

CUMULATIVE
SUPPLEMENT 11
JAN'97-NOV'97

APPROVED DRUG PRODUCTS

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

17TH EDITION

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
PUBLIC HEALTH SERVICE
FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF INFORMATION TECHNOLOGY
DIVISION OF DATA MANAGEMENT AND SERVICES

1997

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Suppl



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APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

17TH EDITION

Cumulative Supplement 11

NOVEMBER 1997

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Approved drug products with
therapeutic equivalence
C:355661 M:174736 O:12937927
APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

17TH EDITION

CUMULATIVE SUPPLEMENT 11
NOVEMBER 1997

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, 17th 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and products that have never been marketed, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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.

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. Patent and Exclusivity information for a specific drug is

indicated by an abbreviation followed by the date upon which the patent and/or exclusivity expires. Refer to the Patent and Exclusivity Terms section in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations. Information regarding drug patents and exclusivity for an approved product will appear in this addendum as it is received and may not correspond with the month the drug product is approved and published in the Rx/OTC sections of the Cumulative Supplement.

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.)

New additions to the Prescription Drug Product List, OTC Drug Product List, and 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, OTC Drug Product List, and the Patent and Exclusivity Data are indicated by the symbol >**DLT**> (DELETE) to the left of the line. The >**DLT**> symbol is dropped in subsequent Cumulative Supplements for that item. The shaded print remains in the Prescription Drug Product List and OTC Drug Product Lists in all Cumulative Supplements (hard copy only) for this edition. However, the overstruck data in the Patent and Exclusivity Data is dropped in subsequent Cumulative Supplements (hard copy).

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 17th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 18th Edition.

1.2 COURT ORDER AFFECTING URUGUAY ROUND AGREEMENTS ACT-EXTENDED PATENTS

As a result of the April 4, 1996, decision of the United States Court of Appeals for the Federal Circuit in Merck, et al. v. Kessler, patent expiration dates for certain patents subject to patent term extensions under the Uruguay Round Agreements Act and to the patent term extension provisions at 35 U.S.C. § 156 may be changed. FDA has published a notice in the March 14, 1997, *Federal Register* advising NDA and NADA applicants that patent expiration dates changed by the Merck decision must be submitted within 60 days. Because there may be changes in listed patents as a result of the Merck decision, users of this publication should consult the most recent supplement, and are encouraged to confirm that patent information upon which they intend to rely is current. (See the *Patent and Exclusivity Addendum to the Approved Drug Products with Therapeutic Equivalence Evaluations*, 17th Edition that explains the background information on this court decision).

1.3 APPLICANT NAME CHANGES

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

It is also not practical to identify each and every product involved when an applicant name is changed to meet internal publication standards (e.g., MSD or Zenith [Former Abbreviated Names] are changed, respectively, to Merck Sharp Dohme or Zenith Labs [New Abbreviated Names]). When this occurs, each product involved (either currently in the Cumulative Supplement or in the following year's edition) will automatically 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. However, when the applicant name change is one which may not be easily recognized or located in the listing (e.g., White Towne Paulsen [Former Abbreviated Name] is changed to Whiteworth Towne PLSN [New Abbreviated Name], the name change will appear in this section and will be identified with an asterisk.

APPLICANT NAME CHANGES

FORMER APPLICANT NAME
(FORMER ABBREVIATED NAME)

NEW APPLICANT NAME
(NEW ABBREVIATED NAME)

CIBA GEIGY CORP
(CIBA GEIGY)

NOVARTIS PHARMACEUTICALS CORP
(NOVARTIS)

CIBA GEIGY CORP PHARMACEUTICALS DIV
(CIBA GEIGY)

NOVARTIS PHARMACEUTICALS CORP
(NOVARTIS)

CIBA PHARMACEUTICAL CO
DIV CIBA GEIGY CORP
(CIBA)

NOVARTIS PHARMACEUTICALS CORP
(NOVARTIS)

CIBA SELF MEDICATION INC
DIV CIBA GEIGY CORP
(CIBA)

NOVARTIS CONSUMER HEALTH INC
(NOVARTIS)

CIBA VISION CORP
(CIBA)

CIBA VISION CORPORATION A
NOVARTIS COMPANY
(CIBA)

CIBA VISION OPHTHALMICS
DIV CIBA VISION CORP
(CIBA)

CIBA VISION CORPORATION A
NOVARTIS COMPANY
(CIBA)

FERRING LABORATORIES INC
(FERRING)

FERRING PHARMACEUTICALS INC
(FERRING)

GEIGY PHARMACEUTICALS
DIV CIBA GEIGY CORP
(GEIGY)

NOVARTIS PHARMACEUTICALS CORP
(NOVARTIS)

LEMMON CO SUB TAG PHARMACEUTICAL INC
(LEMMON)

BIOCRAFT LABORATORIES INC
(BIOCRAFT)
THEN CHANGED TO
TEVA PHARMACEUTICALS USA
(TEVA)

REXAR PHARMACAL
(REXAR)

SHIRE RICHWOOD INC
(SHIRE RICHWOOD)

RICHWOOD PHARMACEUTICAL CO INC
(RICHWOOD PHARM)

SHIRE RICHWOOD INC
(SHIRE RICHWOOD)

APPLICANT NAME CHANGES

<u>FORMER APPLICANT NAME (FORMER ABBREVIATED NAME)</u>	<u>NEW APPLICANT NAME (NEW ABBREVIATED NAME)</u>
SANDOZ CONSUMER HEALTH CARE GROUP DIV SANDOZ PHARMACEUTICALS (SANDOZ)	NOVARTIS CONSUMER HEALTH INC (NOVARTIS)
SANDOZ PHARMACEUTICALS CORP DIV SANDOZ INC (SANDOZ)	NOVARTIS PHARMACEUTICALS CORP (NOVARTIS)
SANDOZ RESEARCH INSTITUTE INC (SANDOZ)	NOVARTIS PHARMACEUTICALS CORP (NOVARTIS)
SANOFI WINTHROP INC (SANOFI WINTHROP)	SANOFI PHARMACEUTICAL INC (SANOFI)
SURVIVAL TECHNOLOGY INC (SURVIVAL TECH)	MERIDIAN MEDICAL TECHNOLOGIES INC (MERIDIAN MEDCL TECHN)

1.4 ACYCLOVIR 200MG TABLET-REFERENCE LISTED DRUG

Novopharm's single source acyclovir tablets have been declared to be a reference listed drug for the 200 mg tablet in addition to the acylcovir (Zovirax) 800 mg tablet of the innovator. A generic firm wishing to submit an ANDA for a duplicate of the 200 mg acyclovir tablet will be eligible for a waiver of the *in vivo* determination of bioequivalence (1) if their product is proportionally similar in its active and inactive ingredients to their own 800 mg acyclovir tablet and (2) by doing an acceptable comparative dissolution test (dissolution profile) against Novopharm's 200 mg acyclovir reference listed drug.

Before a waiver of the *in vivo* determination of bioequivalence can be granted for the 200 mg acyclovir tablet, the generic firm must have completed an acceptable fasting and fed study comparing their acyclovir 800 mg tablet against the Zovirax 800 mg tablet.

For further information on the study designs, you should contact the Division of Bioequivalence, Office of Generic Drugs.

1.5 FOLLITROPIN ALFA AND BETA

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

1.6 AVAILABILITY OF THE PUBLICATION AND UPDATING PROCEDURES

The *Approved Drug Products with Therapeutic Equivalence Evaluations* (Prescription Drug Products, OTC Drug Products and the Discontinued Drug Product Lists) is available on diskette, on a quarterly basis, from the National Technical Information Service. The telephone number for the Subscription Department is (703) 487-6430. Written inquiries regarding this subscription may be forwarded to 5285 Port Royal Road Springfield, VA 22161.

The following *Approved Drug Products with Therapeutic Equivalence Evaluations* files are now available on Internet and are updated each October and April: Prescription Drug Product List; OTC Drug Products; Discontinued Drug Products; Prescription and OTC Drug Product Patent and Exclusivity Data; and Appendices. The update in October will include drug products that have been approved through August and the update in April will include drug products that have been approved through December.

These files may be accessed on the Internet's World Wide Web. FDA's Internet site replaced the Agency's electronic bulletin board. To access the CDER Home Page, use this Uniform Resource Locator (URL): <http://www.fda.gov/cder>. You do not need an Internet connection to reach the FDA Home Page; you can use the free dial-up connection (800) 222-0185 for text based, non-graphical use only. For further assistance, please call (301) 443-4908.

1.7 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 1996) 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

CATEGORIES COUNTED	DEC 1996*	MAR 1997	JUN 1997	SEP 1997
DRUG PRODUCTS LISTED	9392	9493	9533	9501
SINGLE SOURCE	2383 (25.4%)	2387 (25.1%)	2388 (25.0%)	2420 (25.5%)
MULTISOURCE	6905 (73.5%)	6991 (73.7%)	7031 (73.8%)	6970 (73.3%)
THERAPEUTICALLY EQUIVALENT	6463 (68.8%)	6549 (69.0%)	6626 (69.5%)	6587 (69.3%)
NOT THERAPEUTICALLY EQUIVALENT	442 (4.7%)	442 (4.7%)	405 (4.3%)	383 (4.0%)
EXCEPTIONS	104 (1.1%)	115 (1.2%)	114 (1.2%)	111 (1.2%)
NEW MOLECULAR ENTITIES APPROVED	--	6	8	14
NUMBER OF APPLICANTS	650	662	682	701

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

*Exceptions were originally included in the total count of the Multisource Drug Products. Beginning with December 1996, exceptions were no longer included in the Multisource Drug Products total count, but included in the total count of the Drug Products Listed.

X

PRESCRIPTION DRUG PRODUCT LIST

17TH EDITION

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN' 97 - NOV' 97

ACARBOSE

TABLET; ORAL
PRECOSE
BAYER
> DLT >
> DLT >
> ADD >
> ADD >

N20482 0.04
MAY 29, 1997
N20482 0.04
MAY 29, 1997

25MG
@

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BUCET
MALLINCKRODT

650MG; 50MG
JUN 28, 1985

BUTALBITAL AND ACETAMINOPHEN
GRAHAM DM

650MG; 50MG
JUN 28, 1985

CONTIN
GRAHAM DM

650MG; 50MG

TENCON
MALLINCKRODT

650MG; 50MG

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

TRIAD
MALLINCKRODT

N889023 001

JUN 19, 1985

TABLET; ORAL

REPAN
GRAHAM DM

N889023 001

JAN 24, 1985

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE
MIKART

N40109 001

AUG 26, 1997

325MG; 50MG; 1.0MG

N88904 001

JAN 24, 1985

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE
DURAMED

N40223 001

NOV 18, 1997

300MG; 15MG

N40223 002

NOV 18, 1997

356 . 4MG; 1.6MG

N40223 003

NOV 18, 1997

356 . 4MG; 1.6MG

N88904 001

MAR 04, 1986

N88904 001

JUN 19, 1985

ACETAMINOPHEN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE
DURAMED

N88904 001

JUL 08, 1982

300MG; 3.0MG

300MG; 6.0MG

N88906 001

JUL 19, 1985

300MG; 5MG

N88906 001

AUG 09, 1985

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE
GRAHAM DM

N88743 001

JUL 18, 1985

325MG; 50MG; 4.0MG

N88765 001

MAR 27, 1985

325MG; 50MG; 4.0MG

N89067 001

APR 19, 1985

325MG; 50MG; 4.0MG

N89102 001

JUN 19, 1985

325MG; 50MG; 4.0MG

N88743 001

JUL 18, 1985

325MG; 50MG; 4.0MG

N88765 001

MAR 27, 1985

325MG; 50MG; 4.0MG

N89066 001

JUN 19, 1985

325MG; 50MG; 4.0MG

N88906 001

AUG 09, 1985

325MG; 50MG; 4.0MG

N89066 001

JUN 19, 1985

325MG; 50MG; 4.0MG

N88743 001

JUL 08, 1982

N88956 001

JUL 19, 1985

325MG; 50MG

N88906 001

AUG 09, 1985

325MG; 50MG

N88906 001

JUN 19, 1985

325MG; 50MG

N88743 001

JUL 18, 1985

N88956 001

JUL 19, 1985

325MG; 50MG

N88906 001

AUG 09, 1985

325MG; 50MG

N88906 001

JUN 19, 1985

325MG; 50MG

N88743 001

JUL 18, 1985

N88956 001

JUL 19, 1985

325MG; 50MG

N88906 001

AUG 09, 1985

325MG; 50MG

N88906 001

JUN 19, 1985

325MG; 50MG

N88743 001

JUL 18, 1985

N88956 001

JUL 19, 1985

325MG; 50MG

N88906 001

AUG 09, 1985

325MG; 50MG

N88906 001

JUN 19, 1985

ACETAMINOPHEN; HYDROCODONE BITARTRATE

<u>ACETAMINOPHEN; HYDROCODONE BITARTRATE</u>						
<u>CAPSULE; ORAL</u>						
<u>AA</u>	<u>MALLINCKRODT</u>	<u>500MG; 5MG</u>				
		N89006 001	AUG 09, 1985	<u>AA</u>	+ UCB	<u>500MG; 10MG</u>
<u>AA</u>	<u>LORCET-HD</u>	<u>500MG; 5MG</u>				
	+ MALLINCKRODT	N87336 001	JUL 08, 1982			325MG; 5MG
<u>AA</u>	<u>ZYDONE</u>	<u>500MG; 5MG</u>				
	MALLINCKRODT	N88956 001	JUL 19, 1985	+ WATSON LABS		325MG; 10MG
<u>TABLET; ORAL</u>						
<u>AA</u>	<u>ACETAMINOPHEN AND HYDROCODONE BITARTRATE</u>	<u>500MG; 5MG</u>				
	GRAHAM DM	N87722 001	JUL 09, 1982			
<u>AA</u>	<u>HYDROCODONE BITARTRATE AND ACETAMINOPHEN</u>	<u>500MG; 5MG</u>				
	® BARR	N88577 001	DEC 21, 1984			
<u>AA</u>	EON	500MG; 5MG	N40149 001			
<u>AA</u>		750MG; 7.5MG	JAN 27, 1997			
<u>AA</u>	HALSEY	500MG; 5MG	N40149 002			
<u>AA</u>		650MG; 7.5MG	JAN 27, 1997			
<u>AA</u>		650MG; 7.5MG	SEP 25, 1997	<u>AA</u>	* DUPONT MERCK	325MG; 5MG
<u>AA</u>		650MG; 10MG	N40236 001	<u>AA</u>	+ ENDO PHARMS	325MG; 5MG
<u>AA</u>		750MG; 7.5MG	NOV 24, 1997			
<u>AA</u>		500MG; 10MG	N40240 002			
<u>AA</u>		500MG; 2.5MG	NOV 26, 1997			
<u>AA</u>		650MG; 7.5MG	N40236 002			
<u>AA</u>		500MG; 10MG	SEP 25, 1997			
<u>AA</u>		500MG; 10MG	N40210 001			
<u>AA</u>		500MG; 2.5MG	AUG 13, 1997			
<u>AA</u>		650MG; 7.5MG	N40144 002			
<u>AA</u>		500MG; 5MG	APR 25, 1997			
<u>AA</u>		500MG; 5MG	N40155 001			
<u>AA</u>		500MG; 5MG	APR 14, 1997			
<u>AA</u>		500MG; 10MG	N40148 002			
<u>AA</u>		500MG; 10MG	FEB 14, 1997			
<u>AA</u>		500MG; 5/ ACETAMINOPHEN				
<u>AA</u>	BARR	500MG; 5MG				
<u>AA</u>	LORTAB	N88577 001	DEC 21, 1984	<u>AB</u>	TARO	12.5MG
<u>AA</u>	* GRAHAM DM	500MG; 10MG		<u>AB</u>		2.50MG
<u>AA</u>	MALLINCKRODT	N40100 001	JAN 26, 1996			
<u>AA</u>		500MG; 5MG	JUL 09, 1982			
<u>CAPSULE; ORAL</u>						
<u>AA</u>	<u>OXYCODEONE AND ACETAMINOPHEN</u>	<u>500MG; 5MG</u>				
	ROYCE LABS	N40234 001	OCT 30, 1997			
<u>CAPSULE; ORAL</u>						
<u>AA</u>	<u>OXYCODEONE AND ACETAMINOPHEN</u>	<u>500MG; 5MG</u>				
	ROYCE LABS	N40171 001	OCT 30, 1997			
<u>TABLET; ORAL</u>						
<u>AA</u>	<u>OXYCODEONE AND ACETAMINOPHEN</u>	<u>325MG; 5MG</u>				
	ROYCE LABS	N88106 002				
		N85106 002				
<u>CAPSULE; ORAL</u>						
<u>AA</u>	<u>PROPOXYPHENE NAPSYLATE</u>	<u>650MG; 100MG</u>				
	VINTAGE PHARMS	N74843 001	FEB 12, 1997			
<u>TABLET; ORAL</u>						
<u>AA</u>	<u>PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN</u>	<u>650MG; 100MG</u>				
	VINTAGE PHARMS	N40195 001	MAY 28, 1997			
<u>TABLET; ORAL</u>						
<u>AA</u>	<u>ACETAZOLAMIDE</u>	<u>12.5MG</u>				
		N40195 002	MAY 28, 1997			
<u>TABLET; ORAL</u>						
<u>AA</u>	<u>ACETAZOLAMIDE</u>	<u>2.50MG</u>				
		N40195 002	MAY 28, 1997			

