HCL BUTENAFINE HYDROCHLORIDE; MENTAX CALCITRIOL; CALCIJEX						
		4308264 6051567 4308264*PED 6051567*PED 6265392 6274169 6274169*PED 6265392*PED	JAN 28, AUG 02, JUL 28, FEB 02, AUG 02, AUG 02, FEB 02, FEB 02,	2001 2001 2020 2019 2019 2019 2020 2020	I-333 JUN 06, 2004		

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME ; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL USE CODE	EXCLUS CODE	EXCLUS EXPIRES
018874 002	CALCITRIOL;CNLCIJEX	4308254 6051567 430824*PED 605157*PED	JAN 28, 2001 AUG 02, 2019 JUL 28, 2001 FEB 02, 2020			
019976 001	CALCIUM ACETATE ; PHOSLO	6265332 6274159 6274159*PED	AUG 02, AUG 02, FEB 02,	2019 2019 2020		
021160 001	CALCIUM ACETATE ; PHOSLO	6265392*PED	FEB 02,	2020		
021160 002	CALCIUM ACETATE ; PHOSLO	4870105	APR 07,	2007	U-381	
021160 003	CALCIUM ACETATE ; PHOSLO GELCAPS	4870105	APR 07,	2007	U-381	
021160 004	CALCIUM ACETATE ; PHOSLO GELCAPS	4870105	APR 07,	2007	U-381	
020896 001	CAPECITABINE;XELODA				I-341	APR 30, 2004
020896 002	CAPECITABINE;XELODA				I-341	SEP 07, 2004
020297 001	CARVEDILOL;COREG	5952300 5378804 5514650 5792246 6136783	MAR 28, 2017 MAR 16, 2013 MAR 16, 2013 MAR 16, 2013 MAR 28,	2017 2013 2013 2013 2017	NCE	JAN 26, 2006
020297 002	CARVEDILOL;COREG				I-343	NOV 01, 2004
020297 003	CARVEDILOL;COREG				I-343	NOV 01, 2004
020297 004	CARVEDILOL;COREG				I-343	NOV 01, 2004
021227 001	CASPOFUNGIN ACETATE;CANCIDAS				I-343	NOV 01, 2004
020998 001	CELECOXIB;CELEBREX	5792246 6136783	MAR 16, 2013 MAY 28,	2013 2017		
020998 002	CELECOXIB;CELEBREX				I-343	NOV 01, 2004
021197 001	CETRORELIIX;CETROTIDE	6319192 6319192 4762709	APR 23, APR 23, AUG 09,	2018 2018 2005	U-426 U-426 NP	OCT 17, 2004 OCT 17, 2004 SEP 20, 2003
021197 002	CETRORELIIX;CETROTIDE				I-341	OCT 17, 2004
>ADD>	CHLORPHENIRAMINE POLISTREX;TUSSIONEX	5767251 4957730 4847265 6177101	JUN 16, SEP 18, NOV 17, JUN 11,	2015 2007 2011 2018		
>ADD>	CHLOROGONADOTROPIN ALFA;OVIDREL					
019111 001	CICLOPIROX;PENIAC					
021149 001	CLOPIDOGREL BISULFATE;PLAVIX					
021022 001	DELAVIRDINE MESYLATE;RESPRITOR					
020839 001	DEXMEDETOMIDINE;PREDEX					
020705 001	DEXMETHYLPHENIDATE HYDROCHLORIDE ; DEXMETHYLPHENIDATE H	4910214 5922736 5908850 6255325 5922736 5908850	JUL 15, DEC 04, DEC 04, DEC 04, DEC 04, DEC 04,	2008 2015 2015 2015 2015 2015	U-421 U-423 U-422 U-424 U-423 U-422	NOV 13, 2004 NOV 13, 2004 NOV 13, 2004 NOV 13, 2004 NOV 13, 2004 U-424
021038 001	DEXMETHYLPHENIDATE HYDROCHLORIDE ; DEXMETHYLPHENIDATE H					
021278 001	DEXMETHYLPHENIDATE HYDROCHLORIDE ; DEXMETHYLPHENIDATE H					
021278 002	DEXMETHYLPHENIDATE HYDROCHLORIDE ; DEXMETHYLPHENIDATE H	6255325	DEC 04,	2015		

PRESCRIPTION AND OTC DRUG PRODUCT

PREScription AND UIC DRUG PRESCRIPTIONS

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

THE JOURNAL OF CLIMATE

PRESCRIPTION AND OTC DRUG PRODUCT

*PED and PED represent Pediatric Exclusivity

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	USE CODE	EXCLUS EXPIRES
020623 001	DOLASETRON MESYLATE MONOHYDRATE; ANZEMET	4906755	JUL 02, 2011			
020623 002	DOLASETRON MESYLATE MONOHYDRATE; ANZEMET	4906755	JUL 02, 2011			
020624 001	DOLASETRON MESYLATE MONOHYDRATE; ANZEMET	4906755	JUL 02, 2011			
020690 001	DONEPEZIL HYDROCHLORIDE; ARICEPT	6140321	DEC 30, 2016			
020690 002	DONEPEZIL HYDROCHLORIDE; ARICEPT	624511	DEC 01, 2018			
		5988864	DEC 30, 2016			
		6140321	DEC 30, 2016			
020869 001	DORZOLAMIDE HYDROCHLORIDE; COSOPT	6245911	DEC 01, 2018			
021098 001	DROSPRENONE; YASMIN	5988864	DEC 30, 2016			
021319 001	DUTASTERIDE; DUTASTERIDE	6248735	APR 17, 2011			
020706 001	EMEDASTINE DIFUMARATE; EMADINE	4430343	AUG 14, 2005			
020668 001	ENALAPRIL MALEATE; LEXXEL	5441958	DEC 08, 2013			
		4264611	JUN 19, 2001			
		4803081	APR 03, 2007			
		4264611*PED	DEC 19, 2001			
		4803081*PED	OCT 03, 2007			
		4374829	DEC 30, 2001			
		447230	SEP 18, 2001			
		4701038	OCT 07, 2005			
		4803081	APR 03, 2007			
		4264611	JUN 19, 2001			
		4264611*PED	DEC 19, 2001			
020668 002	ENALAPRIL MALEATE; LEXXEL	4803081*PED	OCT 03, 2007			
018998 001	ENALAPRIL MALEATE; VASOTEC	4803081*PED	OCT 03, 2007	M-7		
018998 002	ENALAPRIL MALEATE; VASOTEC			PED		
018998 003	ENALAPRIL MALEATE; VASOTEC			M-7		
018998 005	ENALAPRIL MALEATE; VASOTEC			PED		
020164 002	ENOXAPARIN SODIUM; LOVENOX	4486420	DEC 04, 2001			
		4692435	DEC 24, 2004			
		5389618	FEB 14, 2012			
		4486420	DEC 04, 2001			
		4692435	DEC 24, 2004			
		5389618	FEB 14, 2012			
		4486420	DEC 04, 2001			
		4692435	DEC 24, 2004			
020164 003	ENOXAPARIN SODIUM; LOVENOX	5389618	FEB 14, 2012			
020164 004	ENOXAPARIN SODIUM; LOVENOX	4486420	DEC 04, 2001			
		4692435	DEC 24, 2004			
		5389618	FEB 14, 2012			
		4486420	DEC 04, 2001			
		4692435	DEC 24, 2004			
020164 005	ENOXAPARIN SODIUM; LOVENOX	5389618	FEB 14, 2012			

**PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA**
and PED represent Pediatric Exclusivity

*PED and PED represent Pediatric Exclusivity

and PED represent Pediatric Exclusivity

**PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA**

PRESCRIPTION AND OTC DRUGS PRODUCED

PATENT AND EXCLUSIVITY DATA
AND PRESCRIBITION AND OTC DRUG PRODUCTION

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME / TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	USE CODE	EXCLUSIVITY CODE	EXCLUSIVITY EXPIRES
020870 001	ESTRADIOL; COMBIPATCH	5474783	DEC 12, 2012				
		565286	AUG 12, 2014				
		5958446	DEC 12, 2012				
020870 002	ESTRADIOL; COMBIPATCH	6024976	JAN 07, 2014				
		5474783	DEC 12, 2012				
		5656286	AUG 12, 2014				
020538 005	ESTRADIOL; VIVELLE-DOT	5958446	DEC 12, 2012				
		6024976	JAN 07, 2014				
020538 006	ESTRADIOL; VIVELLE-DOT	5474783	DEC 12, 2012				
		565286	AUG 12, 2014				
020538 007	ESTRADIOL; VIVELLE-DOT	5958446	DEC 12, 2012				
		6024976	JAN 07, 2014				
020538 008	ESTRADIOL; VIVELLE-DOT	5474783	DEC 12, 2012				
		565286	AUG 12, 2014				
020130 002	ETHINYL ESTRADIOL; ESTROSTEP FE	5951446	DEC 12, 2012				
020130 001	ETHINYL ESTRADIOL; ESTROSTEP 21	5010070	APR 23, 2008	I-331	JUL 01, 2004		
021187 001	ETHINYL ESTRADIOL; NUVARING	5010070	APR 23, 2008	I-331	JUL 01, 2004		
021180 001	ETHINYL ESTRADIOL; ORTHO EVRA			NP	OCT 03, 2004		
020946 001	ETHINYL ESTRADIOL; PREVYN EMERGENCY CON			NP	NOV 20, 2004		
020584 001	ETODOLAC; LODINE XL	6156742	DEC 05, 2020	U-374			
020584 002	ETODOLAC; LODINE XL						
020584 003	ETODOLAC; LODINE XL						
020457 001	ETOPOSIDE PHOSPHATE; ETOPOPHOS PRESERVATI	RE25524	MAY 17, 2010				
020906 001	ETOPOSIDE PHOSPHATE; ETOPOPHOS PRESERVATI	5041424	AUG 20, 2008	U-135			
020906 002	ETOPOSIDE PHOSPHATE; ETOPOPHOS PRESERVATI	RE35524	MAY 17, 2010				
		5041424	AUG 20, 2008	U-135			
		RE35524	MAY 17, 2010				
075312 001	FAMOTIDINE; FAMOTIDINE			PC	NOV 28, 2001		
020902 001	FAMOTIDINE; PEPCID AC			D-47	NOV 09, 2001		
019834 001	FELODIPINE; PLENDIL	4264611	JUN 19, 2001	U-3	MAY 09, 2002		
		4803081*PED	APR 03, 2007				
		4803081*PED	OCT 03, 2007				
		4264611*PED	DEC 19, 2001				
				U-3			

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019834 002	FELODIPINE; PLENDIL	4264611	JUN 19, 2001	U-3		
		4803081	APR 03, 2007			
		4244611*PED	DEC 19, 2001	U-3		
		4803081*PED	OCT 03, 2007			
019834 004	FELODIPINE; PLENDIL	4264611	JUN 19, 2001	U-3		
		4803081	APR 03, 2007			
		4264611*PED	DEC 19, 2001	U-3		
>ADD2	021203 001	FENOFIBRATE; TRICOR	4803081*PED	OCT 03, 2007		
>ADD2	021203 003	FENOFIBRATE; TRICOR	6277405	JAN 09, 2018		
>ADD2	020625 001	FEKOGENADINE HYDROCHLORIDE; ALLEGRA	6277405	JAN 09, 2018		
>ADD2	020872 001	FEKOGENADINE HYDROCHLORIDE; ALLEGRA	6074670	JAN 09, 2018		
>ADD2	020872 002	FEKOGENADINE HYDROCHLORIDE; ALLEGRA	6074670	JAN 09, 2018		
>ADD2	020872 004	FEKOGENADINE HYDROCHLORIDE; ALLEGRA	6074670	JAN 09, 2018		
>ADD2	020876 001	FEKOGENADINE HYDROCHLORIDE; ALLEGRA-D	6074670	JAN 09, 2018		
>ADD2	019452 001	FLUCCINOLONE ACETONIDE; DERMA-SMOOTHÉ / FS	6074670	JAN 09, 2018		
>ADD2	020985 001	FLUOROURACIL; CARAC	6074670	JAN 09, 2018		
>ADD2	074803 001	FLUOXETINE HYDROCHLORIDE; FLUOXETINE	4895726	MAY 11, 2012	U-138	
>ADD2	075049 001	FLUOXETINE HYDROCHLORIDE; FLUOXETINE	4895726	MAY 11, 2012	U-138	
>ADD2	075465 003	FLUOXETINE HYDROCHLORIDE; FLUOXETINE	4895726	MAY 11, 2012	U-138	
>ADD2	075506 001	FLUOXETINE HYDROCHLORIDE; FLUOXETINE HCL	4895726	MAY 11, 2012	U-138	
>ADD2	075755 001	FLUOXETINE HYDROCHLORIDE; FLUOXETINE HCL	4895726	MAY 11, 2012	U-138	
>ADD2	075755 002	FLUOXETINE HYDROCHLORIDE; PROZAC WEEKLY	4895726	MAY 11, 2012	U-138	
>ADD2	021235 001	FLUOXETINE HYDROCHLORIDE; PROZAC	4690825	OCT 04, 2005	I-340	OCT 10, 2004
>ADD2	018936 007	FLUOXETINE HYDROCHLORIDE; SARAFEM	5910319	MAY 29, 2017	PC	JAN 23, 2002
>ADD2	021077 001	FLUTICASONE PROPIONATE; ADVAIR DISKUS 100/50	5985322*PED	NOV 29, 2017	PC	JAN 23, 2002
>ADD2	021077 002	FLUTICASONE PROPIONATE; ADVAIR DISKUS 250/50	5985322*PED	NOV 29, 2017	PC	JAN 23, 2002
>ADD2	021077 003	FLUTICASONE PROPIONATE; ADVAIR DISKUS 500/50	5985322*PED	NOV 29, 2017	PC	JAN 23, 2002
>ADD2	020549 001	FLUTICASONE PROPIONATE; FLOVENT	5270305	SEP 07, 2010	U-387	
>ADD2	020549 002	FLUTICASONE PROPIONATE; FLOVENT	5290815	MAR 01, 2011	U-386	
>ADD2	020549 003	FLUTICASONE PROPIONATE; FLOVENT	5270305	SEP 07, 2010	U-387	
>ADD2	020833 001	FLUTICASONE PROPIONATE; FLOVENT DISKUS 50	5290815	MAR 01, 2011	U-386	
>ADD2	020833 002	FLUTICASONE PROPIONATE; FLOVENT DISKUS 100	5270305	SEP 07, 2010	U-387	
>ADD2	020833 003	FLUTICASONE PROPIONATE; FLOVENT DISKUS 250	5290815	MAR 01, 2011	U-386	

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020261 001	FLUVASTATIN SODIUM; LESCOL	5356896	DEC 12, 2011			
020261 002	FLUVASTATIN SODIUM; LESCOL	5356896	DEC 12, 2011			
021192 001	FLUVASTATIN SODIUM; LESCOL XL	5354772	OCT 11, 2011	U-413		
020831 001	FORMOTEROL FUMARATE; FORADIL	5356896	DEC 12, 2011			
021279 001	FORMOTEROL FUMARATE; FORADIL					
021006 001	FROVATRIPTAN SUCCINATE; FROVA	5464864	NOV 07, 2012	U-72	NCE	FEB 16, 2006
		5616603	APR 01, 2014	U-72	NCE	SEP 25, 2004
		5637611	JUN 10, 2014	U-72	NCE	FEB 16, 2006
		5827871	OCT 27, 2015	U-72		
021129 001	GABAPENTIN; NEURONTIN	5962501	DEC 16, 2013	U-72		
021169 001	GALANTAMINE HYDROBROMIDE; REMINYL	4894476	MAY 02, 2008			
021169 002	GALANTAMINE HYDROBROMIDE; REMINYL	5084479	JAN 02, 2010	U-258		
021169 003	GALANTAMINE HYDROBROMIDE; REMINYL	4693318	JAN 05, 2006			
021224 001	GALANTAMINE HYDROBROMIDE; REMINYL	6099863	JUN 06, 2017			
021061 001	GATIFLOXACIN; TEQUIN	4663318	JAN 15, 2006			
021061 002	GATIFLOXACIN; TEQUIN	6099863	JUN 06, 2017			
021062 001	GATIFLOXACIN; TEQUIN	4663318	JAN 15, 2006			
021062 002	GATIFLOXACIN; TEQUIN	6099863	JUN 06, 2017			
021178 001	GLYBURIDE; GLUCOVANCE	6303146	JUL 14, 2019			
021178 002	GLYBURIDE; GLUCOVANCE	6303146	JUL 14, 2019			
021178 003	GLYBURIDE; GLUCOVANCE	6303146	JUL 14, 2019			
020239 001	GRANISETRON HYDROCHLORIDE; KYTRIL	6294548	MAY 04, 2019			
020239 002	GRANISETRON HYDROCHLORIDE; KYTRIL	4886808	DEC 29, 2007	U-89		
021238 001	GRANISETRON HYDROCHLORIDE; KYTRIL	6294548	MAY 04, 2019			
020387 001	HYDROCHLOROTHIAZIDE; HYZZAR	4886808	DEC 29, 2007			
020387 002	HYDROCHLOROTHIAZIDE; HYZZAR	5608075	MAR 04, 2014			
019778 001	HYDROCHLOROTHIAZIDE; PRINZIDE	5608075	MAR 04, 2014			
019778 002	HYDROCHLOROTHIAZIDE; PRINZIDE	4374829	DEC 29, 2001	U-3		
019778 003	HYDROCHLOROTHIAZIDE; PRINZIDE	4374829*	PED JUN 29, 2002			
019888 001	HYDROCHLOROTHIAZIDE; ZESTORETIC	4374829	DEC 29, 2001	U-3		
019888 002	HYDROCHLOROTHIAZIDE; ZESTORETIC	4374829	DEC 29, 2001	U-3		
		4374829*	PED JUN 29, 2002			