<u>ACETAZOLAMIDE SODIUM</u>		<u>ACYCLOVIR</u>	
INJECTABLE; INJECTION <u>ACETAZOLAMIDE SODIUM</u>	<u>EQ</u> 500MG BASE/VIAL	N40108 001 OCT 30, 1995 N40108 001 OCT 30, 1995	AB + <u>ZOVIRAX</u> WELLCOME <u>ZOVIRAX</u> WELLCOME 2.00MG
AP ABBOTT			
AP SANOFI WINTHROP	<u>EQ</u> 500MG BASE/VIAL		SUSPENSION; ORAL <u>ACYCLOVIR</u> ALPHARMA 2.00MG/5ML
ACETIC ACID, GLACIAL; DESONIDE			
SOLUTION/DROPS; AURICULAR (OTIC) TRIDESILON ④ BAYER	2% ; 0.05%	N17914 001	AB + <u>ZOVIRAX</u> WELLCOME 2.00MG/5ML
ACYCLOVIR			
SOLUTION/DROPS; OTIC TRIDESILON BAYER	2% ; 0.05%	N17914 001	TABLET; ORAL <u>ACYCLOVIR</u> AESGEN 4.00MG
ACYCLOVIR		> ADD > > ADD > > ADD >	AB AB AB
CAPSULE; ORAL <u>ACYCLOVIR</u> AESGEN	<u>200MG</u>	N74833 001 APR 22, 1997 N74889 001 OCT 31, 1997 N74914 001 NOV 26, 1997 N74872 001 APR 22, 1997 N74750 001 APR 22, 1997 N74828 001 APR 22, 1997 N74727 001 APR 22, 1997 N74578 001 APR 22, 1997 N74906 001 AUG 26, 1997 N74570 002 APR 22, 1997 N74828 001 APR 22, 1997 N74674 001 APR 22, 1997	4.00MG 8.00MG 8.00MG 4.00MG 8.00MG 8.00MG 4.00MG 8.00MG 8.00MG 4.00MG 8.00MG 8.00MG 4.00MG 8.00MG 8.00MG 4.00MG 8.00MG 8.00MG 4.00MG 8.00MG 8.00MG 4.00MG
AB	AB		
AB	AB		
> ADD > > ADD >	AB		
AB	COPLEY PHARM		
AB	ESTI LEDERLE		
AB	LEK PHARM		
AB	LEMON		
AB	MYLAN		
AB	NOVOPHARM		
AB	PUREPAC PHARM		
AB	ZENITH GOLDLINE		
AB	ZENITH GOLDLINE		
AB			

N20089 001
APR 30, 1991

<u>ACYCLOVIR</u>				
TABLET; ORAL <u>ZOVIRAX</u> AB + GLAXO WELLCOME	800MG N20089 002 APR 30, 1991			
<u>ACYCLOVIR SODIUM</u>				
INJECTABLE; INJECTION <u>ACYCLOVIR SODIUM</u> ABBOTT	EQ 500MG BASE/VIAL N74663 001 APR 22, 1997			
AP	EQ 500MG BASE/VIAL N74758 001 APR 22, 1997			
AP	EQ 1GM BASE/VIAL N74663 002 APR 22, 1997			
AP	EQ 1GM BASE/VIAL N74758 002 APR 22, 1997			
AP	EQ 500MG BASE/VIAL N74596 002 APR 22, 1997			
AP	EQ 1GM BASE/VIAL N74596 001 APR 22, 1997			
AP	EQ 500MG BASE/VIAL N74913 001 OCT 15, 1997			
AP	EQ 1GM BASE/VIAL N74913 002 OCT 15, 1997			
AP	EQ 25MG BASE/ML N74720 001 APR 22, 1997			
+ FAULDING	EQ 500MG BASE/VIAL N74969 001 AUG 26, 1997			
AP	GENSIA	EQ 1GM BASE/VIAL N74969 002 AUG 26, 1997		
AP	SANOFI	EQ 500MG BASE/VIAL N74663 001 APR 22, 1997		
AP	ZOVIRAX AB + GLAXO WELLCOME	EQ 1GM BASE/VIAL N74663 002 APR 22, 1997		
<u>ALBUTEROL</u>				
AEROSOL, METERED; INHALATION <u>ALBUTEROL</u> ALPHARMA	AB 0.09MG/INH AUG 19, 1997			
<u>ALBUTEROL SULFATE</u>				
SOLUTION; INHALATION <u>ALBUTEROL SULFATE</u> NEPHRON	AN EQ 0.083% BASE SEP 17, 1997			
<u>ALENDRONATE SODIUM</u>				
TABLET; ORAL FOSAMAX MERCK	EQ 5MG BASE N20560 003 APR 25, 1997			
<u>ALPRAZOLAM</u>				
TABLET; ORAL <u>ALPRAZOLAM</u> ROYCE LABS	AB 0.25MG JAN 21, 1997			
<u>ALPROSTADIL</u>				
INJECTABLE; INJECTION <u>CAVERJECT</u>	AP PHARMacia AND UPJOHN 0.005MG/VIAL N20379 003 JUN 27, 1996			
EQ 1GM BASE/VIAL N18603 002 JUN 29, 1989	AP 0.01MG/VIAL N20379 001 JUL 06, 1995			
AP +	AP 0.02MG/VIAL N20379 002 JUL 06, 1995			
AP +	AP 0.04MG/VIAL N20379 004 MAY 19, 1997			

ALPROSTADILINJECTABLE; INJECTIONCAVERJECT

PHARMACIA AND UPJOHN 0.005MG/ML

0.01MG/ML

0.02MG/ML

0.04MG/ML

0.05MG/ML

0.06MG/ML

0.07MG/ML

0.08MG/ML

0.09MG/ML

0.10MG/ML

0.11MG/ML

0.12MG/ML

0.13MG/ML

0.14MG/ML

0.15MG/ML

0.16MG/ML

0.17MG/ML

0.18MG/ML

0.19MG/ML

0.20MG/ML

0.21MG/ML

0.22MG/ML

0.23MG/ML

0.24MG/ML

0.25MG/ML

0.26MG/ML

0.27MG/ML

0.28MG/ML

0.29MG/ML

0.30MG/ML

0.31MG/ML

0.32MG/ML

0.33MG/ML

0.34MG/ML

0.35MG/ML

0.36MG/ML

0.37MG/ML

0.38MG/ML

0.39MG/ML

0.40MG/ML

0.41MG/ML

0.42MG/ML

0.43MG/ML

0.44MG/ML

0.45MG/ML

0.46MG/ML

0.47MG/ML

0.48MG/ML

0.49MG/ML

0.50MG/ML

0.51MG/ML

0.52MG/ML

0.53MG/ML

0.54MG/ML

0.55MG/ML

0.56MG/ML

0.57MG/ML

0.58MG/ML

AMIKACIN SULFATEINJECTABLE; INJECTIONAMIKACIN SULFATE

SANOFTI MINTHROP

N64098 001

JUN 26, 1995

N64099 001

JUN 20, 1995

N64099 004

JUN 20, 1995

N64146 001

APR 02, 1997

N64146 002

APR 03, 1997

N64146 003

APR 03, 1997

N64146 004

APR 03, 1997

N64146 005

APR 03, 1997

N64146 006

APR 03, 1997

N64146 007

APR 03, 1997

N64146 008

APR 03, 1997

N64146 009

APR 03, 1997

N64146 010

APR 03, 1997

N64146 011

APR 03, 1997

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUMAMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUMAMANTADINE HYDROCHLORIDECAPSULE; ORALSYMADINE

SYNTHAX

100MG

SEP 04, 1986

N71000 001

AMIKACIN SULFATEAMIKACIN SULFATE

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

INJECTABLE; INJECTION AMINOSYN II 4.25% W/ CALCIUM IN PLASTIC CONTAINER ④ ABBOTT	4.25%; 36 .8MG/100ML; 261MG/100ML; 51MG/100ML; 22 .4MG/100ML; 261IMG/100ML; 205MG/100ML	SEP 12, 1988 N19714 002	CLINIMIX E 2.75/10 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	2.75%; 33MG/100ML; 10GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML	MAR 26, 1997 N20678 002
AMINOSYN II 4.25% W/ CALCIUM IN PLASTIC CONTAINER ④ ABBOTT	4.25%; 36 .8MG/100ML; 261MG/100ML; 51MG/100ML; 22 .4MG/100ML; 261IMG/100ML; 205MG/100ML	SEP 12, 1988 N19714 004	CLINIMIX E 2.75/5 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	2.75%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML	MAR 26, 1997 N20678 005
AMINOSYN II 5% W/ IN PLASTIC CONTAINER ④ ABBOTT	5%; 36 .8MG/100ML; 25GM/100ML; 51MG/100ML; 22 .4MG/100ML; 261MG/100ML; 205MG/100ML	SEP 12, 1988 N19714 004	CLINIMIX E 4.25/10 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	4.25%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 217MG/100ML; 112MG/100ML	MAR 26, 1997 N20678 001
AMINOSYN II 5% W/ IN PLASTIC CONTAINER ④ ABBOTT	5%; 36 .8MG/100ML; 25GM/100ML; 51MG/100ML; 22 .4MG/100ML; 261MG/100ML; 205MG/100ML	NOV 07, 1988 N19683 004	CLINIMIX E 4.25/20 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	4.25%; 33MG/100ML; 297MG/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	MAR 26, 1997 N20678 009
AMINOSYN II 5% W/ IN PLASTIC CONTAINER ④ ABBOTT	5%; 36 .8MG/100ML; 25GM/100ML; 51MG/100ML; 22 .4MG/100ML; 261MG/100ML; 205MG/100ML	SEP 12, 1988 N19714 003	CLINIMIX E 4.25/20 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	4.25%; 33MG/100ML; 20GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	MAR 26, 1997 N20678 011
AMINOSYN II 5% W/ IN PLASTIC CONTAINER ④ ABBOTT	5%; 36 .8MG/100ML; 25GM/100ML; 51MG/100ML; 22 .4MG/100ML; 261MG/100ML; 205MG/100ML	NOV 07, 1988 N19683 004	CLINIMIX E 4.25/25 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	4.25%; 33MG/100ML; 20GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	MAR 26, 1997 N20678 012
AMINOSYN II 5% W/ IN PLASTIC CONTAINER ④ ABBOTT	5%; 36 .8MG/100ML; 25GM/100ML; 51MG/100ML; 22 .4MG/100ML; 261MG/100ML; 205MG/100ML	SEP 12, 1988 N19714 003	CLINIMIX E 4.25/5 SULFITE-FREE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE	4.25%; 33MG/100ML; 5GM/100ML; 51MG/100ML; 261MG/100ML; 297MG/100ML; 77MG/100ML	MAR 26, 1997 N20678 008

<u>AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DI BASIC; SODIUM ACETATE; SODIUM CHLORIDE;</u>	<u>AMINO ACIDS; DEXTROSE</u>
INJECTABLE; INJECTION CLINIMIX E 5/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE 5%; 3.3MG/100ML; 10GM/100ML; 51MG/100ML; 26.1MG/100ML; 34.0MG/100ML; 59MG/100ML N20678 016 MAR 26, 1997	INJECTABLE; INJECTION AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER ABBOTT 3 .5%; 5GM/100ML NOV 07, 1986
CLINIMIX E 5/15 SULFITE-FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE 5%; 3.3MG/100ML; 15GM/100ML; 51MG/100ML; 26.1MG/100ML; 34.0MG/100ML; 59MG/100ML N20678 017 MAR 26, 1997	AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER ABBOTT 3 .5%; 5GM/100ML NOV 07, 1986
CLINIMIX E 5/20 SULFITE-FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE 5%; 3.3MG/100ML; 20GM/100ML; 51MG/100ML; 26.1MG/100ML; 34.0MG/100ML; 59MG/100ML N20678 018 MAR 26, 1997	AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER ABBOTT 3 .5%; 5GM/100ML NOV 07, 1986
CLINIMIX E 5/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE 5%; 3.3MG/100ML; 25GM/100ML; 51MG/100ML; 26.1MG/100ML; 34.0MG/100ML; 59MG/100ML N20678 019 MAR 26, 1997	AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER ABBOTT 3 .5%; 20GM/100ML NOV 07, 1986
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER + BAXTER HLTHCARE 5%; 3.3MG/100ML; 35GM/100ML; 51MG/100ML; 26.1MG/100ML; 34.0MG/100ML; 59MG/100ML N20678 021 MAR 26, 1997	AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER ABBOTT 3 .5%; 25GM/100ML NOV 07, 1986
AMINO ACIDS; DEXTROSE	CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER BAXTER HLTHCARE 2 .75%; 10% SEP 09, 1988
INJECTABLE; INJECTION AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER ABBOTT 3 .5%; 25GM/100ML NOV 07, 1986	CLINIMIX 2 .75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER BAXTER HLTHCARE 2 .75%; 25GM/100ML NOV 07, 1986
② 3 .5%; 25GM/100ML SEP 09, 1988	CLINIMIX 2 .75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER BAXTER HLTHCARE 2 .75%; 5GM/100ML NOV 07, 1986
② 3 .5%; 25GM/100ML SEP 09, 1988	N20734 001 SEP 29, 1997
AMINO ACIDS; DEXTROSE	N20734 002 SEP 29, 1997
INJECTABLE; INJECTION AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER ABBOTT 3 .5%; 25GM/100ML NOV 07, 1986	N20734 005 SEP 29, 1997
② 3 .5%; 25GM/100ML SEP 09, 1988	N20734 006 SEP 29, 1997
② 3 .5%; 25GM/100ML SEP 09, 1988	N20734 007 SEP 29, 1997

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER	BAXTER HLTHCARE	4 .25%;10GM/100ML	N20734 008	> DLT >	AMINOSYN II 4 .25% W/ DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 4 .25% W/ DEXTROSE AND ADJUSTED PHOSPHATE IN PLASTIC CONTAINER
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER	BAXTER HLTHCARE	4 .25%;20GM/100ML	N20734 010	> DLT >	AMINOSYN II 4 .25% W/ DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 4 .25% W/ DEXTROSE AND ADJUSTED PHOSPHATE IN PLASTIC CONTAINER
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER	BAXTER HLTHCARE	4 .25%;25GM/100ML	N20734 011	> DLT >	AMINOSYN II 4 .25% W/ DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 4 .25% W/ DEXTROSE AND ADJUSTED PHOSPHATE IN PLASTIC CONTAINER
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;10GM/100ML	N20734 014	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;15GM/100ML	N20734 015	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;20GM/100ML	N20734 016	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;25GM/100ML	N20734 017	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;35GM/100ML	N20734 018	@	AMINOSYN II 3 .5% M IN DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 10% IN PLASTIC CONTAINER

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

INJECTABLE; INJECTION

CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER	BAXTER HLTHCARE	4 .25%;10GM/100ML	N20734 008	> DLT >	AMINOSYN II 4 .25% W/ DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 4 .25% W/ DEXTROSE AND ADJUSTED PHOSPHATE IN PLASTIC CONTAINER
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER	BAXTER HLTHCARE	4 .25%;20GM/100ML	N20734 010	> DLT >	AMINOSYN II 4 .25% W/ DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 4 .25% W/ DEXTROSE AND ADJUSTED PHOSPHATE IN PLASTIC CONTAINER
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER	BAXTER HLTHCARE	4 .25%;25GM/100ML	N20734 011	> DLT >	AMINOSYN II 4 .25% W/ DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 4 .25% W/ DEXTROSE AND ADJUSTED PHOSPHATE IN PLASTIC CONTAINER
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;10GM/100ML	N20734 014	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;15GM/100ML	N20734 015	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;20GM/100ML	N20734 016	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;25GM/100ML	N20734 017	> ADD >	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 5% IN PLASTIC CONTAINER
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER	BAXTER HLTHCARE	5%;35GM/100ML	N20734 018	@	AMINOSYN II 3 .5% M IN DEXTROSE 10% IN PLASTIC CONTAINER	AMINOSYN II 3 .5% M IN DEXTROSE 10% IN PLASTIC CONTAINER

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE	AP * SEARLE	25MG/ML	N87243 001
AMINOPHYLLINE	AP @	25MG/ML	N87243 001
AMINOPHYLLINE	AP EKINS SINK	25MG/ML	N87243 001

AMOXICILLINAMOXIL

CAPSULE; ORAL

SMITHKLINE BEECHAM

AB	*	250MG	N50459 001
AB		250MG	N62216 001
AB	+	250MG	N62216 001
AB	+	500MG	N62216 002
AB	*	500MG	N62216 004
AB	+	500MG	N62216 004
AB	(@)	500MG	N50459 001
AB	(@)	500MG	N50459 002

POWDER FOR RECONSTITUTION; ORAL

AMOILL

AB	*	SMITHKLINE BEECHAM	1.25MG/5ML	N50460 001
AB			1.25MG/5ML	N62226 001
AB	+		1.25MG/5ML	N62226 001
AB	+		2.50MG/5ML	N50460 002
AB	*		2.50MG/5ML	N50460 005
AB	+		5.0MG/ML	N62226 002
AB	+		2.50MG/5ML	N62226 002
AB	*		2.50MG/5ML	N62226 005
AB	+		5.0MG/ML	N62226 005
AB	*		12.5MG/5ML	N50460 001
AB	+		25.0MG/5ML	N50460 002
AB	(@)		5.0MG/ML	N50460 005
AB	(@)		5.0MG/ML	N50460 006

AB	*	SMITHKLINE BEECHAM	5.0MG/ML	N50460 006
AB	(@)		5.0MG/ML	N50460 006

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATETABLET; ORAL

ADDERALL 10

RICHWOOD PHARM

SHIRE RICHWOOD

ADDERALL 20

RICHWOOD PHARM

SHIRE RICHWOOD

ADDERALL 30

SHIRE RICHWOOD

AB	*	250MG	N50459 001	TABLET; ORAL ADDERALL 5 SHIRE RICHWOOD	1.25MG; 1.25MG; 1.25MG; 1.25MG	N11522 009 MAY 12, 1997
AB	+	250MG	N62216 001			
AB	+	500MG	N62216 001			
AB	*	500MG	N50459 002			
AB	+	500MG	N62216 004			
AB	*	500MG	N62216 004			
AB	(@)	500MG	N50459 001	<u>AMPHOTERICIN B</u>		
AB	(@)	500MG	N50459 002	<u>INJECTABLE; INJECTION AMPHOTERICIN B</u>	50MG/VIAL 50MG/VIAL	N64414 001 DEC 23, 1996
AB	*	1.25MG/5ML	N50460 001			N64414 001 DEC 23, 1996
AB		1.25MG/5ML	N62226 001			
AB	+	1.25MG/5ML	N62226 001			
AB	*	2.50MG/5ML	N50460 002			
AB	+	2.50MG/5ML	N50460 005	<u>INJECTABLE, LIPOSOMAL; INJECTION AMBISOME</u>	50MG/VIAL	N50740 001 AUG 11, 1997
AB	*	2.50MG/5ML	N62226 002	+ FUJISAWA		
AB	+	2.50MG/5ML	N62226 002			
AB	*	5.0MG/ML	N62226 005			
AB	+	5.0MG/ML	N62226 005	<u>OINTMENT; TOPICAL FUNGIZONE + APOTHECON</u>	3% 3%	N50313 001 N50313 001
AB	*	5.0MG/ML	N50460 006	<u>AMPICILLIN/AMPICILLIN TRIHYDRATE</u>		
AB	(@)	5.0MG/ML	N50460 006	CAPSULE; ORAL <u>AMPICILLIN TRIHYDRATE</u>		N60765 001
AB	*	5.0MG/ML	N50460 006	@ ZENITH GOLDLINE	EQ 250MG BASE	N60765 002
AB		5.0MG/ML	N50460 006	@ ZENITH LABS	EQ 500MG BASE	N60765 002
AB	+	5.0MG/ML	N50460 006	AB	EQ 250MG BASE	N60765 002
AB	*	5.0MG/ML	N50460 006		EQ 500MG BASE	
AB	*	5.0MG/ML	N50460 006	<u>ANAGRELIDE HYDROCHLORIDE</u>		
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 007			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	FEB 13, 1996			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 007			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	FEB 13, 1996			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 007			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	FEB 13, 1996			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 007			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	FEB 13, 1996			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 008	<u>CAPSULE; ORAL AGRYLIN ROBERTS LABS</u>	EQ 0.5MG BASE	N20333 001 MAR 14, 1997
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 008			N20333 002 MAR 14, 1997
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	FEB 13, 1996			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	N11522 010			
AB	*	2.5MG; 2.5MG; 2.5MG; 2.5MG	MAY 12, 1997			

ANISOTROPINE METHYLBROMIDE

TABLET; ORAL
 VALPIN 50
 @ DUPONT MERCK
 @ ENDO PHARMS
 > DLT >
 > ADD >

ARBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION
 GENESA
 + GENSTIA
 0 . 05MG/ML

INJECTABLE; INJECTION
 NORMIFLO
 + WYETH AYERST
 5,000 UNITS/0 . 5ML
 +
 10,000 UNITS/0 . 5ML

TABLET; ORAL
 AXOTAL
 + SAVAGE LABS
 ®

650MG; 50MG
 650MG; 50MG

ASPIRIN; BUTALBITAL
 TABLET; ORAL
 AXOTAL
 + SAVAGE LABS
 ®

N88305 001
 OCT 13, 1983
 N88305 001
 OCT 13, 1983

ASPIRIN; BUTALBITAL; CAFFEINE
 TABLET; ORAL
 BUTALBITAL COMPOUND

N88305 002
 OCT 31, 1984
3.25MG; 50MG; 40MG

AB

ZENITH LABS

ASPIRIN; CAFFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL
 ORPHEGENESTIC
 @ PAR PHARM
 ORPHEGENESTIC FORTE
 @ PAR PHARM
 N13428 001
 N13428 001
 JUN 23, 1987
 N71643 001
 JUN 23, 1987

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

N20420 001 SEP 12, 1997	CAPSULE; ORAL PROPOXYPHENE COMPOUND 65 ® ZENITH GOLDLINE	3.89MG; 32 . 4MG; 6.5MG	N83077 002 DEC 07, 1984
MAY 23, 1997	MAY 23, 1997	3.89MG; 32 . 4MG; 6.5MG	N83077 002 DEC 07, 1984
N20227 002 MAY 23, 1997	TABLET; ORAL CODoxy HALSEY	3.25MG; 4 . 5MG; 0 . 38MG	N887464 001 JUL 01, 1982
N20227 001 MAY 23, 1997	MAY 23, 1997	3.25MG; 4 . 5MG; 0 . 38MG	N887464 001 JUL 01, 1982
MAY 23, 1997	AA	3.25MG; 4 . 5MG; 0 . 38MG	N887464 001 JUL 01, 1982
PERCODAN DUPONT MERCK	AA	3.25MG; 4 . 5MG; 0 . 38MG	N073337 006 NO7337 006
+ ENDO PHARMS	AA	3.25MG; 4 . 5MG; 0 . 38MG	N073337 006 NO7337 006
PERCODAN-DEMI	AA	3.25MG; 2 . 25MG; 0 . 19MG	N073337 005 NO7337 005
+ DUPONT MERCK	AA	3.25MG; 2 . 25MG; 0 . 19MG	N073337 005 NO7337 005
+ ENDO PHARMS	AA	3.25MG; 2 . 25MG; 0 . 19MG	N073337 005 NO7337 005

ATENOLOL

AB	TABLET; ORAL ATENOLOL	2.5MG	N74499 001 JUL 30, 1997
AB	MUTUAL PHARM	2.5MG	N74101 001 JUL 17, 1997
AB	SIDMAK LABS NJ	2.5MG	N74101 002 JUL 17, 1997
AB	5.0MG	1.00MG	N74101 003 JUL 17, 1997
AB	1.00MG		

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL
ATENOLOL AND CHLORTHALIDONE
 SIDMAK LABS NJ 50MG; 25MG

AB N74107 001 SEP 24, 1997
AB N74107 002 100MG; 25MG SEP 24, 1997

AP + FAULDING 10MG/ML

ATROPOINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL
DIPHENOXYLATE HCL AND ATROPOINE SULFATE

AA @ ROXANE 0.025MG; 2.5MG SEP 24, 1997
AA @ DIPHENOXYLATE HCL W/ ATROPOINE SULFATE 0.025MG; 2.5MG ICN SEP 24, 1997

AA @ 0.025MG; 2.5MG FEB 16, 1982
AA @ KV PHARM 0.025MG; 2.5MG FEB 16, 1982
AA @ ZENITH GOLDLINE 0.025MG; 2.5MG NBS659 001
AA ZENITH LABS 0.025MG; 2.5MG N86727 001

ATROPOINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION
ATRACURIUM BESYLATE

AP @ ABBOTT 10MG/ML DEC 23, 1996
AP BEDFORD 10MG/ML JUL 18, 1997
AP ESI LEDERLE 10MG/ML N74824 001 SEP 30, 1997
AP FAULDING 10MG/ML N74740 001 MAR 28, 1997
AP GENSIA 10MG/ML N74784 001 JUN 11, 1997
AP OHMEDA 10MG/ML N74753 001 JAN 23, 1997
AP ABEOTT 10MG/ML N74633 001 DEC 23, 1996
AP FAULDING 10MG/ML N74639 001 MAR 25, 1997
AP BEDFORD 10MG/ML N74900 001 JUL 18, 1997
AP ESI LEDERLE 10MG/ML N74825 001 SEP 30, 1997
AP FAULDING 10MG/ML N74741 001 MAR 28, 1997
AP OHMEDA 10MG/ML N74768 001 JAN 23, 1997
AP TRACRIUM 10MG/ML N18831 001 NOV 23, 1983
AP + GLAXO WELLCOME 10MG/ML N18831 002 NOV 20, 1983

AP + TRACRIUM PRESERVATIVE FREE 10MG/ML N18831 001 NOV 23, 1983

AUBANOFIN CAPSULE; ORAL RIDAURA 3MG
AP + CONNECTICS * SMITHKLINE BEECHAM 3MG
AP N18689 001 MAY 24, 1985
AP N18689 001 MAY 24, 1985

<u>AZTHROMYCIN DIHYDRATE</u>		<u>BECLOMETHASONE DIPROPIONATE MONOHYDRATE</u>	
INJECTABLE; INJECTION	ZITHROMAX + PFIZER	EQ 500MG BASE/VIAL	N50733 001 JAN 30, 1997
AZLICILLIN SODIUM			
INJECTABLE; INJECTION			
AZLIN + BAYER	EQ 2GM BASE/VIAL	N50562 001 SEP 03, 1982	
*	EQ 3GM BASE/VIAL	N50562 002 SEP 03, 1982	
*	EQ 4GM BASE/VIAL	N50562 003 SEP 03, 1982	
@	EQ 2GM BASE/VIAL	N50562 001 SEP 03, 1982	
@	EQ 3GM BASE/VIAL	N50562 002 SEP 03, 1982	
@	EQ 4GM BASE/VIAL	N50562 003 > ADD > SEP 03, 1982	
BACITRACIN			
INJECTABLE; INJECTION			
BACIM AP PHARMA TEK	50,000 UNITS/VIAL	N64153 001 MAY 09, 1997	
BACTRACIN AP + PHARMACIA AND UPJOHN	50,000 UNITS/VIAL	N60733 002	
BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE			
OINTMENT; TOPICAL			
CORTISPORIN + GLAXO WELLCOME	400 UNITS/GM; 1% EQ 3.5MG BASE/GM; 5,000 UNITS/GM	N50168 002 MAY 04, 1984	
+ MONARCH PHARMS	400 UNITS/GM; 1% EQ 3.5MG BASE/GM; 5,000 UNITS/GM	N50168 002 MAY 04, 1984	
BECLOMETHASONE DIPROPIONATE			
SUPPOSITORY; RECTAL			
EMETIC CON + ROBERT	EQ 100MG BASE EQ 100MG BASE	N16818 006 N16818 006	
BENZQUINAMIDE HYDROCHLORIDE			
BETAMETHASONE DIPROPIONATE			
CREAM; TOPICAL			
BETAMETHASONE DIPROPIONATE AB CLAY PARK	> ADD > " "	EQ 0.05% BASE	N74579 001 NOV 26, 1997
LOTION; TOPICAL			
BETAMETHASONE DIPROPIONATE ④ ALPHARMA	AB NMC	EQ 0.05% BASE	N71085 001 FEB 03, 1983
BETAMETHASONE DIPROPIONATE AB	NMC	EQ 0.05% BASE	N71085 001 FEB 03, 1987
BETAMETHASONE VALERATE			
CREAM; TOPICAL			
BETAMETHASONE VALERATE FOUGERA	AB	EQ 0.1% BASE	N18861 001 AUG 31, 1983
BETAMETHASONE VALERATE FOUGERA	AB +	EQ 0.1% BASE	N18861 001 AUG 31, 1983
VALISONE AB + SCHERING	AB	EQ 0.1% BASE	N16322 001 N16322 002
BETAMETHASONE VALERATE FOUGERA	AB	EQ 0.1% BASE	N16322 002 N16322 001 N16322 001 AUG 31, 1983

BETAMETHASONE VALERATE

<u>LOTION; TOPICAL</u>	<u>BETAMETHASONE VALERATE</u>	<u>EQ 0.1% BASE</u>
<u>AB + FOUGERA</u>		
<u>AB * SCHERRING</u>		

OINTMENT; TOPICAL

<u>BETAMETHASONE VALERATE</u>	<u>EQ 0.1% BASE</u>
<u>FOUGERA</u>	
<u>AB +</u>	<u>EQ 0.1% BASE</u>
<u>AB * SCHERRING</u>	

BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE

<u>SUSPENSION/DROPS; OPHTHALMIC</u>	
<u>BETOPTIC PILO</u>	<u>EQ 0.25% BASE; 1.75%</u>

N20619 001

APR 17, 1997

N20535 001

JUL 15, 1997

BETHANECHOL CHLORIDE

<u>TABLET; ORAL</u>	
<u>BETHANECHOL CHLORIDE</u>	

N84689 001

<u>BUDESONIDE</u>		<u>BUTABARBITAL SODIUM</u>	
POWDER, METERED;	INHALATION	TABLET; ORAL	N83484 001
PULMICORT ® ASTRA	0 .32MG/INH	SODIUM BUTABARBITAL @ ZENITH GOLDLINE	N84040 001
<u>BUMETANIDE</u>		@ ZENITH LABS	N83484 001
		@	N84040 001
<u>INJECTABLE; INJECTION</u>		<u>BUTOCONAZOLE NITRATE</u>	
<u>BUMETANIDE</u>		CREAM; VAGINAL	N19881 001
AP ABBOTT	0 .25MG/ML	FEMSTAT ONE	FEB 07, 1997
	0 .25MG/ML	+ SYNTEX	2%
AP GENSIA	0 .25MG/ML		
> ADD > AP >	0 .25MG/ML		
SANOFI WINTHROP	0 .25MG/ML		
<u>BUPRENORPHINE HYDROCHLORIDE</u>		<u>INJECTABLE; INJECTION</u>	
		<u>BUTORPHANOL TARTRATE PRESERVATIVE FREE</u>	
		AP ABBOTT [®]	N74620 001
			JAN 22, 1997
		AP	N74626 001
			JAN 23, 1997
		AP	N74620 002
			JAN 22, 1997
		AP	N74626 002
			JAN 23, 1997
<u>BUPROPION HYDROCHLORIDE</u>		<u>INJECTABLE; INJECTION</u>	
		<u>BUTORPHANOL TARTRATE PRESERVATIVE FREE</u>	
		AP ABBOTT [®]	N74137 001
			JUN 03, 1996
		AP	N74137 001
			JUN 03, 1996
		AP	N74137 001
			JUN 03, 1996
<u>TABLET, EXTENDED RELEASE; ORAL</u>		<u>CALCIPOTRIENE</u>	
		<u>WELLBUTRIN</u>	N20358 002
		+ GLAXO WELLCOME	OCT 04, 1996
	150MG		N20358 003
			OCT 04, 1996
			N20711 002
ZYBAN	100MG		MAY 14, 1997
GLAXO WELLCOME	150MG		N20711 003
			MAY 14, 1997
			N20711 002
	100MG		MAY 14, 1997
			N20711 003
			MAY 14, 1997
<u>ZYBAN</u>		<u>SOLUTION; TOPICAL</u>	
GLAXO WELLCOME	100MG	+ DOVONEX	N20611 001
	150MG	+ BRISTOL MYERS SQUIBB 0 .005%	MAR 03, 1997

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

CAPTOPRIL

<u>DEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER</u>	<u>MCGAW</u>	4MG/100ML; 4GM/100ML; 6MG/100ML; 12.0MG/100ML; 6.2MG/100ML	N19624 002 FEB 24, 1988
<u>DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER</u>	<u>MCGAW</u>	41MG/100ML; 4GM/100ML; 6MG/100ML; 12.0MG/100ML; 6.2MG/100ML	N19634 002 FEB 24, 1988
<u>DEXTROSE 10% IN LACTATED RINGER'S IN PLASTIC CONTAINER</u>	<u>MCGAW</u>	20MG/100ML; 2GM/100ML; 30MG/100ML; 600MG/100ML; 31.0MG/100ML	N17310 001 FEB 24, 1988
<u>CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE</u>	<u>MCGAW</u>	20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML; 20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML	N18023 001 N18023 001 N18023 001

WOCKHARDT

N18343	002
N18343	002
N74748	004
RAY 29,	1997
N74748	002
RAY 29,	1997
N74748	001
RAY 29,	1997
N74748	003
RAY 29,	1997
N74418	001
EBB 13,	1996
N74418	001
EBB 13,	1996