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	EXCLUSIVITY CODE	EXCLUSIVITY EXPIRES
019888 003	HYDROCHLOROTHIAZIDE; ZESTORETIC	4 374 829 4 374 829 *PED	DEC 29, 2001 JUN 29, 2002	U-3 U-3	NP	MAR 16, 2003
020402 002	IBUPROFEN POTASSIUM; ADVIL MIGRAINE LIQUI	6 211 246	JUN 10, 2019			
021128 001	IBUPROFEN CHILDREN'S MOTRIN CO	5 521 184	MAY 28, 2013	NCE	MAY 10, 2006	
021335 001	IMATINIB MESYLATE; GLEEVEC	5 521 184	MAY 28, 2013	ODE	MAY 10, 2008	
021335 002	IMATINIB MESYLATE; GLEEVEC	5 521 184	MAY 28, 2013	NCE	MAY 10, 2006	
021172 001	INSULIN ASPART RECOMBINANT; NOVOLOG MIX 70/30	5 656 722	SEP 12, 2014	ODE	MAY 10, 2008	
021081 001	INSULIN GLARGINE; LANTUS	5 656 722 *PED	MAR 12, 2015	NC	NOV 01, 2004	
020394 001	IPRATROPIUM BROMIDE; ATROVENT	4 464 394	AUG 07, 2001	I-327	OCT 27, 2003	
018662 002	ISOTRETINOIN; ACCUTANE	4 464 394 *PED	FEB 07, 2002			
018662 003	ISOTRETINOIN; ACCUTANE	4 464 394	AUG 07, 2001			
018662 004	ISOTRETINOIN; ACCUTANE	4 464 394 *PED	FEB 07, 2002			
020657 001	ITRACONAZOLE; SPORANOX	4 791 111	DEC 23, 2005	I-332	MAY 09, 2004	
020966 001	ITRACONAZOLE; SPORANOX	4 454 151	MAR 22, 2002	I-332	MAY 09, 2004	
019700 001	KETOROLAC TROMETHAMINE; ACULAR	5 110 93	MAY 05, 2009	U-75		
		4 454 151 -PED	SEP 22, 2002	U-75		
		5 110 93 *PED	NOV 05, 2009	U-75		
020811 001	KETOROLAC TROMETHAMINE; ACULAR PRESERVATIVE	4 454 151	MAR 22, 2002	I-339	AUG 16, 2004	
020857 001	LAMIVUDINE; COMBIVIR	4 454 151 *PED	SEP 22, 2002	PED	FEB 16, 2005	
020564 001	LAMIVUDINE; EPIVIR	6 180 639	JAN 30, 2018			
020596 001	LAMIVUDINE; EPIVIR	6 180 639 *PED	JUL 30, 2018			
021003 001	LAMIVUDINE; EPIVIR-HBV	6 180 639	JAN 30, 2018			
021004 001	LAMIVUDINE; EPIVIR-HBV	6 180 639 *PED	JUL 30, 2018			
021281 001	LANSOPRAZOLE; PREVACID	6 180 639 *PED	JUL 30, 2018			
021281 002	LANSOPRAZOLE; PREVACID			D-42	JUL 20, 2001	
020905 001	LEFLUNOMIDE; ARAVA			I-316	NOV 30, 2003	
				M-1	JUL 06, 2002	
				D-42	JUL 20, 2001	
				M-1	JUL 06, 2002	
				D-42	JUL 20, 2001	

4284786 DEC 13, 2001

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT PED EXPIRES	PATENT EXCL CODE	USE CODE	EXCLUS CODE	EXCLUS CODE EXPIRES
020905 002	LEFLUNOMIDE;ARAVA	4284786	DEC 13, 2001				
020905 003	LEFLUNOMIDE;ARAVA	4284786	DEC 13, 2001	JUN 03,	2011	U-203	
020726 001	LETROZOLE;FEMARA	4978672		NOV 01,	2004		
019732 001	LEUPROLIDE ACETATE;LUPRON DEPOT	5631021		NOV 01,	2004		
		5476663		NOV 01,	2004		
		5476663		NOV 01,	2004		
020011 001	LEUPROLIDE ACETATE;LUPRON DEPOT	5631020		NOV 01,	2004		
		5476663		NOV 01,	2004		
020517 001	LEUPROLIDE ACETATE;LUPRON DEPOT	5575987		SEP 02,	2013		
		5575987		SEP 02,	2013		
020263 002	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5643607		JAN 02,	2013		
		5643607		JAN 02,	2013		
020263 003	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5631021		NOV 01,	2004		
		5476663		NOV 01,	2004		
020263 004	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5476663		NOV 01,	2004		
		5575987		NOV 01,	2004		
020263 005	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5631021		NOV 01,	2004		
		5575987		NOV 01,	2004		
020263 006	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5476663		NOV 01,	2004		
		5575987		NOV 01,	2004		
020708 001	LEUPROLIDE ACETATE;LUPRON DEPOT-3	5631021		NOV 01,	2004		
		5476663		NOV 01,	2004		
020517 002	LEUPROLIDE ACETATE;LUPRON DEPOT-4	5643607		JAN 02,	2013		
		5814342		FEB 01,	2011		
		5643607		NOV 01,	2004		
		5814342		FEB 01,	2011		
		6036976		DEC 13,	2016		

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
* PED and PED represent Pediatric Exclusivity

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	EXCLUS EXPIRES
021121 002	METHYLPHENIDATE HYDROCHLORIDE; CONCERTA	4783337	SEP 16, 2003	U-372	AUG 01, 2003
021121 003	METHYLPHENIDATE HYDROCHLORIDE; CONCERTA	4783337	SEP 16, 2003	NDF	APR 03, 2004
021259 001	METHYLPHENIDATE HYDROCHLORIDE; METADATE CD	4927640	MAY 22, 2007	I-194	FEB 05, 2004
019962 001	METOPROLOL SUCCINATE; TOPROL-XL	5246714	SEP 21, 2010		
019962 002	METOPROLOL SUCCINATE; TOPROL-XL	4927640	MAY 22, 2007	I-194	FEB 05, 2004
019962 003	METOPROLOL SUCCINATE; TOPROL-XL	5246714	SEP 21, 2010	I-194	FEB 05, 2004
019962 004	METOPROLOL SUCCINATE; TOPROL-XL	4927745	SEP 18, 2007	U-107	NS
		5001161	MAR 19, 2008	U-107	FEB 05, 2004
		5081154	JAN 14, 2009	U-107	
021308 001	MICONAZOLE NITRATE; MONISTAT 1 COMBINATI	4927640	MAY 22, 2007		
019436 001	MILRINONE LACTATE; PRIMACOR	5246714	SEP 21, 2010		
020343 001	MILRINONE LACTATE; PRIMACOR IN DEXTROSE	6153335	NOV 28, 2020		
020343 002	MILRINONE LACTATE; PRIMACOR IN DEXTROSE	5514698	MAR 21, 2014		
020343 003	MILRINONE LACTATE; PRIMACOR IN DEXTROSE	4313951*PED	NOV 26, 2001		
021208 001	MIRTAZAPINE; REMERON SOLTAB	4313951	NOV 26, 2001		
021208 002	MIRTAZAPINE; REMERON SOLTAB	4313951*PED	MAY 26, 2002		
021208 003	MIRTAZAPINE; REMERON SOLTAB	4313951	NOV 26, 2001		
019297 001	MITOXANTRONE HYDROCHLORIDE; NOVANTRONE	4313951*PED	MAY 26, 2002		
020762 001	MOMETASONE FUROATE MONOHYDRATE; NASONEX	4313951	NOV 26, 2001		
021208 001	MIRTAZAPINE; REMERON SOLTAB	4313951*PED	MAY 26, 2002		
021208 002	MIRTAZAPINE; REMERON SOLTAB	5178878	JAN 12, 2010	NCE	JUN 14, 2001
021208 003	MIRTAZAPINE; REMERON SOLTAB	5178878	JAN 12, 2010	NCE	JUN 14, 2001
019297 001	MITOXANTRONE HYDROCHLORIDE; NOVANTRONE	4617319	JUN 13, 2005	U-390	OCT 13, 2003
020762 001	MOMETASONE FUROATE MONOHYDRATE; NASONEX	4472393	SEP 18, 2001	I-285	DEC 02, 2002
019543 001	MOMETASONE FUROATE; ELOCON	5837769	JAN 27, 2014	U-249	PED
019625 001	MOMETASONE FUROATE; ELOCON	6127353	OCT 03, 2017		JUN 02, 2003
019796 001	MOMETASONE FUROATE; ELOCON	4472393*PED	MAR 18, 2002		
		5837769*PED	JUL 27, 2014	U-249	
		6127353*PED	APR 03, 2018		
		4472393*PED	MAR 18, 2002		
		4472393*PED	OCT 02, 2006		
		4808610	SEP 18, 2001		
		4472393	MAR 18, 2002		
		4472393*PED	APR 02, 2007		
		4808610*PED	APR 02, 2007		
		477529	MAY 21, 2007		
		4472393	SEP 18, 2001		
		4472393*PED	MAR 18, 2002		
		477529*PED	NOV 21, 2007		

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL USE CODE	EXCLUS EXCLUS CODE EXPIRES
				EXPIRES
020829 002	MONTLUKAST SODIUM;SINGULAIR	5565473 FEB 03, 2012	U-228 NCE	FEB 20, 2003
		5565473 *PED AUG 03, 2012	U-228 PED	AUG 20, 2003
020830 001	MONTLUKAST SODIUM;SINGULAIR	5565473 FEB 03, 2012	U-228 NCE	FEB 20, 2003
		5565473 *PED AUG 03, 2012	U-228 PED	AUG 20, 2003
020830 002	MONTLUKAST SODIUM;SINGULAIR	5565473 FEB 03, 2012	U-228 I-300	MAR 03, 2003
		5565473 *PED AUG 03, 2012	U-228 NS	MAR 03, 2003
			NCE	FEB 20, 2003
			PED	SEP 03, 2003
			PED	AUG 20, 2003
			PED	SEP 03, 2003
			PED	APR 27, 2004
			NCE	DEC 10, 2004
			NDF	NOV 30, 2004
			I-329	APR 27, 2004
			PC.	FEB 23, 2002
			PC	FEB 16, 2002
			PC	FEB 23, 2002
021085 001	MOXIFLOXACIN HYDROCHLORIDE;AVELOX	4420639 DEC 13, 2002		
021277 001	MOXIFLOXACIN HYDROCHLORIDE;AVELOX IV	4420639 *PED JUN 13, 2003		
075179 001	NABUMETONE;NABUMETONE	4420639 DEC 13, 2002		
075189 001	NABUMETONE;NABUMETONE	4420639 *PED JUN 13, 2003		
075189 002	NABUMETONE;NABUMETONE	RE34878 MAR 28, 2006		
019583 001	NABUMETONE;RELAFEN	5443116 OCT 21, 2012		
019583 002	NABUMETONE;RELAFEN	5448150 JAN 30, 2013		
021204 001	NATEGLINIDE;STARLIX	RE34878 MAR 28, 2006		
021204 002	NATEGLINIDE;STARLIX	5443116 OCT 21, 2012		
020920 001	NESSRITIDE;NATRECOR	5488150 JAN 30, 2013		
020165 004	NICOTINE;NICODERM CQ	5114923 MAY 19, 2009		
020165 005	NICOTINE;NICODERM CQ	5674710 OCT 07, 2014		
020165 006	NICOTINE;NICODERM CQ	6115497 JUN 14, 2008	U-388	
075269 002	NIFEDIPINE;NIFEDIPINE	5633008 JUN 14, 2008	U-389	
019667 001	OCTREOTIDE ACETATE; SANDOSTATIN	5633008 JUN 14, 2008	U-389	
019667 002	OCTREOTIDE ACETATE; SANDOSTATIN	5633008 JUN 14, 2008	U-388	
019667 003	OCTREOTIDE ACETATE; SANDOSTATIN	6165497 JUN 14, 2008	U-388	
019667 004	OCTREOTIDE ACETATE; SANDOSTATIN	5633008 JUN 14, 2008	U-389	
019667 005	OCTREOTIDE ACETATE; SANDOSTATIN	6165497 JUN 14, 2008	U-388	
021008 001	OCTREOTIDE ACETATE; SANDOSTATIN LAR	5633008 JUN 14, 2008	U-389	
021008 002	OCTREOTIDE ACETATE; SANDOSTATIN LAR	6165497 JUN 14, 2008	U-388	
021008 003	OCTREOTIDE ACETATE; SANDOSTATIN LAR	5753618 JUL 08, 2008	PC	JUN 05, 2001

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020799 001	OFLOXACIN; FLOXIN	5401741	MAR 27, 2012		U-407	
020592 001	OLANZAPINE; ZYPREXA	6251895	SEP 23, 2017			
020592 002	OLANZAPINE; ZYPREXA	6231895	SEP 23, 2017			
020592 003	OLANZAPINE; ZYPREXA	6231895	SEP 23, 2017			
020592 004	OLANZAPINE; ZYPREXA	6251895	SEP 23, 2017			
020592 005	OLANZAPINE; ZYPREXA	5229382	APR 23, 2011	U-149 NCE	SEP 30, 2001	
020592 006	OLANZAPINE; ZYPREXA	5605897	FEB 25, 2014	U-176		
021086 001	OLANZAPINE; ZYPREXA ZYDIS	6251895	SEP 23, 2017			
021086 002	OLANZAPINE; ZYPREXA ZYDIS	6231895	SEP 23, 2017			
021086 003	OLANZAPINE; ZYPREXA ZYDIS	5229382	APR 23, 2011	U-149		
021086 004	OLANZAPINE; ZYPREXA ZYDIS	5605897	FEB 25, 2014	U-176		
020688 001	OLOPATADINE HYDROCHLORIDE; PATANOL	6020487	SEP 23, 2017			
019810 001	OMEPRAZOLE; PRILOSEC	6251895	SEP 23, 2017			
019810 002	OMEPRAZOLE; PRILOSEC	5641805	JUN 06, 2015	U-184	PED	DEC 29, 2001
		6150380	NOV 10, 2018			
		6147103	OCT 09, 2018			
		6166213	OCT 09, 2018			
		6191148	OCT 09, 2018			
		4255131*PED	OCT 05, 2001			
		4636199*PED	JAN 30, 2006			
		4786505*PED	OCT 20, 2007			
		4853130*PED	OCT 20, 2007			
		5093142*PED	AUG 02, 2010			
		5599194*PED	AUG 04, 2014			
		5629305*PED	AUG 04, 2014			
		6147103*PED	APR 09, 2019			
		6150380*PED	MAY 10, 2019			
		6166213*PED	APR 09, 2018			
		6191148*PED	APR 09, 2019			
		4508905	APR 02, 2002			
		4508905*PED	OCT 02, 2002			
		6150380	NOV 10, 2018			
		6147103	OCT 09, 2018			
		6166213	OCT 09, 2018			
		6191148	OCT 09, 2018			
		4255131*PED	OCT 05, 2001			
		4636199*PED	JAN 30, 2006			
		4786505*PED	OCT 20, 2007			
		4853130*PED	OCT 20, 2007			
		5093142*PED	AUG 02, 2010			
		5599194*PED	AUG 04, 2014			
		5629305*PED	AUG 04, 2014			
		6147103*PED	APR 09, 2019			
		6150380*PED	MAY 10, 2019			
		6166213*PED	APR 09, 2018			
		6191148*PED	APR 09, 2019			
		4508905	APR 02, 2002			
		4508905*PED	OCT 02, 2002			
		6150380	NOV 10, 2018			
		6147103	OCT 09, 2018			
		6166213	OCT 09, 2018			
		6191148	OCT 09, 2018			
		4255131*PED	OCT 05, 2001			
		4636199*PED	JAN 30, 2006			
		4786505*PED	OCT 20, 2007			
		4853130*PED	OCT 20, 2007			
		5093142*PED	AUG 02, 2010			
		5599194*PED	AUG 04, 2014			
		5629305*PED	AUG 04, 2014			
		6147103*PED	APR 09, 2019			
		6150380*PED	MAY 10, 2019			
		6166213*PED	APR 09, 2018			
		6191148*PED	APR 09, 2019			
		4508905	APR 02, 2002			