WOCKHARDT

N74532 002
MAR 28, 1997
N74532 003
MAR 28, 1997
N74532 004
MAR 28, 1997

CARBAMAZEPINE

N20712 001
SEP 30, 1997

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL
CARBATROL
+ SHIRE LABS

300MG
N20712 002
SEP 30, 1997

TABLET; ORAL

EPITOL
LEMMON

200MG

200MG

100MG

100MG

100MG

100MG

TABLET, CHEWABLE; ORAL

EPITOL
LEMMON

100MG

CARVEDILOL

TABLET; ORAL

COREG

SMITHKLINE BEECHAM

3 . 125MG

N20297 004

MAY 29, 1997

CEFACLOR

AB CAPSULE; ORAL

CEFACLOR

RANBAXY

SEP 17, 1996

N70541 001

SEP 17, 1996

N70541 001

SEP 17, 1986

N73524 001

JUL 29, 1992

N73524 001

JUL 29, 1992

AB

POWDER FOR RECONSTITUTION; ORAL

CEFACLOR

RANBAXY

EQ 250MG BASE

EQ 500MG BASE

CEFAZOLIN SODIUM

AB INJECTABLE; INJECTION

CEFAZOLIN SODIUM

LEMMON

AP

EQ 250MG BASE/VIAL

EQ 500MG BASE/VIAL

EQ 1GM BASE/VIAL

EQ 5GM BASE/VIAL

EQ 10GM BASE/VIAL

EQ 250MG BASE/VIAL

EQ 500MG BASE/VIAL

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EQ 500MG BASE/VIAL

EQ 1

<u>CEFAZOLIN SODIUM</u>	
INJECTABLE; INJECTION <u>CEFAZOLIN SODIUM</u>	
<u>AP</u> <u>TEVA</u>	<u>EQ 5GM BASE/VIAL</u>
	N63018 001 MAR 05, 1990
<u>AP</u>	<u>EQ 10GM BASE/VIAL</u>
	N63018 002 MAR 05, 1990
<u>CEFTRIAXONE SODIUM; LIDOCAINE</u>	
INJECTABLE; INJECTION ROCEPHIN KIT + <u>ROCHE</u>	<u>EQ 1GM BASE/VIAL, N/A; N/A,</u> 1% MAY 08, 1996
	N50585 006
+ <u>CEFTIN</u> + <u>GLAXO WELLCOME</u>	<u>EQ 500MG BASE/VIAL, N/A; N/A,</u> 1% MAY 08, 1996
	N50585 007 MAY 08, 1996
<u>CEFUROXIME AXETIL</u>	
POWDER FOR RECONSTITUTION; ORAL CEFTIN + <u>GLAXO WELLCOME</u>	<u>EQ 125MG BASE/SML</u> N50672 001 JUN 30, 1994
	N50672 001 JUN 30, 1994
	EQ 125MG BASE/5ML
+ <u>CEFUROXIME AXETIL</u>	<u>EQ 250MG BASE/SML</u> N50672 002 APR 29, 1997
	N50672 002 APR 29, 1997
	EQ 250MG BASE/5ML
	EQ 500MG BASE APR 22, 1987
	EQ 500MG BASE APR 22, 1987
<u>CEFUROXIME SODIUM</u>	
INJECTABLE; INJECTION <u>CEFUROXIME SODIUM</u>	
<u>AB</u> <u>HANFORD GC</u>	<u>EQ 750MG BASE/VIAL</u>
	N64125 001 MAY 30, 1997
<u>AP</u>	<u>EQ 1.5GM BASE/VIAL</u>
	N64125 002 MAY 30, 1997
<u>AP</u>	<u>EQ 7.5GM BASE/VIAL</u>
	N64124 001 MAY 30, 1997
<u>AB</u> <u>KEFUROX</u> <u>LILLY</u>	<u>EQ 750MG BASE/VIAL</u>
	N62591 001 JAN 10, 1986
	N62592 001 JAN 10, 1986

<u>CEFUROXIME SODIUM</u>	
INJECTABLE; INJECTION	
<u>KEFUROX</u> <u>LILLY</u>	<u>EQ 750MG BASE/VIAL</u>
	N62591 001 JAN 10, 1986
<u>AB</u>	<u>EQ 750MG BASE/VIAL</u>
	N62592 001 JAN 10, 1986
<u>CEPHAPIRIN SODIUM</u>	
INJECTABLE; INJECTION	
<u>CEFDYLYL</u> <u>APOTHECON</u>	<u>EQ 1GM BASE/VIAL</u>
	N62724 001 DEC 23, 1986
<u>AB</u>	<u>EQ 2GM BASE/VIAL</u>
	N62724 002 DEC 23, 1986
<u>CEPHAPIRIN SODIUM</u>	
INJECTABLE; INJECTION	
<u>CEFDYL</u> <u>APOTHECON</u>	<u>EQ 1GM BASE/VIAL</u>
	N62724 001 DEC 23, 1986
<u>AB</u>	<u>EQ 2GM BASE/VIAL</u>
	N62724 002 DEC 23, 1986
<u>CEPHAPIRIN SODIUM</u>	
INJECTABLE; INJECTION	
<u>CEFDYL</u> <u>APOTHECON</u>	<u>EQ 1GM BASE/VIAL</u>
	N62724 001 DEC 23, 1986
<u>AB</u>	<u>EQ 2GM BASE/VIAL</u>
	N62724 002 DEC 23, 1986

<u>CEPHRADINE</u>		<u>CHLORDIAZEPOXIDE</u>	
CAPSULE; ORAL CEPHRADINE ④ ZENITH GOLDLINE	250MG 500MG AB AB	TABLET; ORAL LIBRITABS ROCHE N62762 001 MAR 06, 1987 N62762 002 MAR 06, 1987 N62762 001 MAR 06, 1987 N62762 002 MAR 06, 1987	5MG 1.0MG 2.5MG 5MG 1.0MG 2.5MG N85482 001 N85481 001 N85488 001
<u>CERIVASTATIN SODIUM</u>		<u>CHLORDIAZEPOXIDE HCL</u>	
TABLET; ORAL BAYCOL ④ BAYER	0 . 05MG 0 . 1MG 0 . 2MG 0 . 3MG +	N20740 001 JUN 26, 1997 N20740 002 JUN 26, 1997 N20740 003 JUN 26, 1997 N20740 004 JUN 26, 1997	SMG 1.0MG 2.5MG 5MG 1.0MG 2.5MG N83570 001 N83570 001
<u>CHLORAMPHENICOL HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE</u>		<u>SOLUTION; DENTAL CHLORHEXIDINE GLUCONATE</u>	
OINTMENT; OPHTHALMIC OPHTHOCCORT ④ PARKE DAVIS	10MG/GM; 5MG/GM; 10,000 UNITS/GM 10MG/GM; 5MG/GM; 10,000 UNITS/GM ④	N50201 002 N50201 002	0.12% AT TEVA
<u>CHLORAMPHENICOL PALMITATE</u>		<u>CHLOROPROCAINE HYDROCHLORIDE</u>	
SUSPENSION; ORAL CHLOROMYCETIN PALMITATE PARKE DAVIS ④	EQ 150MG BASE/5ML EQ 150MG BASE/5ML	N62301 001 N62301 001	0.12% AT TEVA
<u>CHLORAMPHENICOL PALMITATE</u>		<u>INJECTABLE; INJECTION NESACAIN-MPF</u>	
		AP + ASTRA AP +	2% 2%
			MAY 02, 1996
			N09435 004
			N09435 007
			MAY 02, 1996
			N09435 003
			N09435 004

CHLORPHENIRAMINE MALEATE

TABLET; ORAL

CHLORPHENIRAMINE MALEATE

<u>AA</u>	KV PHARM	<u>4MG</u>	N87164 001
@	ZENITH GOLDLINE	4MG	N87164 001
@	ZENITH LABS	4MG	N80779 001
@	ZENITH LABS	4MG	N80779 001
<u>AA</u>	KLOROMIN HALSEY	<u>4MG</u>	N83629 001
@	HALSEY	4MG	N83629 001

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

CHLORPROMAZINE HCL

@	ZENITH GOLDLINE	10MG	N83549 001
@	ZENITH GOLDLINE	25MG	N83549 002
@	ZENITH GOLDLINE	50MG	N83549 003
@	ZENITH GOLDLINE	100MG	N83574 001
@	ZENITH GOLDLINE	200MG	N83575 001
<u>BP</u>	ZENITH LABS	10MG	N83549 001
<u>BP</u>	ZENITH LABS	25MG	N83549 002
<u>BP</u>	ZENITH LABS	50MG	N83549 003
<u>BP</u>	ZENITH LABS	100MG	N83574 001
<u>BP</u>	ZENITH LABS	200MG	N83575 001

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

@	ZENITH GOLDLINE	100MG	N88840 001
OCT 25, 1984			N87353 001
N87353 001			N88840 001
OCT 25, 1984			N87353 001
<u>AB</u>	ZENITH LABS	250MG	N88840 001
<u>AB</u>	ZENITH LABS	100MG	N87353 001
<u>AB</u>	ZENITH LABS	250MG	N87353 001

CHLORZOXAZONE

TABLET; ORAL

<u>AA</u>	MONARCH PHARMS	25MG	N19574 001
+		15MG	N19574 001
@		25MG	DEC 20, 1982
			N88051 001
			NOV 12, 1982
			N19574 001
			DEC 20, 1982
			N19574 002
			FEB 12, 1992
			N19574 001
			DEC 20, 1982
			N88051 001
			NOV 12, 1982

CHLORTHALIDONE

TABLET; ORAL

<u>AA</u>	LEMNON	<u>500MG</u>	N88859 001
<u>AA</u>	TEVA	500MG	MAY 04, 1988
<u>AA</u>	PAPAFLEX	500MG	N88859 001
<u>AA</u>	* JOHNSON RN	500MG	MAY 04, 1988
<u>AB</u>	PUREPAC PHARM	250MG	N11300 003
<u>AB</u>	PUREPAC PHARM	250MG	N11300 003

CHLORTHALIDONE

TABLET; ORAL

CHLORTHALIDONE

<u>AA</u>	PUREPAC PHARM	<u>500MG</u>	N88859 001
<u>AA</u>	LEMNON	500MG	N88859 001
<u>AA</u>	TEVA	500MG	MAY 04, 1988
<u>AA</u>	PAPAFLEX	500MG	N88859 001
<u>AA</u>	* JOHNSON RN	500MG	MAY 04, 1988
<u>AB</u>	PUREPAC PHARM	250MG	N11300 003
<u>AB</u>	PUREPAC PHARM	250MG	N11300 003

ECHOES OF STYAMNE

**POWDER; ORAL
CHOLESTYRAMINE**

<u>AB</u>	<u>BAKER NORTON</u>	<u>EQ 4 GM RESIN/PACKET</u>
<u>AB</u>		<u>EQ 4 GM RESIN/SCOOPFUL</u>

CIMETINE

TABLET. OBAT

CIMETIDINE

<u>AB</u>	SIDMAK LABS NJ	<u>3.00MG</u>	N74566 002
JUL 09, 1997			FEB 27, 1997
<u>AB</u>		<u>4.00MG</u>	N74568 003
N74771 002			FEB 27, 1997
JUL 09, 1997			N74566 001
<u>AB</u>		<u>8.00MG</u>	FEB 27, 1997
			N74365 001
<u>AB</u>	TEVA	<u>2.00MG</u>	FEB 28, 1995
			N74365 002
<u>AB</u>		<u>3.00MG</u>	FEB 28, 1995
			N74365 003
<u>AB</u>		<u>4.00MG</u>	FEB 28, 1995
			N74365 004
<u>AB</u>		<u>8.00MG</u>	
N20519 001			
JUL 21, 1997			

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CIMETIDINE HYDROCHLORIDE

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<u>INJECTABLE; INJECTION CIMETIDINE HCL</u>	<u>EQ</u> <u>300MG BASE/2ML</u>	N74296 001
ABBOTT		MAR 28, 1997
	<u>EQ</u> <u>300MG BASE/2ML</u>	N74412 001
ENDO LABS	<u>EQ</u> <u>300MG BASE/2ML</u>	MAR 28, 1997
ENDO PHARMS	<u>EQ</u> <u>300MG BASE/2ML</u>	N74005 001
GENSIA	<u>EQ</u> <u>300MG BASE/2ML</u>	AUG 31, 1994
SANOFI	<u>EQ</u> <u>300MG BASE/2ML</u>	N74005 001
	<u>EQ</u> <u>300MG BASE/2ML</u>	AUG 31, 1994
	<u>EQ</u> <u>300MG BASE/2ML</u>	N74252 001
	<u>EQ</u> <u>300MG BASE/2ML</u>	NOV 26, 1997
	<u>EQ</u> <u>300MG BASE/2ML</u>	N74296 001
	<u>EQ</u> <u>300MG BASE/2ML</u>	MAR 28, 1997
	<u>EQ</u> <u>300MG BASE/2ML</u>	N74412 001
	<u>EQ</u> <u>300MG BASE/5ML</u>	MAR 28, 1997
<u>SOLUTION; ORAL CIMETIDINE HCL</u>	<u>EQ</u> <u>300MG BASE/5ML</u>	N74251 001
ENDO LABS		DEC 22, 1994
ENDO PHARMS		N74251 001
HI TECH PHARMA		DEC 22, 1994
MORTON GROVE		N74664 001
		OCT 28, 1997
		N74757 001
		OCT 17, 1997

CIMETIDINE HYDROCHLORIDE

SOLUTION; ORAL
CIMETIDINE HCL
PHARM ASSOC
AA ROXANE
AA

EQ 300MG BASE/5ML
 N74553 001 JAN 27, 1997
 N74541 001 AUG 05, 1997

CIPROFLOXACIN

POWDER FOR RECONSTITUTION; ORAL
CIPRO
BAVER
 +

250MG/5ML
 N20780 001 SEP 26, 1997
 500MG/5ML
 N20780 002 SEP 26, 1997

CISAPRIDE MONOHYDRATE

TABLET; ORAL
PROPULSID QUICKSOLV
 + JANSSSEN
 > ADD >
 > ADD >
 > ADD >

EQ 20MG BASE
 N20767 001 NOV 07, 1997

CLEMASTINE FUMARATE

SYRUP; ORAL
CLEMASTINE FUMARATE
AA LEMMON
AA TEVA
 &

EQ 0.5MG BASE/5ML
 N73399 001 JUN 30, 1994
 EQ 0.5MG BASE/5ML
 N73399 001 JUN 30, 1994

CLORETASOL PROPIONATE

CREAM; TOPICAL
CORMAX
AB HEALTHPOINT
 &

GEL; TOPICAL
CLORETASOL PROPIONATE
STIEFEL
AB
 &

0.05%
TEMOVATE
AB + GLAXO WELLCOME
 &

TABLET; ORAL
CLEMASTINE FUMARATE
AB LEMMON
 &

2.68MG
 N73283 001 JAN 31, 1992
 N73282 001 JAN 31, 1992
 N73283 001 JAN 31, 1992
 N73282 001 JAN 31, 1992
 N73283 001 JAN 31, 1992
 N73282 001 JAN 31, 1992

CLOCORTOLONE PIVALATE
CREAM; TOPICAL
CLODERM
 * CTR LABS
 + EM IND'S
AB TEVA
 &

N17765 001
 N17765 001

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL
CLOMIPRAMINE HCL
INVAMED

<u>AB</u>	<u>2.5MG</u>	N74953 001 JUN 25, 1997
<u>AB</u>	<u>5.0MG</u>	N74953 002 JUN 25, 1997
<u>AB</u>	<u>7.5MG</u>	N74953 003 JUN 25, 1997 > <u>ADD</u> >
<u>AB</u>	<u>2.5MG</u>	N74958 001 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>5.0MG</u>	N74958 002 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>7.5MG</u>	N74958 003 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>2.5MG</u>	N74849 001 AUG 26, 1997
<u>AB</u>	<u>5.0MG</u>	N74849 002 APR 04, 1997
<u>AB</u>	<u>7.5MG</u>	N74849 003 APR 04, 1997

CLONAZEPAM

TABLET; ORAL
CLONAZEPAM
ALPHAPHARM

<u>AB</u>	<u>0.5MG</u>	N74940 001 OCT 30, 1997
<u>AB</u>	<u>1MG</u>	N74940 002 OCT 30, 1997
<u>AB</u>	<u>2MG</u>	N74940 003 OCT 30, 1997
<u>AB</u>	<u>0.5MG</u>	N74979 001 AUG 29, 1997
<u>AB</u>	<u>1MG</u>	N74979 002 AUG 29, 1997
<u>AB</u>	<u>2MG</u>	N74979 003 AUG 29, 1997
<u>AB</u>	<u>0.5MG</u>	N74925 001 SEP 30, 1997
<u>AB</u>	<u>1MG</u>	N74925 002 SEP 30, 1997
<u>AB</u>	<u>2MG</u>	N74925 003 SEP 30, 1997
<u>AB</u>	<u>0.125MG</u>	N17533 005 APR 09, 1997

KLONOPIN
+ ROCHE

CLONAZEPAM

TABLET; ORAL
KLONOPIN
ROCHE

<u>AB</u>	<u>0.25MG</u>	N17533 006 APR 09, 1997
<u>AB</u>	<u>5.0MG</u>	N74953 001 JUN 25, 1997
<u>AB</u>	<u>7.5MG</u>	N74953 002 JUN 25, 1997
<u>AB</u>	<u>2.5MG</u>	N74953 003 JUN 25, 1997 > <u>ADD</u> >
<u>AB</u>	<u>5.0MG</u>	N74958 001 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>7.5MG</u>	N74958 002 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>2.5MG</u>	N74958 003 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>5.0MG</u>	N74958 004 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>7.5MG</u>	N74958 005 AUG 26, 1997 > <u>ADD</u> >
<u>AB</u>	<u>2.5MG</u>	N74849 001 AUG 26, 1997
<u>AB</u>	<u>5.0MG</u>	N74849 002 APR 04, 1997
<u>AB</u>	<u>7.5MG</u>	N74849 003 APR 04, 1997
<u>AB</u>	<u>2.5MG</u>	N71925 001 APR 25, 1988
<u>AB</u>	<u>5.0MG</u>	N71925 002 APR 25, 1988
<u>AB</u>	<u>7.5MG</u>	N71925 003 APR 25, 1988
<u>AB</u>	<u>15MG</u>	N71926 001 APR 25, 1988
<u>AB</u>	<u>15MG</u>	N71926 002 APR 25, 1988
<u>AB</u>	<u>15MG</u>	N71926 003 APR 25, 1988
<u>AB</u>	<u>15MG</u>	N72330 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72331 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72332 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72333 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72334 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72335 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72336 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72337 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72338 001 AUG 08, 1988
<u>AB</u>	<u>15MG</u>	N72339 001 AUG 08, 1988

<u>CROMOLYN SODIUM</u>					
<u>AEROSOL, METERED; INHALATION</u>					
INTAL	+ FISON'S	0.8MG/INH	N18887 001 DEC 05, 1985	N18887 001 DEC 05, 1985	
	+ RHONE POULENC RORER	0.8MG/INH			
NOV 26, 1997	N74949 002				
NOV 26, 1997	N74949 001				
NOV 26, 1997	N74949 001				
> ADD >	AB	<u>25MG</u>			
> ADD >	AB	<u>100MG</u>			
> ADD >	AB				
> ADD >	AB				
<u>CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE</u>					
<u>SYRUP; ORAL</u>					
<u>PHEBRAZINE W/ CODEINE</u>					
HALSEY	<u>1.0MG/5ML; 6.25MG/5ML</u>	N88739 001 DEC 23, 1988			
AA		N88739 001 DEC 23, 1988			
@					
<u>PROMETHAZINE HCL AND CODEINE PHOSPHATE</u>					
HI TECH PHARMA	<u>10MG/5ML; 6.25MG/5ML</u>	N40151 001 AUG 26, 1997			
AA					
<u>CORTICOTROPIN</u>					
<u>INJECTABLE; INJECTION</u>					
<u>ACTHAR</u>					
AP	+ RHONE POULENC RORER	<u>40 UNITS/VIAL</u>			
		25 UNITS/VIAL	N07504 003 N07504 002		
		25 UNITS/VIAL	N07504 002 N07504 003		
		40 UNITS/VIAL	N07504 003		
<u>CORTICOTROPIN</u>					
AP	<u>40 UNITS/VIAL</u>		N88772 001 NOV 21, 1984		
	40 UNITS/VIAL		N88772 001 NOV 21, 1984		
<u>CYTANOCOBALAMIN</u>					
<u>INJECTABLE; INJECTION</u>					
<u>COBAVITE</u>					
AP	<u>STERIS</u>				
<u>CORTISONONE ACETATE</u>					
<u>TABLET; ORAL</u>					
<u>CORTISONONE ACETATE</u>					
<u>PUREPAC PHARM</u>					
BP	(@ ZENITH GOLDLINE	<u>2.5MG</u>	N80493 001 N80493 001		
	(@ ZENITH LABS	2.5MG	N80630 001 N83536 001		
	(@	2.5MG	N80630 001 N83536 001		
		2.5MG	N83536 001		

CYANOCOBALAMIN; CYANOCOBALAMIN, CO-57; CYANOCOBALAMIN, CO-58

N/A; N/A
DICOPAC KIT
MEDI PHYSICS
NYCOMED AMERSHAM
> DLT >
> ADD >

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS: OPHTHALMIC
AK-PENTOLATE
AKORN
1%
④ AKORN
AKORN
1%

AT
AT
AT
AT
AT
AT
2%
CYCLOGYL
+ ALCON
2%

CYPROHEPTADINE HYDROCHLORIDE

N40164 001
JAN 13, 1997
N40165 001
JAN 13, 1997
N84108 001
JUL 03, 1995
④

CYPROHEPTADINE HCL

AK
HALSEY
4 MG

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION
CYTOKAN
AP + BRISTOL MYERS SQUIBB 2GM/VIAL
N17406 001
N17406 001
④ 100MG/VIAL
④ 200MG/VIAL
④ 500MG/VIAL
④ 1GM/VIAL
④ 2GM/VIAL
N12142 005
AUG 30, 1982
N12142 001
N12142 002
N12142 003
N12142 004
AUG 30, 1982
N12142 005
AUG 30, 1982
N12142 005
AUG 30, 1982

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION
CYTOKAN
AP ASTA 100MG/VIAL
AP 200MG/VIAL
AP 500MG/VIAL
AP 1GM/VIAL
AP ELKINS SINN 100MG/VIAL
AP 200MG/VIAL
AP 500MG/VIAL
AP 1GM/VIAL
AP CYTOKAN
AP + BRISTOL MYERS SQUIBB 100MG/VIAL
AP + 200MG/VIAL
AP + 500MG/VIAL
AP + 1GM/VIAL

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION
CERUBIDINE
AP RHONE POULENC RORER EQ 20MG BASE/VIAL
N88371 001 JUL 03, 1986 EQ 20MG BASE/VIAL
N88372 001 JUL 03, 1986 EQ 20MG BASE/VIAL
AP WYETH AYERST EQ 20MG BASE/VIAL
N88373 001 JUL 03, 1986 EQ 20MG BASE/VIAL
AP DAUNORUBICIN HCL
BEDFORD EQ 20MG BASE/VIAL
N88374 001 SEP 24, 1986 EQ 20MG BASE/VIAL
N88371 001 JUL 03, 1986 FEB 03, 1995
N88372 001 JUL 03, 1986 N64103 001
N88373 001 JUL 03, 1986 FEB 03, 1995
N88374 001 JUL 03, 1986
N88375 001 JUL 03, 1986
N88376 001 JUL 03, 1986
N88377 001 JUL 03, 1986
N88378 001 JUL 03, 1986
N88379 001 JUL 03, 1986
N88374 001 SEP 24, 1986
DELAVIRDINE MESYLATE
TABLET; ORAL
RESCRIPTOR
+ PHARMACIA AND UPJOHN 100MG
N20705 001
AP APR 04, 1997

<u>DESERPIDINE</u>								
TABLET; ORAL HARMONYL + ABBOTT ④	N10796 002 N10796 002	0.25MG 0.25MG						
<u>DESIPRAMINE HYDROCHLORIDE</u>								
TABLET; ORAL <u>DESIPRAMINE HCL</u> SIDMAK LABS NJ	N71803 001 N71804 001	100MG 150MG	AT AT	MAY 29, 1997 MAY 29, 1997	NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE BAUSCH AND LOMB	0.1% EQ 3.5MG BASE/GM; 10,000 UNITS/GM	N64063 001 JUL 25, 1994	
<u>DESMOPRESSIN ACETATE</u>								
INJECTABLE; INJECTION DDAVP + RHONE POULENC RORER	N18938 001 N74888 001	0.004MG/ML 0.004MG/ML	AP ④	MAR 30, 1984 OCT 15, 1997	NECROCK SHARP DOWME ④	EQ 0.1% PHOSPHATE EQ 0.1% PHOSPHATE	N11983 002 N11983 002	
<u>DESMOPRESSIN ACETATE</u>								
INJECTABLE; INJECTION GENSIA	N17922 002 N17922 002	0.004MG/ML 0.004MG/ML	AP ④	FEB 06, 1989 FEB 06, 1989	DEXTROSE; POTASSIUM CHLORIDE STERIS	EQ 4MG PHOSPHATE/ML EQ 4MG PHOSPHATE/ML	N84355 001 N84355 001	
<u>SPRAY, METERED; NASAL</u>								
DDAVP + RHONE POULENC RORER	N17922 003 N17922 003	0.01MG/SPRAY 0.01MG/SPRAY	AP ④	AUG 07, 1996	POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% IN PLASTIC CONTAINER MCGRAW	5GM/100ML; 37MG/100ML 5GM/100ML; 37MG/100ML	N19699 001 N19699 001	
<u>DEXAMETHASONE</u>								
ELIXIR; ORAL <u>DEXAMETHASONE</u> ALEPHARMA	N88997 001 N88997 001	0.5MG/5ML 0.5MG/5ML	AP ④	OCT 10, 1986 OCT 10, 1986	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER MCGRAW	5GM/100ML; 75MG/100ML 5GM/100ML; 75MG/100ML	N19699 002 N19699 002	
<u>GEL; TOPICAL</u>								
DECADERM + MERCK SHARP DOHME	N13538 001	0.1%						

DEXTOSE; POTASSIUM CHLORIDE

<u>INJECTABLE; INJECTION POTASSIUM CHLORIDE 0.1% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>AP</u>	<u>5GM/100ML; 110MG/100ML SEP 29, 1989 N19699 003</u>	<u>5GM/100ML; 75MG/100ML; 330MG/100ML N18268 011</u>
<u>POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>AP</u>	<u>5GM/100ML; 220MG/100ML SEP 29, 1989 N19699 005</u>	<u>5GM/100ML; 150MG/100ML; 330MG/100ML N18268 012</u>
<u>POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	<u>AP</u>	<u>5GM/100ML; 220MG/100ML SEP 29, 1989 N19699 005</u>	<u>5GM/100ML; 150MG/100ML; 330MG/100ML N18268 012</u>
<u>DEXTROSE 5% SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 150MG/100ML; 330MG/100ML N18268 013</u>	<u>5GM/100ML; 150MG/100ML; 330MG/100ML N18268 013</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 75MG/100ML; 200MG/100ML N19268 009</u>	<u>5GM/100ML; 220MG/100ML; 330MG/100ML N18268 013</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 75MG/100ML; 200MG/100ML N18268 009</u>	<u>5GM/100ML; 220MG/100ML; 330MG/100ML N18268 013</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 150MG/100ML; 200MG/100ML N18268 004</u>	<u>5GM/100ML; 300MG/100ML; 330MG/100ML N18268 014</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 150MG/100ML; 200MG/100ML N18268 004</u>	<u>5GM/100ML; 300MG/100ML; 330MG/100ML N18268 014</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 150MG/100ML; 200MG/100ML N18268 005</u>	<u>5GM/100ML; 300MG/100ML; 330MG/100ML N18268 014</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 220MG/100ML; 200MG/100ML N18268 005</u>	<u>5GM/100ML; 300MG/100ML; 330MG/100ML N18268 014</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 220MG/100ML; 200MG/100ML N18268 006</u>	<u>5GM/100ML; 300MG/100ML; 330MG/100ML N18268 014</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 150MG/100ML; 330MG/100ML N18268 006</u>	<u>4.50MG/1.00ML 4.50MG/1.00ML N18268 001</u>
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE</u>	<u>AP</u>	<u>5GM/100ML; 220MG/100ML; 330MG/100ML N18268 011</u>	<u>4.50MG/1.00ML 4.50MG/1.00ML N18268 010</u>

DEXTOSE; POTASSIUM CHLORIDE; SODIUM CHLORIDEDIAZEPAM

INJECTABLE; INJECTION
DEXTOSE 5% SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE
0.22% IN PLASTIC CONTAINER

⑧ MCGAW
5GM/100ML; 220MG/100ML;

DEXTOSE 5% SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE
0.3% IN PLASTIC CONTAINER

⑧ MCGAW
5GM/100ML; 300MG/100ML;

DEXTOSE 5% SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE
0.3% IN PLASTIC CONTAINER

⑧ MCGAW
5GM/100ML; 300MG/100ML;

DEXTOSE 5% SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE
0.3% IN PLASTIC CONTAINER

⑧ MCGAW
450MG/100ML;

DEXTOSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

⑧ MCGAW
10GM/100ML; 200MG/100ML;

DEXTOSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

⑧ MCGAW
5GM/100ML; 900MG/100ML;

DEXTOSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

⑧ MCGAW
5GM/100ML; 900MG/100ML;

DEXTHYROXINE SODIUM

⑧ KROLL PHARM
1MG

DEXTHYROXINE SODIUM

⑧ KROLL PHARM
2MG

DEXTHYROXINE SODIUM

⑧ KROLL PHARM
4MG

DIAZEPAM

⑧ ATHENA
2.5MG/0.5ML

DIAZEPAM

⑧ ATHENA
5MG/ML

DIAZEPAM

GEL; RECTAL
DIASTAT
ATHENA

N20648 004
JUL 29, 1997

N20648 005
JUL 29, 1997

N20648 004
DEC 20, 1988

N20648 001
AUG 28, 1986

N20648 001
N70912 001

N20648 001
AUG 28, 1986

N20648 001
DEC 20, 1988

N72079 001
AUG 15, 1986

<u>DICLOFENAC SODIUM</u>		<u>DIFLUNISAL</u>	
TABLET, DELAYED RELEASE; ORAL		TABLET; ORAL	
<u>DICLOFENAC SODIUM</u>		<u>DIFLUNISAL</u>	
<u>AB</u>	<u>COPLEY PHARM</u>	<u>AB</u>	<u>DANBURY PHARMA</u>
<u>25MG</u>	<u>N74459 001</u>	<u>250MG</u>	<u>N74400 001</u>
<u>AB</u>	<u>JUN 25, 1997</u>	<u>JUL 17, 1997</u>	<u>JUL 17, 1997</u>
<u>50MG</u>	<u>N74459 002</u>	<u>500MG</u>	<u>N74400 002</u>
<u>AB</u>	<u>JUN 25, 1997</u>		<u>JUL 17, 1997</u>
<u>75MG</u>	<u>N74459 003</u>		
	<u>JUN 25, 1997</u>		
		<u>DIGOXIN</u>	
		INJECTABLE; INJECTION	
		<u>DIGOXIN</u>	
		<u>ABBOTT</u>	
		<u>0.25MG/ML</u>	
		<u>AP</u>	
		<u>SANOFI WINTHROP</u>	
		<u>0.25MG/ML</u>	
		<u>AP</u>	
		<u>DIGOXIN PEDIATRIC</u>	
		<u>ABBOTT</u>	
		<u>0.1MG/ML</u>	
		<u>AP</u>	
		<u>SANOFI WINTHROP</u>	
		<u>0.1MG/ML</u>	
		<u>AP</u>	
		TABLET; ORAL	
		<u>LANOXIN</u>	
		<u>@ GLAXO WELLCOME</u>	
		<u>0.0625MG</u>	
		<u>N84479 001</u>	
		<u>N84479 001</u>	
		<u>SEP 30, 1997</u>	
		<u>N20405 002</u>	
		<u>SEP 30, 1997</u>	
		<u>N20405 003</u>	
		<u>SEP 30, 1997</u>	
		<u>N20405 004</u>	
		<u>SEP 30, 1997</u>	
		<u>N20405 005</u>	
		<u>SEP 30, 1997</u>	
		<u>N20405 006</u>	
		<u>SEP 30, 1997</u>	
		<u>DILTIAZEM HYDROCHLORIDE</u>	
		<u>CAPSULE, EXTENDED RELEASE; ORAL</u>	
		<u>AB</u>	
		<u>* CARDIZEM SR</u>	
		<u>HOECHST MARION RUSSELL</u>	
		<u>60MG</u>	
		<u>AB1</u>	
		<u>+ 60MG</u>	
		<u>STILBESTROL TAB</u>	
		<u>SMITH MILLER PATCH</u>	
		<u>25MG</u>	
		<u>N83004 001</u>	
		<u>N83004 001</u>	
		<u>NO6194 001</u>	
		<u>JAN 23, 1989</u>	
		<u>N19471 001</u>	
		<u>N19471 001</u>	
		<u>JAN 23, 1989</u>	

<u>DILTIAZEM HYDROCHLORIDE</u>		<u>DILTIAZEM HYDROCHLORIDE</u>	
CAPSULE, EXTENDED RELEASE; ORAL	<u>CARDIZEM SR</u>	CAPSULE, EXTENDED RELEASE; ORAL	<u>DILTIAZEM HCL</u>
<u>AB</u> + HOECHST MARION RSSL	<u>90MG</u>	<u>AB1</u> TEVA	<u>120MG</u>
<u>AB1</u> +	<u>90MG</u>		
<u>AB</u> +	<u>120MG</u>		
<u>AB1</u> +	<u>120MG</u>		
<u>AB2</u> + <u>DILACOR XR</u>	<u>120MG</u>	INJECTABLE; INJECTION <u>CARDIZEM</u>	
<u>AB2</u> + <u>WATSON LABS</u>	<u>120MG</u>	<u>AP</u> + HOECHST MARION RSSL	<u>100MG/VIAL</u>
<u>AB2</u> +	<u>180MG</u>		
<u>AB2</u> +	<u>240MG</u>		
		<u>DILTIAZEM HCL</u>	<u>5MG/ML</u>
		<u>AP</u> GENSTA	
		<u>DIMENHYDRINATE</u>	
		<u>AP</u> ELKINS SIANN	<u>50MG/ML</u>
			<u>50MG/ML</u>
<u>AB2</u> ANDRX	<u>120MG</u>		
<u>AB2</u>	<u>180MG</u>		
<u>AB2</u>	<u>240MG</u>		
<u>AB</u> MYLAN	<u>60MG</u>	INJECTABLE; INJECTION BAL	
<u>AB1</u>	<u>60MG</u>	<u>+</u> AKORN	<u>1.0%</u>
<u>AB</u>	<u>90MG</u>	<u>*</u> BECTON DICKINSON	<u>1.0%</u>
<u>AB1</u>	<u>90MG</u>		
<u>AB</u>	<u>120MG</u>		
<u>AB1</u>	<u>120MG</u>	<u>DIPHENHYDRAMINE HYDROCHLORIDE</u>	
<u>AB</u>	<u>120MG</u>	CAPSULE; ORAL	
<u>AB</u>	<u>120MG</u>	<u>DIPHENHYDRAMINE HCL</u>	
<u>AB1</u>	<u>120MG</u>	<u>EON</u>	<u>25MG</u>
<u>AB</u> PROGRAPHARM	<u>60MG</u>	<u>AA</u>	<u>50MG</u>
<u>AB</u>	<u>90MG</u>	<u>AA</u>	<u>50MG</u>
<u>AB</u>	<u>120MG</u>	<u>AA</u>	<u>50MG</u>
<u>AB1</u> TEVA	<u>60MG</u>	<u>AA</u>	<u>2.5MG</u>
<u>AB1</u>	<u>90MG</u>	<u>AA</u>	<u>50MG</u>
		<u>ZENITH LABS</u>	
		<u>AA</u>	<u>50MG</u>
		<u>ZENITH</u>	
		<u>AA</u>	<u>50MG</u>
		<u>INJECTABLE; INJECTION</u>	
		<u>BENADRYL</u>	
		<u>AP</u> PARKE DAVIS	<u>10MG/ML</u>
		<u>AP</u> *	<u>50MG/ML</u>