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	USE CODE	EXCLUSIVITY CODE	EXCLUSIVITY EXPIRES
019810 003	OMEPRAZOLE; PRILOSEC	4508905*PED 6150380 6147103 6166213 6191448	OCT 02, NOV 10, OCT 09, NOV 10, OCT 09,	2002 2018 2018 2018 2018	I-229 PED	JUN 29, 2001 DEC 29, 2001
020781 001 021246 001	ONDANSETRON; ZOFTRAN ODT OSELTAMIVIR PHOSPHATE; TAMIFLU	4255431*PED 4636499*PED 4786505*PED 4853230*PED 503342*PED	OCT 05, JAN 30, OCT 20, OCT 20, AUG 02,	2001 2006 2007 2007 2010	U-108	
020897 001	OXYBUTYNIN CHLORIDE; DITROPAN XL	5999794*PED 5629305*PED 6147103*PED	AUG 04, AUG 04, APR 09,	2014 2014 2019	U-166 U-188	
020897 002	OXYBUTYNIN CHLORIDE; DITROPAN XL	6150380*PED 6166213*PED	MAY 10, APR 09,	2019 2018		
020897 003	OXYBUTYNIN CHLORIDE; DITROPAN XL	6191448*PED 4508905	APR 09, APR 02,	2019 2002		
020897 004	OXYCODONE HYDROCHLORIDE; OXYCONTIN	4508905*PED 4753789 5763483 5866601 5952375	OCT 02, JUN 24, DEC 27, FEB 02, FEB 02,	2002 2006 2016 2016 2016	U-330 I-317 NDF NCE	NOV 17, 2003 DEC 14, 2003 OCT 27, 2004
020553 005	OXYCODONE HYDROCHLORIDE; OXYCONTIN	6124355 6262115 6124355 6262115 6124355 6262115	MAY 22, MAY 22, MAY 22, MAY 22, MAY 22, MAY 22,	2015 2015 2015 2015 2015 2015	U-378 U-393 U-378 U-393 U-378 U-393	
020262 001	PACLITAXEL; TAXOL	4861598 4970075 5266331 5549912 5508042 5656295 6096331	AUG 29, NOV 13, FEB 05, FEB 05, FEB 05, FEB 22,	2006 2007 2008 2008 2008 2013		
020036 001 020036 003 020036 004 075290 001	PAMDIDRONATE DISODIUM; AREDIA PAMDIDRONATE DISODIUM; AREDIA PAMDIDRONATE DISODIUM; AREDIA PAMDIDRONATE DISODIUM; PAMIDRONATE DISODIUM	5266331 5549912 5508042 5656295 6096331	FEB 05, FEB 05, APR 16, FEB 05, FEB 22,	2008 2008 2013 2008 2013	D-68 D-68 D-68 D-68 PC	AUG 20, 2004 AUG 20, 2004 AUG 20, 2004 MAY 05, 2002

PRESCRIPTION AND OTC DRUG PRODUCT

PATENT AND EXCLUSIVITY DATA

* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUSIVITY CODE	EXCLUSIVITY EXPIRES	
075290 003 020987 001 020988 001	PAMIDRONATE DISODIUM;PAMIDRONATE DISODIUM PANTOPRAZOLE SODIUM;PROTONIX PANTOPRAZOLE SODIUM;PROTONIX IV	4758579	JUL 19, 2005	PC I-330 NDF NCE	MAY 05, JUN 12, MAR 22, FEB 02,	2002 2004 2004 2005	
020819 001 020031 001 020031 002 020031 003 020031 004 020031 005 020936 003	PARICALCITOL; ZEMPLAR PAROXETINE HYDROCHLORIDE; PAXIL PAROXETINE HYDROCHLORIDE; PAXIL PAROXETINE HYDROCHLORIDE; PAXIL PAROXETINE HYDROCHLORIDE; PAXIL PAROXETINE HYDROCHLORIDE; PAXIL CR	5246925	APR 17, 2012	U-314 I-337	I-326 I-326 I-326 I-326 I-326	APR 13, APR 13, APR 13, APR 13, APR 13,	2004 2004 2004 2004 2004
021064 001	PERFLUTREN,DEFINITY						
021073 001	PIOGLITAZONE HYDROCHLORIDE; ACTOS						
021073 002	PIOGLITAZONE HYDROCHLORIDE; ACTOS						
021073 003	PIOGLITAZONE HYDROCHLORIDE; ACTOS						

PREScription AND oTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
PED and PED represent Pediatric Exclusivity

PED and PED represent Pediatric Exclusivity

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
*PED and PED represent Pediatric Exclusivity

APPL./PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	USE CODE	EXCLUS CODE	EXCLUS EXPRES
021071 004 020692 001	ROSTIGLITAZONE MALEATE; AVANDIA SALMETEROL XINAFOATE; SEREVENT	6288095 5290815 5196438	FEB 11, MAR 01, NOV 19,	2017 2011 2010	U-420 U-386	M-11 M-11 M-11	AUG 06, AUG 06, AUG 06,
020828 001 019839 001	SAQUINAVIR; FORTOVASE SERTALINE HYDROCHLORIDE; ZOLOFT						2004 2004 2004
019839 002 019839 005	SERTALINE HYDROCHLORIDE; ZOLOFT SERTALINE HYDROCHLORIDE; ZOLOFT						AUG 06, AUG 06,
020990 001 020478 001	SERTALINE HYDROCHLORIDE; ZOLOFT SEVOFLURANE; ULTRANE	6288127 6288127*PED	JAN 27, JUL 27,	2017 2018	D-65	FEB 16, FEB 16,	2004 2004
020632 001	SIBUTRAMINE HYDROCHLORIDE; MERIDIA					M-9	FEB 16,
020632 002	SIBUTRAMINE HYDROCHLORIDE; MERIDIA					D-65	2004
020632 003	SIBUTRAMINE HYDROCHLORIDE; MERIDIA					M-9	2004
021097 001 020280 006	SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; VISICOL SOMATROPIN RECOMBINANT; GENOTROPIN	5616346 5633352	MAY 18, MAY 27,	2013 2014	U-359	JUL 25, JUL 25,	2004 2008
020280 007	SOMATROPIN RECOMBINANT; GENOTROPIN	5633352	MAY 27,	2014	I-334	JUL 25,	2004
020280 001	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	6152897 5633352	JUN 11, MAY 27,	2018 2014	ODE JUL 25,	JUL 25,	2004 2008
020280 002	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	6152897 5633352	JUN 11, MAY 27,	2018 2014	JUL 25, <td>JUL 25,</td> <td>2004 2008</td>	JUL 25,	2004 2008
020280 003	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	6152897 5633352	JUN 11, MAY 27,	2018 2014	JUL 25, <td>JUL 25,</td> <td>2004 2008</td>	JUL 25,	2004 2008
020280 004	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	5633352	MAY 27,	2014	JUL 25, <td>JUL 25,</td> <td>2004 2008</td>	JUL 25,	2004 2008
020280 005	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	6152897 5633352	JUN 11, MAY 27,	2018 2014	JUL 25, <td>JUL 25,</td> <td>2004 2008</td>	JUL 25,	2004 2008
020280 008	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	6152897 5633352	JUN 11, MAY 27,	2018 2014	JUL 25, <td>JUL 25,</td> <td>2004 2008</td>	JUL 25,	2004 2008
020280 009	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	6152897 5633352	JUN 11, MAY 27,	2018 2014	JUL 25, <td>JUL 25,</td> <td>2004 2008</td>	JUL 25,	2004 2008
020280 010	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	5633352 5633352	MAY 27,	2014	ODE JUL 25,	JUL 25,	2004 2008
020280 011	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	5633352	MAY 27,	2014	I-334	JUL 25,	2004
020280 012	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	5633352	MAY 27,	2014	ODE NP	JUL 25, FEB 22,	2004 2003
020280 013	SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVAT	5633352	MAY 27,	2014	I-334	JUL 25,	2004
021151 001	SOTALOL HYDROCHLORIDE; BETAPACE AF				PED	AUG 22,	