<u>DILTIAZEM HYDROCHLORIDE</u>		<u>DILTIAZEM HYDROCHLORIDE</u>	
CAPSULE, EXTENDED RELEASE; ORAL	<u>CARDIZEM SR</u>	CAPSULE, EXTENDED RELEASE; ORAL	<u>DILTIAZEM HCL</u>
<u>AB</u> + HOECHST MARION RSSL	<u>90MG</u>	<u>AB1</u> TEVA	<u>120MG</u>
<u>AB1</u> +	<u>90MG</u>		
<u>AB</u> +	<u>120MG</u>		
<u>AB1</u> +	<u>120MG</u>		
<u>AB2</u> + <u>DILACOR XR</u>	<u>120MG</u>	INJECTABLE; INJECTION <u>CARDIZEM</u>	
<u>AB2</u> + <u>WATSON LABS</u>	<u>120MG</u>	<u>AP</u> + HOECHST MARION RSSL	<u>100MG/VIAL</u>
<u>AB2</u> +	<u>180MG</u>		
<u>AB2</u> +	<u>240MG</u>		
		<u>DILTIAZEM HCL</u>	<u>5MG/ML</u>
		<u>AP</u> GENSTA	
		<u>DIMENHYDRINATE</u>	
		<u>AP</u> ELKINS SIANN	<u>50MG/ML</u>
			<u>50MG/ML</u>
<u>AB2</u> ANDRX	<u>120MG</u>		
<u>AB2</u>	<u>180MG</u>		
<u>AB2</u>	<u>240MG</u>		
<u>AB</u> MYLAN	<u>60MG</u>	INJECTABLE; INJECTION BAL	
<u>AB1</u>	<u>60MG</u>	<u>+</u> AKORN	<u>1.0%</u>
<u>AB</u>	<u>90MG</u>	<u>*</u> BECTON DICKINSON	<u>1.0%</u>
<u>AB1</u>	<u>90MG</u>		
<u>AB</u>	<u>120MG</u>		
<u>AB1</u>	<u>120MG</u>	<u>DIPHENHYDRAMINE HYDROCHLORIDE</u>	
<u>AB</u>	<u>120MG</u>	CAPSULE; ORAL	
<u>AB</u>	<u>120MG</u>	<u>DIPHENHYDRAMINE HCL</u>	
<u>AB1</u>	<u>120MG</u>	<u>EON</u>	<u>25MG</u>
<u>AB</u> PROGRAPHARM	<u>60MG</u>	<u>AA</u>	<u>50MG</u>
<u>AB</u>	<u>90MG</u>	<u>AA</u>	<u>50MG</u>
<u>AB</u>	<u>120MG</u>	<u>AA</u>	<u>50MG</u>
<u>AB1</u> TEVA	<u>60MG</u>	<u>AA</u>	<u>2.5MG</u>
<u>AB1</u>	<u>90MG</u>	<u>AA</u>	<u>50MG</u>
		<u>ZENITH LABS</u>	
		<u>AA</u>	<u>50MG</u>
		<u>ZENITH</u>	
		<u>AA</u>	<u>50MG</u>
		<u>INJECTABLE; INJECTION</u>	
		<u>BENADRYL</u>	
		<u>AP</u> PARKE DAVIS	<u>10MG/ML</u>
		<u>AP</u> *	<u>50MG/ML</u>

<u>DIPHENHYDRAMINE HYDROCHLORIDE</u>	<u>DOBUTAMINE HYDROCHLORIDE</u>					
INJECTABLE; INJECTION	INJECTABLE; INJECTION					
<u>BENADRYL</u>	<u>DOBUTAMINE HCL</u>					
@ PARKE DAVIS	SANOFI WINTRUP					
10MG/ML	EQ 12.5MG BASE/ML					
<u>BENADRYL PRESERVATIVE FREE</u>	N74292 001					
AP + PARKE DAVIS	FEB 16, 1995					
50MG/ML						
<u>DIPHENHYDRAMINE HCL</u>						
AP FUJISAWA	N80586 002					
INTL MEDICATION	NO 6146 001					
STERIS	N09486 001					
AP +	N84094 001					
50MG/ML	N80873 001					
INTL MEDICATION	N83533 001					
STERIS	N80873 003					
+ @ DIPHENHYDRAMINE HCL PRESERVATIVE FREE	N80586 002					
AP FUJISAWA	N84094 001					
INTL MEDICATION	N80873 003					
50MG/ML	N80873 001					
50MG/ML	N83533 001					
<u>DIPYRIDAMOLE</u>						
INJECTABLE; INJECTION	<u>DOXAZOSIN MESYLATE</u>					
AP DIPYRIDAMOLE	N74952 001					
GENSIA	NOV 26, 1997					
5MG/ML						
> ADD > AP >	<u>TABLET; ORAL</u>					
> ADD >	CARDURA					
	PFIZER					
	EQ 1MG BASE					
TABLET; ORAL	N19668 001					
DIPYRIDAMOLE	NOV 02, 1990					
CHELSEA LABS	N19668 004					
50MG	NOV 02, 1990					
	N19668 001					
	NOV 02, 1990					
<u>DIRITHROMYCIN</u>	N19668 004					
TABLET, DELAYED RELEASE; ORAL	NOV 02, 1990					
DYNABAC	N50678 001					
@ LIXIX	JUN 19, 1995					
250MG	N50678 001					
250MG	JUN 19, 1995					
+ AB	<u>DOXE PIN HCL</u>					
	FUREPAC PHARM					
	EQ 100MG BASE					
<u>DOBUTAMINE HYDROCHLORIDE</u>	N72110 001					
INJECTABLE; INJECTION	SEP 08, 1988					
DOBUTAMINE HCL	N72110 001					
ABBOTT	SEP 08, 1988					
EQ 12.5MG BASE/ML	FEB 16, 1995					

DOXYCYCLINE HYCLATE

CAPSULE, COATED PELLETS; ORAL
DORYX
 PARKE DAVIS

AB EQ 100MG BASE
AB WARNER CHILCOTT
 EQ 100MG BASE

INJECTABLE; INJECTION
DROPERIDOL
 ABBOTT

2.5MG/ML
AP SANOFI WINTHROP
 2.5MG/ML

ECONAZOLE NITRATE

CREAM; TOPICAL
 SPECTAZOLE
 + J AND J
 * JOHNSON & W.

1%
 1%

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION
ALPHACINE HCL N/ EPINEPHRINE
 CARLISIE

> DLT > AP
 > DLT > AP
 > DLT > AP
 > ADD > @
 > ADD > @
 > ADD >

ERGOLOID MESYLATES

TABLET; SUBLINGUAL
 HYDROGENATED ERGOT ALKALOIDS
 @ ZENITH GOLDLINE
 @ ZENITH LABS

N87186 001
 N87186 001

ERYTHROMYCIN

GEL; TOPICAL
ERYTHROMYCIN
 ALTANA

AB OCT 30, 1985
AB N62653 001
 OCT 30, 1985

SOLUTION; TOPICAL
ERYTHROMYCIN
 ALTANA

AB OCT 30, 1985
AB N62653 001
 OCT 30, 1985

AB AUG 31, 1995
AB N72272 001
 AUG 31, 1995

AB AUG 31, 1995
AB N72272 001
 AUG 31, 1995

AB EQ 250MG BASE
AB EQ 250MG BASE

ESTAZOLAM

TABLET; ORAL
ESTAZOLAM
 ROYCE LABS

AB DEC 23, 1982
AB N18751 001
 DEC 23, 1982

AB DEC 23, 1982
AB N18751 001
 DEC 23, 1982

AB DEC 23, 1982
AB N18751 001
 DEC 23, 1982

N74818 001
 N74818 001
 N74818 002
 N74818 002
 N74818 002

AB TEVA
AB ZENITH GOLDLINE
AB ZENITH LABS

AB EQ 250MG BASE
AB EQ 250MG BASE

N62237 001
 N62237 001

N19080 001
 DEC 26, 1990

N19080 002
 DEC 26, 1990

<u>ESTRADIOL</u>		<u>ETODOLAC</u>	
AB	TABLET; ORAL <u>ESTRADIOL</u> BARR	0.5MG	N40197 001 OCT 22, 1997
AB		1MG	N40197 002 OCT 22, 1997
AB		2MG	N40197 003 OCT 22, 1997
			OCT 22, 1997
<u>ETHINYL ESTRADIOL; LEVONORGESTREL</u>		<u>ETODOLAC</u>	
	TABLET; ORAL-21 ALESESE + WYETH AYERST	0 . 02MG; 0 . 1MG	N20683 001 MAR 27, 1997
<u>TABLET; ORAL-28 ALESESE WYETH AYERST</u>		<u>ETODOLAC</u>	
<u>CAPSULE; ORAL <u>ETODOLAC</u></u>		<u>ETODOLAC</u>	
AB	ENDO PHARMS	200MG	N74842 001 JUL 17, 1997
AB	GENEVA PHARMS	300MG	N74842 002 JUL 17, 1997
AB		200MG	N74840 001 AUG 29, 1997
AB		300MG	N74840 002 AUG 29, 1997
AB	INVAMED	200MG	N74942 001 SEP 30, 1997
AB		300MG	N74942 002 SEP 30, 1997
AB	MYLAN	200MG	N74932 001 MAY 16, 1997
AB	ZENITH GOLDLINE	300MG	N74932 002 MAY 16, 1997
AB		200MG	N74899 001 JUL 08, 1997
AB		300MG	N74899 002 JUL 08, 1997
<u>ESTRADIOL</u>		<u>LODINE</u>	
<u>CAPSULE; ORAL <u>LODINE</u></u>		<u>LODINE</u>	
AB	WYETH AYERST	200MG	N18922 001 JAN 31, 1991
AB		300MG	N18922 003 JAN 31, 1991
<u>ETODOLAC</u>		<u>LODINE</u>	
<u>CAPSULE; ORAL <u>LODINE</u></u>		<u>LODINE</u>	
AB	WYETH AYERST	400MG	N74927 001 OCT 30, 1997
AB	ENDO PHARMS	400MG	N74841 001 JUN 27, 1997
AB	EON	400MG	N74903 001 APR 11, 1997
AB	GENEVA PHARMS	400MG	N74839 001 JUL 11, 1997
AB	INVAMED	400MG	N74846 001 FEB 28, 1997
AB	PUREPAC PHARM	400MG	N74819 001 FEB 28, 1997
AB	ROYCE LABS	400MG	N74892 001 APR 16, 1997
AB	TEVA	400MG	N75009 001 NOV 26, 1997
> ADD >	AB	ZENITH GOLDLINE	N74883 001 FEB 28, 1997
> ADD >	AB		
<u>ETOPOSIDE</u>		<u>ETOPOSIDE</u>	
<u>CAPSULE; INJECTION <u>ETOPOSIDE IMPUREX</u></u>		<u>ETOPOSIDE IMPUREX</u>	
AB		200MG/ML	N74513 001 MAR 14, 1996
AB		200MG/ML	N74813 001 JUL 09, 1997
AB		200MG/ML	N74513 001 MAR 14, 1996
AB			
<u>CAPSULE; INJECTION <u>PIERRE FABRE</u></u>		<u>PIERRE FABRE</u>	
<u>CAPSULE; INJECTION <u>SUPERGEN</u></u>		<u>SUPERGEN</u>	

FAMOTIDINE

INJECTABLE; INJECTION
PEPCID IN PLASTIC CONTAINER
* MERCK
0 .4MG/ML
N20249 001
FEB 18, 1994

PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER
+ MERCK
0 .4MG/ML
N20249 001
FEB 18, 1994

> ADD >

> DLT >

> DLT >

> DLT >

> ADD >

> ADD >

> ADD >

N19950 001
JAN 29, 1990N19950 003
SEP 29, 1992N19950 004
JUL 08, 1994N19950 003
SEP 29, 1992N19950 003
SEP 29, 1992FLUCONAZOLE

INJECTABLE; INJECTION
DIFLUCAN
+ PFIZER
200MG/100ML

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER
200MG/100ML

DIFLUCAN IN SODIUM CHLORIDE 0 .9%
200MG/100ML

DIFLUCAN IN SODIUM CHLORIDE 0 .9% IN PLASTIC CONTAINER
200MG/100ML

DIFLUCAN IN SODIUM CHLORIDE 0 .9%
200MG/100ML

FENTANYL CITRATE

INJECTABLE; INJECTION
FENTANYL CITRATE
ABOTT

EQ 0 .05MG BASE/ML
EQ 0 .05MG BASE/ML

FLECAINIDE ACETATE

TABLET; ORAL
TAMBOCOR
® 3M

50MG
100MG
150MG
50MG

100MG
150MG
100MG
150MG

FLUNISOLIDE

SPRAY, METERED; NASAL
ANCORON
ICN

250MG
500MG
250MG
500MG

FLUOCET

CREAM; TOPICAL
ALPHARMA
®

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

0 .025%

<u>FLUOCINOLONE ACETONIDE</u>	CREAM; TOPICAL <u>FLUOCET</u> NNC	AT 0.025%	N88360 001 JPN 16, 1984	AB <u>LIDEX-E</u> SYNTEX	AB 0.05%	N16908 003
<u>FLUOCINOLONE ACETONIDE</u>	@ ALPHARMA	AT 0.01%	N88361 001 JPN 16, 1984	AB <u>FLUOCINONIDE</u> TARO	AB 0.05%	N74935 001 JUL 29, 1997
AT NNC	AT + MEDICIS	AT 0.01%	N88361 001 JPN 16, 1984	AB <u>LIDEX</u> + MEDICIS SYNTEX	AB 0.05%	N17373 001 N17373 001
AT NNC	AT + SYNTEX	AT 0.01%	N12787 004 N12787 002	AB <u>FLUOCINONIDE</u> ALTANA	AB 0.05%	N17373 001 N17373 001
AT NNC	AT + SYNTEX	AT 0.025%	N12787 005 N12787 002	AB <u>LIDEX</u> + MEDICIS SYNTEX	AB 0.05%	N74905 001 AUG 26, 1997
AT NNC	AT + SYNTEX	AT 0.025%	N12787 005 N12787 002	AB <u>FLUOCINONIDE</u> ALTANA	AB 0.05%	N16909 002 N16909 002
AT NNC	AT + MEDICIS SYNTEX	AT 0.2%	N16161 002 N16161 002	AB <u>LIDEX</u> + MEDICIS SYNTEX	AB 0.05%	N16909 002 N16909 002
AT NNC	AT + MEDICIS SYNTEX	AT 0.2%	N16161 002 N16161 002	SOLUTION; TOPICAL <u>LIDEX</u>	AT 0.05%	N18849 001 APR 06, 1984
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N13960 001 N13960 001	AT + MEDICIS SYNTEX	AT 0.05%	N18849 001 APR 06, 1984
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N13960 001 N13960 001	SOLUTION; TOPICAL <u>LIDEX</u>	AT 0.05%	N18849 001 APR 06, 1984
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	AT + MEDICIS SYNTEX	AT 0.05%	N13790 001 N13790 001
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	FLURANDRENOLIDE <u>CORDRAN</u>	AT 0.05%	N13790 001 N13790 001
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	CREAM; TOPICAL CORDRAN SP + LILLY	AT 0.025% 0.05%	N12806 003 N12806 002
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	CREAM; TOPICAL NEO-SYNALAR + OCLASSEN	AT 0.025% 0.05%	N12806 003 N12806 002
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	LOTION; TOPICAL <u>CORDRAN</u>	AT 0.05%	N13790 001 N13790 001
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	CREAM; TOPICAL CORDRAN + LILLY	AT 0.025% 0.05%	N12806 004 N12806 004
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	OINTMENT; TOPICAL + OCLASSEN	AT 0.025% 0.05%	N12806 004 N12806 004
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	OINTMENT; TOPICAL + OCLASSEN	AT 0.025% 0.05%	N12806 004 N12806 004
AT NNC	AT + MEDICIS SYNTEX	AT 0.25%	N15296 001 N15296 001	OINTMENT; TOPICAL + OCLASSEN	AT 0.025% 0.05%	N12806 004 N12806 004

<u>FLURANDRENOLIDE</u>	<u>FOLIC ACID</u>	
OINTMENT; TOPICAL CORDRAN + OCCLASSEN	TABLET; ORAL <u>FOLIC ACID</u> DARMBURY PHARMA AA + AA AA + LEADERLINE AA + @	NB0680 001 N80680 001 N05897 004 N05897 004
TAPE; TOPICAL CORDRAN + LILLY OCCLASSEN		N12806 001 N16455 001 N16455 001 0.004MG/SQ CM 0.004MG/SQ CM
<u>FLURAZEPAM HYDROCHLORIDE</u>	<u>FOLLITROPIN ALFA/BETA</u>	
CAPSULE; ORAL <u>FLURAZEPAM HCL</u> HALSEY	INJECTABLE; INJECTION FOLLISTIM ORGANON	N20582 001 SEP 29, 1997 N20582 002 SEP 29, 1997
AB AB	1.5MG 3.0MG	N71808 001 JAN 07, 1988 N71809 001 JAN 07, 1988 N71808 001 JAN 07, 1988 N71809 001 JAN 07, 1988
@ @	1.5MG 3.0MG	BX BX GONAL-F SERONO BX BX
		75 IU/VIAL 150 IU/VIAL 75 IU/VIAL 150 IU/VIAL
<u>FLURBIPROFEN</u>	<u>FUROSEMIDE</u>	
TABLET; ORAL <u>FLURBIPROFEN</u> SIDWAK LABS NJ	INJECTABLE; INJECTION <u>FUROSEMIDE</u> ABBOTT	N70578 001 JUL 08, 1987 N72080 001 AUG 13, 1991 N74337 001 OCT 31, 1994 N70578 001 JUL 08, 1987 N72080 001 AUG 13, 1991
AB	5.0MG	N74647 001 APR 01, 1997 N74647 002 APR 01, 1997 N74560 002 MAY 16, 1997
AB	100MG	AP
AB	100MG	AP
		SANOFI WINTHROP STERLING WINTHROP
		100MG/ML 100MG/ML
<u>FLUTICASSONE PROPIONATE</u>		
POWDER; INHALATION FLOVENT + GLAXO WELLCOME		N20549 001 NOV 07, 1997 N20549 002 NOV 07, 1997 N20549 003 NOV 07, 1997
> ADD > > ADD > > ADD > > ADD > > ADD > > ADD >	0.044MG/INH 0.088MG/INH 0.22MG/INH	> DLT > BS NEOSCAN MEDI PHYSICS 2mcg/ML