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
* PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL USE CODE	EXCLUS CODE	EXCLUS EXPIRES
021151 002	SOTALOL HYDROCHLORIDE; BETAPACE AF				NP	FEB 22, 2003
021151 003	SOTALOL HYDROCHLORIDE; BETAPACE AF			PED	AUG 22,	2003
020412 001	STAVUDINE; ZERIT	4978655	JUN 24, 2008	NP	FEB 22,	2003
020412 002	STAVUDINE; ZERIT	4978655*PED	DEC 24, 2008	PED	AUG 22,	2003
020412 003	STAVUDINE; ZERIT	4978655	JUN 24, 2008	NP	FEB 22,	2003
020412 004	STAVUDINE; ZERIT	4978655*PED	DEC 24, 2008	PED	AUG 22,	2003
020412 005	STAVUDINE; ZERIT	4978655	JUN 24, 2008	NP	FEB 22,	2003
021184 002	TAZAROTENE; TAZORAC	4978655*PED	DEC 24, 2008	NP	FEB 22,	2003
021356 001	TENOFOVIR DISPROXIL FUMARATE; VIREAD	5977039	JUL 25, 2017	U-248	NCE	OCT 26, 2006
		6043230	JUL 25, 2017	U-248		
		5935946	JUL 25, 2017	U-248		
		4808716	APR 25, 2006	U-248		
		6057305	MAY 02, 2017	U-248		
		5922655	JUL 25, 2017	U-248		
		4358449	NOV 09, 2001			
		5674711	AUG 31, 2010			
		5602006	FEB 11, 2014			
		5658760	AUG 19, 2014			
		5240832	AUG 31, 2010			
		6114144	NOV 24, 2015			
		5840566	NOV 24, 2015			
		4861760	SEP 25, 2006			
		4861760	SEP 25, 2006			
019964 001	TERCONAZOLE; TERAZOL 3	5559269	NOV 05, 2013	U-318	NCE	NOV 27, 2001
020898 001	THYROTROPIN ALFA; THYROGEN	5559269	NOV 05, 2013	U-318		
020330 001	TIMOLOL MALEATE; TIMOPTIC-XE	4861760	SEP 25, 2006			
020330 002	TIMOLOL MALEATE; TIMOPTIC-XE	4861760	SEP 25, 2006			
020397 002	TIZANIDINE HYDROCHLORIDE; ZANAFLEX					
020771 001	TOLTERODINE TARTRATE; DETROL					
020771 002	TOLTERODINE TARTRATE; DETROL					
020505 001	TOPIRAMATE; TOPAMAX					
020505 002	TOPIRAMATE; TOPAMAX					
020505 003	TOPIRAMATE; TOPAMAX					
020505 004	TOPIRAMATE; TOPAMAX					
020505 005	TOPIRAMATE; TOPAMAX					

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME : TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL CODE	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020505 006	TOPIRAMATE; TOPAMAX					ODE	AUG 28, 2008
020844 001	TOPIRAMATE; TOPAMAX SPRINKLE					J-335	AUG 28, 2004
020844 002	TOPIRAMATE; TOPAMAX SPRINKLE					ODE	AUG 28, 2008
020844 003	TOPIRAMATE; TOPAMAX SPRINKLE					I-335	AUG 28, 2004
020528 001	TRANSDOLAPRIL; MAVIK	4933361	JUN 12, 2007			ODE	AUG 28, 2008
020528 002	TRANSDOLAPRIL; MAVIK	4933361	JUN 12, 2007			ODE	AUG 28, 2004
020528 003	TRANSDOLAPRIL; MAVIK	4933361	JUN 12, 2007			I-335	AUG 28, 2004
020591 001	TRANSDOLAPRIL; TARKA	5721244	FEB 24, 2015			ODE	AUG 28, 2008
020591 002	TRANSDOLAPRIL; TARKA	5721244	FEB 24, 2015			I-335	AUG 28, 2004
020591 003	TRANSDOLAPRIL; TARKA	5721244	FEB 24, 2015			ODE	AUG 28, 2008
020591 004	TRANSDOLAPRIL; TARKA	5721244	FEB 24, 2015			I-335	AUG 28, 2004
021257 001	TRAVODROST; TRAVATAN	6011062	DEC 22, 2014	U-382	NCE	MAR 16, 2006	
		5631287	DEC 22, 2014	U-382			
		5849792	DEC 22, 2014	U-382			
		5889032	AUG 03, 2013	U-383			
		6235781	JUN 15, 2019	U-382			
019963 001	TRETINOIN; RENOVA	RE36068	JUL 29, 2003	U-131			
021108 001	TRETINOIN; RENOVA	RE36068	JUL 29, 2003	U-131			
020475 001	TRETINOIN; RETIN-A MICRO	4603146	JUL 29, 2003	U-131			
020468 001	TRIACINOLONE ACETONIDE/NASACORT AQ	5955109	SEP 21, 2016	U-134			
021288 001	TRIPTORELIN PAMOATE; TRELSTAR	6143319	JUL 03, 2016				
020715 001	TRIPTORELIN PAMOATE; TRELSTAR DEPOT	5225705	JUL 20, 2010			NP	JUN 29, 2004
020759 001	TROVAFLOXACIN MESYLATE; TROVAN	5192741	MAR 09, 2010			NCE	JUN 15, 2005
020759 002	TROVAFLOXACIN MESYLATE; TROVAN	5776885	JUL 07, 2015				
020586 002	UREA, C-13; BREATHTEK UBT FOR H-	6187341	JAN 20, 2019				
		6187341	JAN 20, 2019				
		4830010	OCT 27, 2009	U-147			
		514093	AUG 24, 2009	U-147			
019415 004	UROFOLLITROPIN; FERTINEX	5767067	JUN 16, 2015				
		4845077	JUL 04, 2006	U-408			
019415 005	UROFOLLITROPIN; FERTINEX	4725579	FEB 21, 2005	U-408			
		5767067	JUN 16, 2015				
		4845077	JUL 04, 2006	U-408			
		4725579	FEB 21, 2005	U-408			
020550 001	VALACYCLOVIR HYDROCHLORIDE; VALTREX	6083953	JUL 28, 2014	U-384	D-67	JUN 25, 2004	
020550 002	VALACYCLOVIR HYDROCHLORIDE; VALTREX	5399578	MAR 21, 2012	NE	D-67	JUN 25, 2004	
021304 001	VALGANCICLOVIR HYDROCHLORIDE; VALCYTE			MAR 29, 2004			
021283 001	VALSARTAN; DIOVAN			DEC 23, 2001			

PREScription AND OTC DRUG PRODUCT
 PATENT AND EXCLUSIVITY DATA
 * PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXPIRES	EXCL USE CODE	EXCLUS CODE	EXCLUS EXPIRES	
021283 002	VALSARTAN;DIOVAN	5399578	MAR 21, 2012	NCE	DEC 23,	2001	
021283 003	VALSARTAN;DIOVAN	5399578	MAR 21, 2012	NCE	DEC 23,	2001	
020151 001	VENLAFAXINE HYDROCHLORIDE;EFFEXOR			I-325	MAY 02,	2004	
020151 002	VENLAFAXINE HYDROCHLORIDE;EFFEXOR			I-325	MAY 02,	2004	
020151 003	VENLAFAXINE HYDROCHLORIDE;EFFEXOR			I-325	MAY 02,	2004	
020151 004	VENLAFAXINE HYDROCHLORIDE;EFFEXOR			I-325	MAY 02,	2004	
020151 005	VENLAFAXINE HYDROCHLORIDE;EFFEXOR			I-325	MAY 02,	2004	
020151 006	VENLAFAXINE HYDROCHLORIDE;EFFEXOR			I-325	MAY 02,	2004	
020659 001	VENLAFAXINE HYDROCHLORIDE;EFFEXOR XR	6274171	MAR 20, 2017	U-398	I-325	MAY 02,	2004
020659 002	VENTLAFAXINE HYDROCHLORIDE;EFFEXOR XR	5916923	JUN 28, 2013	U-398	I-325	MAY 02,	2004
020659 003	VENTLAFAXINE HYDROCHLORIDE;EFFEXOR XR	6274171	MAR 20, 2017	U-398	I-325	MAY 02,	2004
020659 004	VENTLAFAXINE HYDROCHLORIDE;EFFEXOR XR	5916923	JUN 28, 2013	U-398	I-325	MAY 02,	2004
021119 001	VERTEPORFIN;VISUDYNE	5916923	JUN 28, 2013	U-398	I-325	MAY 02,	2004
020547 001	ZAFIRLUKAST;ACCOLATE			I-328	SEP 17,	2002	
020547 003	ZAFIRLUKAST;ACCOLATE			I-328	SEP 17,	2002	
020859 001	ZALEPLON;SONATA			M-8	FEB 22,	2004	
020859 002	ZALEPLON;SONATA			M-8	FEB 22,	2004	
020825 001	ZIPRASIDONE HYDROCHLORIDE;GEODON	4831031	MAR 02, 2007	NCE	FEB 05,	2006	
020825 002	ZIPRASIDONE HYDROCHLORIDE;GEODON	5312925	SEP 01, 2012				
		4831031	MAR 02, 2007	NCE	FEB 05,	2006	
		5312925	SEP 01, 2012				
020825 003	ZIPRASIDONE HYDROCHLORIDE;GEODON	5312925	MAR 02, 2007	NCE	FEB 05,	2006	
020825 004	ZIPRASIDONE HYDROCHLORIDE;GEODON	4831031	MAR 02, 2007	NCE	FEB 05,	2006	
021223 001	ZOLEDRONIC ACID;ZOMETA	5312925	SEP 01, 2012				
021231 001	ZOLMITRIPTAN;ZOMIG-ZMT	4939130	NOV 13, 2007	U-53	NCE	AUG 20,	2006
		4777163	JUL 24, 2007	ODE	AUG 20,	2008	
				NDF	FEB 13,	2004	

PATENT AND EXCLUSIVITY TERMS

DUE TO SPACE LIMITATIONS IN THE PATENT AND EXCLUSIVITY COLUMNS, ABBREVIATIONS AND REFERENCES HAVE BEEN DEVELOPED. PLEASE REFER BACK TO THE APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 21ST EDITION FOR A FULL LISTING OF PATENT AND EXCLUSIVITY TERMS (ABBREVIATIONS, NEW DOSING SCHEDULE, NEW INDICATIONS AND PATENT USE CODES). THE CUMULATIVE SUPPLEMENT WILL LIST NEW CODES ADDED SINCE THE LAST ANNUAL EDITION.

ABBREVIATIONS

REFERENCES NEW DOSING SCHEDULE

- D-47 PREVENTION OF HEARTBURN SYMPTOMS WHEN ADMINISTERED FROM 15 MINUTES UP TO, BUT NOT INCLUDING, 1 HOUR PRIOR TO A PROVOCATIVE MEAL
- D-65 CHANGE DOSING AND ADMINISTRATION TO INDICATE MAINTENANCE OF WEIGHT LOSS OVER AN 18 MONTH PERIOD THUS EXTENDING THE USE OF THIS DRUG FROM ONE TO TWO YEARS
- D-66 DOSING RECOMMENDATIONS FOR PATIENTS UNDERGOING PCI
- D-67 SHORTER TREATMENT COURSE OF THREE DAYS IN THE TREATMENT OF RECURRENT EPISODES OF GENITAL HERPES
- D-68 CHANGE OF ADMIN RATE FOR INFUSION OF AREDIA FOR TREATMENT OF MODERATE AND SEVERE HYPERCALCEMIA OF MALIGNANCY FROM 24 HOURS TO 2 HOURS UP TO BUT NOT INCLUDING 24 HOURS
- D-69 SHORTENED DOSING REGIMENT TO 5 DAYS FOR THE TREATMENT OF ACUTE EXACERBATION OF CHRONIC BRONCHITIS

NEW INDICATION

- I-321 JUVENILE RHEUMATOID ARTHRITIS
- I-322 USE OF DIPRIVAN IN PATIENTS 3 MONTHS TO 16 YEARS
- I-323 COLORECTAL CANCER
- I-324 REDUCING NEUROLOGIC DISABILITY AND/OR FREQUENCY OF CLINICAL RELAPSES IN PATIENTS WITH SECONDARY (CHRONIC) PROGRESSIVE, PROGRESSIVE RELAPSING, OR WORSENING RELAPSING-REMITTING MULTIPLE SCLEROSIS
- I-325 PREVENTION OF RELAPSE AND RECURRENCE OF DEPRESSION
- I-326 GENERALIZED ANXIETY DISORDER
- I-327 SYMPTOMATIC RELIEF OF RHINOARRHEA ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN PATIENTS 5 YEARS AND OLDER
- I-328 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 5-6 YEARS OF AGE
- I-329 UNCOMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS
- I-330 MAINTENANCE OF HEALING OF EROSIVE ESOPHAGITIS AND CONTROL OF DAYTIME AND NIGHTTIME HEARTBURN SYMPTOMS IN PATIENTS WITH GERD
- I-331 TREATMENT OF MODERATE ACNE VULGARIS
- I-332 EMPIRIC THERAPY IN FEBRILE NEUTROPENIC PATIENTS WITH SUSPECTED FUNGAL INFECTIONS (ETFN)
- I-333 TOPICAL TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR DUE TO MALASSEZIA FURFUR (FORMERLY PITYROSPORUM ORBICULARE)
- I-334 LONG-TERM TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE WHO FAIL TO MANIFEST CATCH-UP GROWTH BY TWO YEARS OF AGE
- I-335 ADJUNCTIVE THERAPY IN PATIENTS TWO YEARS AND OLDER WITH SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME

PATENT AND EXCLUSIVITY TERMS

REFERENCES NEW INDICATION

- I-336 EXPANSION OF INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH PREDOMINATELY CLASSIC SUBFOVEAL CHOROIDAL NEOVASCULARIZATION DUE TO PATHOLOGIC MYOPIA OR PRESUMED OCULAR HISTOPLASMOSIS
- I-337 PATHOLOGICAL HYPERSECRETION ASSOCIATED WITH ZOLLINGER-ELLISON SNYDROME
- I-338 MANAGEMENT OF ACUTE PAIN IN ADULTS AND TREATMENT OF PRIMARY DYSMENORRHEA
- I-339 TREATMENT OF HEPATITIS B IN PEDIATRIC PATIENTS AGES 2-17 YEARS
- I-340 ATOPIC DERMATITIS IN PEDIATRIC PATIENTS AGES 2-5
- I-341 BREAST CANCER COMBINATION THERAPY
- I-342 USE OF FORADIL FOR LONG-TERM, TWICE DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHO-CONSTRICKTION IN PATIENTS WITH COPD INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-343 USE OF COREG FOR SEVERE HEART FAILURE
- I-344 ACNE VULGARIS

MISCELLANEOUS EXCLUSIVITY CODES

- M-6 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH GLUOPHAGE/GLYBURIDE COMBINATION ADDED TO CLIN PHARM AND DOSING AND ADMIN
- M-7 CLINICAL PHARMACOLOGY IN PEDIATRIC PATIENTS; DOSAGE AND ADMINISTRATION INFORMATION
- M-8 ADDITIONAL INFORMATION FOR THE USE OF SONATA CAPSULES FOR UP TO 5 WEEKS (35 NIGHTS) OF TREATMENT IN A CONTROLLED TRIAL SETTING
- M-9 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING OF TEXT AND TWO TABLES CONTAINING INFORMATION FOR THE PRESCRIBING PHYSICIAN ON BLOOD PRESSURE, HEART RATE, AND HEART RATE VARIABILITY
- M-10 INFORMATION REGARDING MAINTENANCE OF AN ANTIDEPRESSANT EFFECT UP TO 1 YEAR OF DOSING
- M-11 USE FOR LONG-TERM TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- M-12 NEW LANGUAGE FOR PEDIATRIC USE