GUANFACINE HYDROCHLORIDE

<u>TABLET; ORAL</u>	<u>GUANFACINE HCL</u>
<u>AB</u>	<u>AMIDE PHARM</u>
<u>AB</u>	<u>EQ 1MG BASE</u>
<u>AB</u>	<u>EQ 2MG BASE</u>
<u>AB</u>	<u>EQ 1MG BASE</u>
<u>AB</u>	<u>EQ 2MG BASE</u>
<u>AB</u>	<u>EQ 1MG BASE</u>
<u>AB</u>	<u>EQ 2MG BASE</u>
<u>AB</u>	<u>ROYCE LABS</u>
<u>AB</u>	<u>EQ 1MG BASE</u>
<u>AB</u>	<u>EQ 2MG BASE</u>
<u>HALCINONIDE</u>	
<u>CREAM; TOPICAL</u>	
<u>HALOG</u>	
<u>WESTWOOD SQUIBB</u>	
<u>@</u>	

HEPARIN SODIUM

<u>HEPARIN SODIUM</u>	
<u>INJECTABLE; INJECTION</u>	
<u>HEP FLUSH KIT IN PLASTIC CONTAINER</u>	<u>100 UNITS/ML</u>
<u>FUJISAWA</u>	<u>100 UNITS/ML</u>
<u>AP</u>	<u>DEC 05, 1985</u>
	<u>DEC 05, 1985</u>
	<u>N17029 018</u>
	<u>DEC 05, 1985</u>
	<u>N17029 017</u>
	<u>DEC 05, 1985</u>
	<u>N17029 016</u>
	<u>DEC 05, 1985</u>
	<u>N17029 015</u>
	<u>DEC 05, 1985</u>
	<u>N17029 014</u>
	<u>DEC 05, 1985</u>
<u>HEPARIN LOCK FLUSH</u>	<u>10 UNITS/ML</u>
<u>ABBOTT</u>	<u>FEB 28, 1995</u>
<u>AP</u>	<u>FEB 28, 1995</u>
	<u>N40082 001</u>
	<u>N88097 001</u>
	<u>N88346 001</u>
	<u>MAY 18, 1983</u>
	<u>N40082 002</u>
	<u>FEB 28, 1995</u>
	<u>N88098 001</u>
	<u>APR 28, 1983</u>
	<u>N88347 001</u>
	<u>MAY 18, 1983</u>
	<u>N40082 003</u>
	<u>FEB 28, 1995</u>
	<u>N88098 002</u>
	<u>APR 28, 1995</u>
	<u>N88347 002</u>
	<u>MAY 18, 1983</u>
	<u>N40082 004</u>
	<u>FEB 28, 1995</u>
	<u>N88098 003</u>
	<u>APR 28, 1983</u>
	<u>N88347 003</u>
	<u>MAY 18, 1983</u>
	<u>N40082 005</u>
	<u>FEB 28, 1995</u>
	<u>N88098 004</u>
	<u>APR 28, 1983</u>
	<u>N88347 004</u>
	<u>MAY 18, 1983</u>
	<u>N40082 006</u>
	<u>FEB 28, 1995</u>
	<u>N88098 005</u>
	<u>APR 28, 1983</u>
	<u>N88347 005</u>
	<u>MAY 18, 1983</u>
	<u>N40082 007</u>
	<u>FEB 28, 1995</u>
	<u>N88098 006</u>
	<u>APR 28, 1983</u>
	<u>N88347 006</u>
	<u>MAY 18, 1983</u>
	<u>N40082 008</u>
	<u>FEB 28, 1995</u>
	<u>N88098 007</u>
	<u>APR 28, 1983</u>
	<u>N88347 007</u>
	<u>MAY 18, 1983</u>
	<u>N40082 009</u>
	<u>FEB 28, 1995</u>
	<u>N88098 008</u>
	<u>APR 28, 1983</u>
	<u>N88347 008</u>
	<u>MAY 18, 1983</u>
	<u>N40082 010</u>
	<u>FEB 28, 1995</u>
	<u>N88098 009</u>
	<u>APR 28, 1983</u>
	<u>N88347 009</u>
	<u>MAY 18, 1983</u>
	<u>N40082 011</u>
	<u>JUL 26, 1996</u>
	<u>N17029 004</u>
	<u>N88100 001</u>
	<u>N40085 001</u>
	<u>JUL 26, 1996</u>
	<u>N17064 002</u>
	<u>N88099 001</u>
	<u>APR 28, 1983</u>

HEPARIN SODIUM

<u>HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE</u>									
<u>TABLET; ORAL</u>									
<u>HYCODAN</u>									
<u>AB</u>	<u>STERLING WINTHROP</u>	<u>5,000 UNITS/ML</u>	<u>N88100 001</u>	<u>JUN 26, 1988</u>	<u>AA</u>	<u>+ DUPONT MERCK</u>	<u>1.5MG; 5MG</u>	<u>N05213 001</u>	<u>JUL 26, 1988</u>
<u>HEPARIN SODIUM CONTAINER</u>	<u>10,000 UNITS IN PLASTIC</u>	<u>APR 28, 1983</u>	<u>N19339 003</u>	<u>MAR 27, 1985</u>	<u>AA</u>	<u>+ ENDO PHARMS</u>	<u>1.5MG; 5MG</u>	<u>N05213 001</u>	<u>JUL 26, 1988</u>
<u>ABBOTT</u>	<u>10,000 UNITS/100ML</u>	<u>N19339 003</u>	<u>MAR 27, 1985</u>	<u>N19339 003</u>	<u>AP</u>	<u>HYDRALAZINE HYDROCHLORIDE</u>	<u>2.0MG/ML</u>	<u>N40136 001</u>	<u>JUN 30, 1997</u>
<u>HEPARIN SODIUM CONTAINER</u>	<u>12,500 UNITS IN PLASTIC</u>	<u>APR 27, 1985</u>	<u>N19339 004</u>	<u>MAR 27, 1985</u>	<u>AP</u>	<u>INJECTABLE; INJECTION HYDRALAZINE HCL LUITPOLD</u>	<u>2.0MG/ML</u>	<u>N40136 001</u>	<u>JUN 30, 1997</u>
<u>ABBOTT</u>	<u>5,000 UNITS/100ML</u>	<u>N19339 004</u>	<u>MAR 27, 1985</u>	<u>N19339 004</u>	<u>AP</u>	<u>TABLET; ORAL HYDRALAZINE HCL HALSEY</u>	<u>2.5MG</u>	<u>N89130 001</u>	<u>JAN 15, 1986</u>
<u>HEPARIN SODIUM CONTAINER</u>	<u>25,000 UNITS IN PLASTIC</u>	<u>APR 27, 1985</u>	<u>N19339 004</u>	<u>MAR 27, 1985</u>	<u>AA</u>	<u>HALSEY</u>	<u>1.00MG</u>	<u>N89178 001</u>	<u>JAN 15, 1986</u>
<u>ABBOTT</u>	<u>5,000 UNITS/100ML</u>	<u>N19339 004</u>	<u>MAR 27, 1985</u>	<u>N19339 002</u>	<u>AA</u>		<u>2.5MG</u>	<u>N89130 001</u>	<u>JAN 15, 1986</u>
<u>ABBOTT</u>	<u>10,000 UNITS/100ML</u>	<u>N19339 002</u>	<u>MAR 27, 1985</u>	<u>N19339 004</u>	<u>AA</u>		<u>1.00MG</u>	<u>N89178 001</u>	<u>JAN 15, 1986</u>
<u>ABBOTT</u>	<u>5,000 UNITS/100ML</u>	<u>N19339 004</u>	<u>MAR 27, 1985</u>	<u>N19339 004</u>	<u>AA</u>		<u>2.5MG</u>	<u>N89177 001</u>	<u>JUL 29, 1983</u>
<u>ABBOTT</u>	<u>10,000 UNITS/100ML</u>	<u>N19339 002</u>	<u>MAR 27, 1985</u>	<u>N19339 002</u>	<u>AA</u>		<u>2.5MG</u>	<u>N89177 001</u>	<u>JUL 29, 1983</u>
<u>ABBOTT</u>	<u>HEPARIN SODIUM PRESERVATIVE FREE</u>	<u>N89522 001</u>	<u>MAY 04, 1987</u>	<u>MAY 04, 1987</u>	<u>AA</u>	<u>PUREPAC PHARM</u>	<u>2.5MG</u>	<u>N88177 001</u>	<u>JUL 29, 1983</u>
<u>ABBOTT</u>	<u>10,000 UNITS/ML</u>	<u>N89522 001</u>	<u>MAY 04, 1987</u>	<u>MAY 04, 1987</u>	<u>AA</u>		<u>2.5MG</u>	<u>N88177 001</u>	<u>JUL 29, 1983</u>
<u>HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE</u>									
<u>SYRUP; ORAL</u>									
<u>HYCODAN</u>									
<u>AB</u>	<u>+ STERLING WINTHROP</u>	<u>1.5MG/5ML; 5MG/5ML</u>	<u>N05213 002</u>	<u>JUL 26, 1988</u>	<u>AA</u>	<u>HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE</u>	<u>1.00MG</u>	<u>N88358 001</u>	<u>APR 10, 1984</u>
<u>AB</u>	<u>+ ENDO PHARMS</u>	<u>1.5MG/5ML; 5MG/5ML</u>	<u>N05213 002</u>	<u>JUL 26, 1988</u>	<u>AA</u>	<u>CAPSULE; ORAL HYDRALAZINE HCL W/ HYDROCHLOROTHIAZIDE</u>	<u>100MG; 50MG</u>	<u>N88358 001</u>	<u>APR 10, 1984</u>
<u>HYDROPAME</u>									
<u>AB</u>	<u>HALSEY</u>	<u>1.5MG/5ML; 5MG/5ML</u>	<u>N88066 001</u>	<u>JUN 28, 1985</u>	<u>AA</u>	<u>@ ZENITH GOLDLINE</u>	<u>100MG; 50MG</u>	<u>N88358 001</u>	<u>APR 10, 1984</u>
<u>AB</u>	<u>HALSEY</u>	<u>1.5MG/5ML; 5MG/5ML</u>	<u>N88066 001</u>	<u>JUN 28, 1985</u>	<u>AA</u>	<u>@ ZENITH LABS</u>	<u>100MG; 50MG</u>	<u>N88358 001</u>	<u>APR 10, 1984</u>

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE, ORAL	
<u>HYDRALAZINE HCL W/ HYDROCHLOROTHIAZIDE 25/25</u>	
④ ZENITH GOLDLINE	25MG ; 25MG
AB	25MG ; 25MG
<u>HYDRALAZINE HCL W/ HYDROCHLOROTHIAZIDE 50/50</u>	
④ ZENITH GOLDLINE	50MG ; 50MG
AB	50MG ; 50MG
<u>HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE</u>	
TABLET, ORAL	
<u>HYDRALAZINE HCL, HYDROCHLOROTHIAZIDE AND RESERPINE</u>	
④ ZENITH GOLDLINE	25MG ; 15MG ; 0.1MG
BP	25MG ; 15MG ; 0.1MG
<u>HYDROSERPINE PLUS (R-H-H)</u>	
④ ZENITH GOLDLINE	25MG ; 15MG ; 0.1MG
BP	25MG ; 15MG ; 0.1MG
ZENITH LABS	25MG ; 15MG ; 0.1MG
UNIPRES	25MG ; 15MG ; 0.1MG
BP	25MG ; 15MG ; 0.1MG
SOLVAY	④
<u>HYDROCHLOROTHIAZIDE</u>	
TABLET, ORAL	
<u>HYDROCHLOROTHIAZIDE</u>	
④ ZENITH GOLDLINE	50MG
④ ZENITH LABS	50MG
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN</u>	
TABLET, ORAL	
<u>IRBESARTAN-HYDROCHLOROTHIAZIDE</u>	
④ SANOFI	12.5MG ; 7.5MG
+	

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET, ORAL	
<u>METHYLDOPA AND HYDROCHLOROTHIAZIDE</u>	
N88356 001	N70853 001
APR 10, 1984	OCT 08, 1986
④ N88356 001	N70688 001
APR 10, 1984	APR 24, 1986
AB	N70854 001
APR 10, 1984	OCT 08, 1986
N88357 001	N70853 001
APR 10, 1984	OCT 08, 1986
④ N88357 001	N70688 001
APR 10, 1984	APR 24, 1986
AB	N70854 001
APR 10, 1984	OCT 08, 1986
N88358 001	N71458 001
APR 10, 1984	MAR 08, 1988
AB	N71459 001
APR 10, 1984	MAR 08, 1988
N84291 001	N71460 001
N84291 001	MAR 08, 1988
④ N84291 001	N71461 001
N83877 001	MAR 08, 1988
N83877 001	N71458 001
④ ZENITH LABS	MAR 08, 1988
N86298 001	N71459 001
N86298 001	MAR 08, 1988
④ N86298 001	N71460 001
N86298 001	MAR 08, 1988
④ N86298 001	N71461 001
N84658 001	MAR 08, 1988
N84658 001	MAR 08, 1988
④ N84658 001	N71461 001
<u>HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE</u>	
TABLET, ORAL	
<u>UNIRETIC</u>	
SCHWARZ PHARMA	12.5MG ; 7.5MG
+	25MG ; 15MG
<u>HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE</u>	
TABLET, ORAL	
<u>PROPRANOLOL HCL AND HYDROCHLOROTHIAZIDE</u>	
④ ZENITH GOLDLINE	25MG ; 40MG

N71552 001
DEC 01, 1988

N20729 001
JUN 27, 1997

N20729 002
JUN 27, 1997

HYDROCHLOROTHIAZIDE: PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL <u>PROPRANOLOL HCL AND HYDROCHLORTIAZIDE</u>	N71553 001 DEC 01, 1988 N71552 001 DEC 01, 1988 N71553 001 DEC 01, 1988	25MG ; 80MG 25MG ; 40MG 25MG ; 80MG	ZENITH LABS AB AB	TABLET; ORAL <u>ZENITH GOLDLINE</u> HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE	N20716 001 SEP 23, 1997	7.5MG ; 200MG
SYRUP; ORAL <u>HYCOMINE</u>				SYRUP; ORAL <u>HYCOMINE</u>		
HYDROCHLORTIAZIDE; RESERPINE				HYDROCHLORTIAZIDE; RESERPINE		
TABLET; ORAL <u>ZENITH GOLDLINE</u>				TABLET; ORAL <u>ZENITH GOLDLINE</u>		
HYDROCHLORTIAZIDE W/ RESERPINE				HYDROCHLORTIAZIDE W/ RESERPINE		
(6) 25MG ; 0.125MG				(6) 25MG ; 0.125MG		
50MG ; 0.1MG				50MG ; 0.125MG		
50MG ; 0.125MG				50MG ; 0.125MG		
ZENITH LABS				ZENITH LABS		
BP				BP		
25MG ; 0.125MG				25MG ; 0.125MG		
25MG ; 0.1MG				25MG ; 0.1MG		
50MG ; 0.1MG				50MG ; 0.1MG		
HYDROCORTISONE				HYDROCORTISONE		
CREAM; TOPICAL <u>DERMACORT</u>				CREAM; TOPICAL <u>DERMACORT</u>		
MONARCH PHARMS				MONARCH PHARMS		
AT AT				AT AT		
SOLVAY				SOLVAY		
HYDROCORTISONE				HYDROCORTISONE		
(® ALPHARM)				(® ALPHARM)		
NMC				NMC		
PROCTOCORT				PROCTOCORT		
MONARCH PHARMS				MONARCH PHARMS		
AT AT				AT AT		
SOLVAY				SOLVAY		
SYACORT				SYACORT		
MEDICIS				MEDICIS		
HYDROCHLORTIAZIDE; SPIRONOLACTONE				HYDROCHLORTIAZIDE; SPIRONOLACTONE		
TABLET; ORAL <u>ZENITH GOLDLINE</u>				TABLET; ORAL <u>ZENITH GOLDLINE</u>		
HYDROCHLORTIAZIDE				HYDROCHLORTIAZIDE		
25MG ; 25MG				25MG ; 25MG		
ZENITH LABS				ZENITH LABS		
HYDROCHLORTIAZIDE; TRIAMTERENE				HYDROCHLORTIAZIDE; TRIAMTERENE		
CAPSULE; ORAL <u>TRIAMTERENE AND HYDROCHLORTIAZIDE</u>				CAPSULE; ORAL <u>TRIAMTERENE AND HYDROCHLORTIAZIDE</u>		
GENEVA PHARMS				GENEVA PHARMS		
25MG ; 37.5MG				25MG ; 37.5MG		
NOVARTIS				NOVARTIS		
25MG ; 37.5MG				25MG ; 37.5MG		
SEP 09, 1997				SEP 09, 1997		
TABLET; ORAL <u>TRIAMTERENE AND HYDROCHLORTIAZIDE</u>				TABLET; ORAL <u>TRIAMTERENE AND HYDROCHLORTIAZIDE</u>		
PAR PHARM				PAR PHARM		
50MG ; 75MG				50MG ; 75MG		
BP				BP		
TABLET; ORAL <u>HYDROCORTISONE PUREPAC PHARM</u>				TABLET; ORAL <u>HYDROCORTISONE PUREPAC PHARM</u>		
100MG				100MG		
MAY 11, 1998				MAY 11, 1998		