PATENT USE CODES

- U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMIN TO PT, 30 MIN PRIOR TO CONSUMPTION BY THE PT THE FOOD/BEVERAGE, A COMPOSITION COMPRISING 10MG FAMOTIDINE
- U-372 METHOD FOR ADMINISTERING A BENEFICIAL DRUG TO THE GI TRACT OF AN ANIMAL, WHICH METHOD COMPRISES ADMITTING AN OSMOTIC DEVICE ORALLY INTO THE ANIMAL...
- U-373 GENERAL USE CLAIM SUBMITTED FOR 12 NEXIUM PATIENTS STATING "PERTINENT TO THE CAPSULE FORMULATION FOR NEXIUM AND ITS INDICATIONS FOR THE TREATMENT OF GERD AND ERADICATION OF H.PYLORI TO REDUCE THE RISK OF DUODENAL ULCER RECURRENT"
- U-374 KIT ADAPTED AND DESIGNED TO PROVIDE BOTH DATA ON THE CURRENT REPRODUCTIVE STATUS OF A PATIENT AND CONTRACEPTION FOR THOSE WHO ARE NOT PREGNANT, BUT RECENTLY ENGAGED IN UNPROTECTED SEX
- U-375 METHOD OF USING RIBAVIRIN FOR TREATING A DISEASE RESPONSIVE TO RIBAVIRIN, E.G. HEPATITIS C
- U-376 TREATMENT OF INFLUENZA
- U-377 METHOD OF TREATING PT WITH CHRONIC HEPATITIS C HAVING HCV GENOTYPE 1 AND VIRAL LOAD GREATER THAN 2 MILLION COPIES/ML TO ERADICATE DETECTABLE HCV-RNA BY ADMIN COMBINATION OF RIBAVIRIN AND INTERFERON ALFA-2B FOR A LEAST 24 WEEKS
- U-378 METHOD FOR TREATING INCONTINENCE
- U-379 METHOD OF TREATINGONYCHROMYCOSIS
- U-380 COMBINATIONS OF TAXOL (PACLITAXEL) AND CISPLATIN WHICH ARE SUITABLE FOR THE TREATMENT OF OVARIAN AND NON-SMALL CELL LUNG CARCINOMAS
- U-381 TREATMENT OF HYPERPHOSPHATEMIA
- U-382 METHOD OF STABILIZING PROSTAGLANDIN
- U-383 METHOD FOR TREATING GLAUCOMA AND OCULAR HYPERTENSION

PATENT AND EXCLUSIVITY TERMS

REFERENCES PATENT USE CODES

- U-384 TREATMENT OF CMV RETINITIS
- U-385 TREATMENT OF PEPTIC ULCERS
- U-386 TREATMENT OF PATIENTS SUFFERING FROM A LATE ASTHMATIC REACTION OR LATE PHASE ASTHMA
- U-387 TREATMENT OF PATIENTS WITH RESPIRATORY DISORDERS
- U-388 SMOKING CESSATION AID APPLIED TO THE SKIN
- U-389 SMOKING CESSATION AID APPLIED TO THE SKIN ON WAKING AND REMOVED PRIOR TO SLEEP AFTER ABOUT 16 HOURS
- U-390 METHOD OF USING THE DRUG TO TREAT NEUROIMMUNOLOGIC DISEASES (INCLUDING MULTIPLE SCLEROSIS)
- U-391 USE OF CASODEX IN COMBINATION WITH LHRH AGONISTS FOR THE TREATMENT OF PROSTATE CANCER
- U-392 TREATMENT OF PATIENTS FOR INFLAMMATION
- U-393 MANAGEMENT OF INCONTINENCE, MGT OF HORMONE REPLACEMENT THERAPY, TREATMENT OF INVOLUNTARY INCONTINENCE, MGT OVERACTIVE BLADDER AND INCREASING COMPLIANCE IN SUCH PT
- U-394 METHOD OF USE OF ALPHAGAN
- U-395 METHOD OF USE OF ALPHAGAN P
- U-396 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION
- U-397 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION WITHOUT AN INCREASE IN NAUSEA
- U-398 TREATMENT OF GENERALIZED ANXIETY DISORDER
- U-399 IN-THE-EYE USE OF CHLORINE DIOXIDE CONTAINING COMPOSITIONS
- U-400 USE OF RIBAVIRIN TO INCREASE TYPE 1 CYTOKINE RESPONSE AND SUPPRESS TYPE 2 CYTOKINE RESPONSE TO LYMPHOCYTES, INCLUDING METHODS THAT TAKE ADVANTAGE OF SUCH MODULATION TO TREAT INFECTIONS AND INFESTATIONS
- U-401 USE OF LOPINAVIR IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS FOR TREATING HIV INFECTION AND IN COMBO WITH OTHER HIV PROTEASE INHIBITORS
- U-402 TREATMENT OF ACTINIC KERATOSES
- U-403 ANTI-ALLERGIC FOR VARIOUS ALLERGIC DISEASES
- U-404 TREATMENT OF ALLERGIC CONJUNCTIVITIS
- U-405 METHOD OF USE OF LOTRONEX
- U-406 METHOD OF USE OF ATOVAQUONE AND PROGUANIL
- U-407 METHOD OF TREATING OTOPATHY
- U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASING FACTOR ANTAGONIST AND RECRUITING OOCYTES FOR IN-VITRO FERTILIZATION
- U-409 METHOD OF TREATING INFLAMMATION USING DRUG SUBSTANCE
- U-410 METHOD OF REDUCING AMOUNT OF RESPECTIVE ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (INCLUDING PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-411 METHOD OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN PREPARATION
- U-412 TREATMENT OF TYPE 2 DIABETES
- U-413 USE OF THE ACTIVE INGREDIENT FOR INHIBITING THE BIOSYNTHESIS OF CHOLESTEROL AND TREATMENT OF ATHEROSCLEROSIS
- U-414 A METHOD OF TREATING GLYCOMETABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE
- U-415 A METHOD FOR REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS
- U-416 A METHOD FOR REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS
- U-417 COMBINATION USE OF AD-4833 WITH A BIGUANIDE
- U-418 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-419 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE
- U-420 METHOD OF TREATMENT OF TYPE II DIABETES

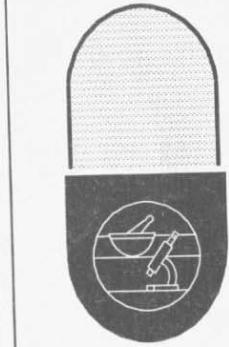
PATENT AND EXCLUSIVITY TERMS

REFERENCES
PATENT USE CODES

- U-421 USE FOR SEDATION
- U-422 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER AND ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-423 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER, ATTENTION DEFICIT HYPERACTIVITY DISORDER, OR AIDS RELATED DEMENTIA
- U-424 FOR ONCE DAILY, BOLUS ADMINISTRATION TO A PATIENT IN ORDER TO ENGENDER TREATMENT FOR A NERVOUS DISORDER FOR SUBSTANTIALLY AN ENTIRE DAY ON A CHRONIC BASIS
- U-425 METHOD OF REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMIN TO A DIABETIC BY ADMIN A CHEMICAL COMPOUND HAVING FORMULA (INCL PIOGLITAZONE) IN COMBINATION WITH AN INSULIN ECRETION ENHANCER
- U-426 PREVENTION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION

*U.S. Government Printing Office: 2002 — 491-214/40005

New 22nd Edition



APPROVED DRUG PRODUCTS

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

22nd EDITION

Superintendent of Documents Subscription Order Form

Order Processing Code

* 8392

**Charge your order.
It's easy!**



Yes, enter my subscription as follows:

Subscriptions of APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, ADP, and the monthly Cumulative Supplements, for \$105.00 per year.

The total cost of my order is \$. Price includes regular shipping and handling and is subject to change. International customers please add 25%.

For privacy protection, check the box below:

Do not make my name available to other mailers.

Please choose method of payment:

Check payable to Superintendent of Documents
 GPO Deposit Account -

VISA or MasterCard

Thank you for your order!

(Credit card expiration date)

Company or personal name

Additional address/attention line

Street address

City, State, ZIP Code

() Daytime phone including area code

Purchase Order No. (optional)

(10/01)

Mail To: Superintendent of Documents, Government Printing Office, P.O. Box 371954 Pittsburgh, PA 15250-7954

To FAX your charge order, call (202) 512-2250.

To charge your subscription call (202) 512-1800.