HYDROCORTISONE

TABLET; ORAL
HYDROCORTISONE
PURÉPAC PHARM
BP @
@ 20MG

N84247 002
N84247 003
AUG 31, 1982
N84247 002

1% ; 10%
@ BIOLAN
@ VIVAN

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; AURICULAR (OTIC)
CORTISPORIN
AT + MONARCH PHARMS

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

OTOCORT
STERIS
@ STERIS

SUSPENSION/DROPS; OTIC
CORTISPORIN
AT + GLAXO WELLCOME

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

OTOCORT
STERIS
@ STERIS

SUSPENSION/DROPS; AURICULAR (OTIC)
OTOCORT
STERIS
@ STERIS

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

PEDIOTIC
MONARCH PHARMS

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

SUSPENSION/DROPS; OTIC
OTOCORT
AT + STERIS

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

PEDIOTIC
GLAXO WELLCOME

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

SEP 29, 1987

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL
CORTISPORIN
AT + GLAXO WELLCOME

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

SEP 29, 1987

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL
CORTISPORIN
AT + GLAXO WELLCOME

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

1% ; EQ 3.5MG BASE/ML;
10,000 UNITS/ML

SEP 29, 1987

N81274 001 JUN 19, 1992	N81274 001 JUN 19, 1992
N86052 001 N86052 001	N86052 001 N86052 001

N9637 001 N9637 002	N9637 001 N9637 002
N83759 001 N83759 002	N83759 001 N83759 002

AUG 09, 1985

AUG 09, 1985

AUG 09, 1985

HYDROCORTISONE BUTEPRATE

CREAM; TOPICAL
 PANDEL
 + SAVAGE LABS
 0.1%

N20453 001
 FEB 28, 1997

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL
 LOCOD LIPOCREAM
 YAMANOUCHI
 0.1%

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION
HYDROCORTISONE SODIUM SUCCINATE
ELKINS SINN
 AP @ EQ 1GM BASE/VIAL

HYDROFLUMETHIAZIDE; RESERPINE

TABLET; ORAL
 RESERPINE AND HYDROFLUMETHIAZIDE
 @ ZENITH GOLDLINE
 50MG; 0.125MG

ZENITH LABS

50MG; 0.125MG

N88932 001
 JAN 11, 1985

N88932 001
 JAN 11, 1985

JAN 11, 1985

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION
HYDROMORPHONE HCL
AP ABBOTT

1.0MG/ML

1.0MG/ML

N74598 001
 JUN 19, 1997
 N74598 001
 JUN 19, 1997

ZENITH GOLDLINE

@ ZENITH GOLDLINE

1.0MG
 2.5MG
 5.0MG

1.0MG
 2.5MG
 5.0MG

1.0MG
 2.5MG
 5.0MG

1.0MG
 2.5MG
 5.0MG

N87416 001
 NOV

N87411 001
 NOV

N87416 001
 NOV

N87411 001
 NOV

N87411 001
 NOV

N87416 001
 NOV

N87411 001
 NOV

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION
HYDROXYZINE HCL
AP
AB
AP
AP

STERLING WINTHROP
TABLET; ORAL

HYDROXYZINE HCL
HALSEY

2.5MG
 5MG

1.0MG
 2.5MG

2.5MG
 5.0MG

1.0MG
 2.5MG

HYDROXYZINE PAMOATECAPSULE; ORAL
HY-PAMAB
EON
HYDROXYZINE PAMOATEAB
EONEQ 25MG HCL
N87479 001
EQ 25MG HCL
N87479 001IBUPROFENSUSPENSION; ORAL
TBUEX KNOLL PHARM
@TABLET; ORAL
IBUPROFEN
PUREPAC PHARMABIDARUBICIN HYDROCHLORIDECAPSULE; INJECTABLE;
IDAMYCIN PFS

+ PHARMACIA AND UPJOHN 1MG/ML

N50734 001
FEB 17, 1997IFOSFAMIDEINJECTABLE; INJECTION
IPEX

+ BRISTOL MYERS SQUIBB 1GM/VIAL

3GM/VIAL

1GM/VIAL

3GM/VIAL

IMIQUIMODCREAM; TOPICAL
ALDARA

+ 3M

5%

N20723 001
FEB 27, 1997INDAPAMIDETABLET; ORAL
INDAPAMIDE

+ NYLAN

1.25MG

NOVOPHARM

1.25MG

N74461 002
MAR 26, 1997N74465 001
APR 04, 1997MOTRIN

MCNEIL

300MG
400MG
600MG
800MGN17463 003
N17463 002
N17463 004
N17463 005

MAY 22, 1985

N17463 003
N17463 002
N17463 004
N17463 005N17463 003
N17463 002
N17463 004
N17463 005

<u>INDAPAMIDE</u>	<u>IODOHIPPURATE SODIUM, I-131</u>		
TABLET; ORAL <u>INDAPAMIDE</u> AB Novopharm	2.5MG	N74665 002 APR 04, 1997	INJECTABLE; INJECTION HIPPOTOP BRACCO ② IODOHIPPURATE SODIUM I 131 ③ CIS SORIN 1.2mCi/VIAL 1.2mCi/VIAL 0.2mCi/ML 0.2mCi/ML
INDIUM IN-111 OXYQUINOLINE			
> DLT >			
> DLT >			
> ADD >			
> ADD >			
INDIUM IN-111 PENTETATE DISODIUM			
> DLT >			
> DLT >			
> ADD >			
> ADD >			
INDOCYANINE GREEN			
> DLT >			
> DLT >			
> ADD >			
> ADD >			
VISIPAQUE 270	55%	N20808 001 AUG 29, 1997	INJECTABLE; INJECTION VISIPAQUE 270 NYCOMED ② IPATROPIUM BROMIDE SOLUTION; INHALATION ATROVENT AN + BOEHRINGER INGELHEIM 0.02%
VISIPAQUE 320	65.2%	N20808 002 AUG 29, 1997	INJECTABLE; INJECTION VISIPAQUE 320 NYCOMED ② GRANULES; ORAL ORAGRAP IN CALCIUM BRACCO ③ 3GM/PACKET 3GM/PACKET
IODIXANOL			
INJECTABLE; INJECTION VISIPAQUE 270	55%	N12968 001 N12968 001	
INJECTABLE; INJECTION VISIPAQUE 320	65.2%	N12968 001 N12968 001	

<u>IPRATROPIUM BROMIDE</u>	<u>ISONIAZID</u>	
SOLUTION; INHALATION <u>IPRATROPIUM BROMIDE</u>	<u>SYRUP; ORAL</u>	
<u>AN</u> <u>DEY</u> <u>0.02%</u>	<u>LANTAZID</u> ® LANNETT	N89243 001 FEB 03, 1986
IRBESARTAN	TABLET; ORAL <u>ISONIAZID</u>	
TABLET; ORAL AVAFRO SANOFI	<u>AA</u> MIKART <u>AA</u>	N40090 001 JUN 26, 1997
75MG SEP 30, 1997 N20757 001	@ ZENITH GOLDLINE <u>AA</u>	N40090 002 JUN 26, 1997
150MG SEP 30, 1997 N20757 002	@ ZENITH LABS <u>AA</u>	N80270 001 N83610 001
300MG SEP 30, 1997 N20757 003	@ ZENITH LABS <u>AA</u>	N80270 001 N83610 001
300MG SEP 30, 1997 N20757 004		N83610 001
<u>ISOETHARINE HYDROCHLORIDE</u>	<u>ISOPROTERENOL HYDROCHLORIDE</u>	
SOLUTION; INHALATION <u>ISOETHARINE HCL</u>	AEROSOL, METERED; INHALATION <u>ISOPROTERENOL HCL</u>	
<u>AN</u> <u>ALPHARMA</u> <u>1%</u>	<u>NN</u> * <u>3M</u> <u>NN</u> + <u>ALPHARMA</u> <u>NN</u> @	N10375 004 N10375 004 N85904 001 N85904 001
<u>ISOFLUROPHATE</u>	INJECTABLE; INJECTION <u>ISUPREL</u>	
OINTMENT; OPHTHALMIC FLOROPRYL * MERCK SHARP DOHME ④	<u>AP</u> * <u>ABBOTT</u> <u>AP</u> * <u>SANOFI WINSTROP</u>	N10515 001 N10515 001
0.025% 0.025%		
N10656 001 N10656 001		
<u>ITRACONAZOLE</u>		
	SOLUTION; ORAL SPORANOX + JANSSEN	10MG/ML
<u>ISONIAZID</u>	<u>KETAMINE HYDROCHLORIDE</u>	
SYRUP; ORAL <u>ISONIAZID</u>	INJECTABLE; INJECTION <u>KETAMINE HCL</u>	
<u>AA</u> + <u>CAROLINA MEDCL</u> *	<u>AP</u> <u>ABBOTT</u>	N20657 001 FEB 21, 1997
50MG/5ML 50MG/5ML		
50MG/5ML 50MG/5ML		
50MG/5ML 50MG/5ML		
<u>LANTAZID</u> LANNETT	<u>EQ 50MG BASE/ML</u>	N74549 001 JUN 27, 1996

<u>KETAMINE HYDROCHLORIDE</u>	<u>KRYPTON, KR-81M</u>					
INJECTABLE; INJECTION	GAS; INHALATION					
<u>KETAMINE HCL</u>	<u>EQ 100MG BASE/ML</u>	<u>AP 100MG BASE/ML</u>	<u>AP 50MG BASE/ML</u>	<u>AP 100MG BASE/ML</u>	<u>AP 100MG BASE/ML</u>	<u>AP 100MG BASE/ML</u>
<u>Abbott</u>	<u>N74549 002</u>	<u>JUN 27, 1996</u>	<u>> DLT ></u>	<u>> ADD ></u>		
<u>Sanofi Winthrop</u>	<u>N74549 001</u>	<u>JUN 27, 1996</u>				
<u>Abbott</u>	<u>N74549 002</u>	<u>JUN 27, 1996</u>				
<u>Abbott</u>	<u>N74549 002</u>	<u>JUN 27, 1996</u>				
<u>KETOROLAC TROMETHAMINE</u>	<u>LAMIVUDINE; ZIDOVUDINE</u>					
INJECTABLE; INJECTION	TABLET; ORAL					
<u>KETOROLAC TROMETHAMINE</u>	<u>+ GLAXO WELLCOME</u>	<u>150MG; 300MG</u>	<u>150MG; 300MG</u>	<u>150MG; 300MG</u>	<u>150MG; 300MG</u>	<u>150MG; 300MG</u>
<u>Abbott</u>	<u>N74801 001</u>	<u>JUN 05, 1997</u>				
<u>Abbott</u>	<u>N74802 001</u>	<u>JUN 05, 1997</u>				
<u>Abbott</u>	<u>N74801 002</u>	<u>JUN 05, 1997</u>				
<u>Abbott</u>	<u>N74802 002</u>	<u>JUN 05, 1997</u>				
<u>TORADOL</u>	<u>15MG/ML</u>	<u>15MG/ML</u>	<u>3.0MG/ML</u>	<u>3.0MG/ML</u>	<u>15MG/ML</u>	<u>15MG/ML</u>
<u>Abbott + Syntex</u>	<u>N19598 001</u>	<u>NOV 30, 1989</u>	<u>N19598 002</u>	<u>NOV 30, 1989</u>	<u>N19598 001</u>	<u>NOV 30, 1989</u>
<u>Abbott + Syntex</u>	<u>3.0MG/ML</u>	<u>3.0MG/ML</u>	<u>3.0MG/ML</u>	<u>3.0MG/ML</u>	<u>3.0MG/ML</u>	<u>3.0MG/ML</u>
SOLUTION/DROPS; OPHTHALMIC	<u>LETROZOLE</u>					
ACULAR PRESERVATIVE FREE	TABLET; ORAL					
+ Syntex	<u>0.5%</u>	<u>N20811 001</u>	<u>NOV 03, 1997</u>	<u>N20811 001</u>	<u>NOV 03, 1997</u>	<u>N20811 001</u>
> ADD >						
> ADD >						
> ADD >						
<u>KETOROLAC TROMETHAMINE</u>	<u>LEUCOVORIN CALCIUM</u>					
TABLET; ORAL	TABLET; ORAL					
<u>KETOROLAC TROMETHAMINE</u>	<u>10MG</u>	<u>10MG</u>	<u>10MG</u>	<u>10MG</u>	<u>10MG</u>	<u>10MG</u>
<u>Chelsea Labs</u>	<u>N74955 001</u>	<u>SEP 19, 1997</u>	<u>N74955 001</u>	<u>SEP 19, 1997</u>	<u>N74955 001</u>	<u>SEP 19, 1997</u>
<u>Mylan</u>	<u>N74761 001</u>	<u>MAY 16, 1997</u>	<u>N74761 001</u>	<u>MAY 16, 1997</u>	<u>N74761 001</u>	<u>MAY 16, 1997</u>
<u>Roxane</u>	<u>N74790 001</u>	<u>JUN 26, 1997</u>	<u>N74790 001</u>	<u>JUN 26, 1997</u>	<u>N74790 001</u>	<u>JUN 26, 1997</u>
<u>Teva</u>	<u>N74754 001</u>	<u>MAY 16, 1997</u>	<u>N74754 001</u>	<u>MAY 16, 1997</u>	<u>N74754 001</u>	<u>MAY 16, 1997</u>
<u>Toradol</u>	<u>EQ 10MG BASE/ML</u>	<u>AP + Abbott</u>	<u>EQ 10MG BASE/ML</u>	<u>AP + Abbott</u>	<u>EQ 10MG BASE/ML</u>	<u>AP + Abbott</u>
<u>Abbott + Syntex</u>	<u>EQ 200MG BASE/VIAL</u>	<u>AP + Bedford</u>	<u>EQ 200MG BASE/VIAL</u>	<u>AP + Bedford</u>	<u>EQ 200MG BASE/VIAL</u>	<u>AP + Bedford</u>

<u>LEUCOVORIN CALCIUM</u>		<u>LIDOCAINE HYDROCHLORIDE</u>	
INJECTABLE; INJECTION		INJECTABLE; INJECTION	
<u>LEUCOVORIN CALCIUM</u>		<u>LIDOCAINE HCL</u>	
+ BEDFORD	EQ 200MG BASE/VIAL	SANOFI WINTHROP	<u>2%</u>
AP GENSIA	EQ 350MG BASE/VIAL		N40078 001 JUN 23, 1995
AP + IMMUNEX	EQ 350MG BASE/VIAL		N40056 001 MAY 23, 1995
AP PHARMACHEMIE	EQ 50MG BASE/VIAL		N40174 001 JUN 12, 1997
AP	EQ 100MG BASE/VIAL		N08107 005 APR 05, 1989
AB PAR PHARM	<u>LEUCOVORIN CALCIUM</u>		N89628 001 APR 17, 1997
	EQ 5MG BASE		N89915 001 APR 17, 1997
AB PHARMACHEMIE	EQ 25MG BASE		N74544 001 AUG 28, 1997
AB	EQ 5MG BASE		N74544 002 AUG 28, 1997
AB	EQ 25MG BASE		N73099 001 MAR 28, 1997
			N73101 001 MAR 28, 1997
<u>LEUPROLIDE ACETATE</u>		<u>LORAZEPAM</u>	
INJECTABLE; INJECTION		INJECTABLE; INJECTION	
LUPRON DEPOT-3	11.25MG/VIAL	<u>LORAZEPAM</u>	<u>2MG/ML</u>
+ TAP HOLDINGS		ABBOTT	APR 12, 1994
LUPRON DEPOT-4			N74243 001 APR 12, 1994
+ TAP HOLDINGS	3.0MG/VIAL		N74300 001 APR 12, 1994
<u>LIDOCAINE HYDROCHLORIDE</u>		<u>2MG/ML</u>	
INJECTABLE; INJECTION		<u>4MG/ML</u>	
<u>LIDOCAINE HCL</u>			N74243 002 APR 12, 1994
AP ABBOTT	<u>1%</u>		N74300 002 APR 12, 1994
AP	<u>2%</u>		N74243 003 APR 12, 1994
AP SANOFI WINTHROP	<u>1%</u>		N74648 001 MAR 18, 1997
			0.5MG/5ML
			+ ROXANE

<u>LOXAPINE HYDROCHLORIDE</u>		<u>MALATHION</u>	
CONCENTRATE; ORAL LOXTANE C + COCENSYS * LISTERINE	EQ 25MG BASE/ML EQ 25MG BASE/ML	> DLT > > DLT > > DLT > > ADD > > ADD >	LOTION; TOPICAL OVIDE * GRANDERM @
INJECTABLE; INJECTION + COCENSYS * LISTERINE	EQ 50MG BASE/ML EQ 50MG BASE/ML	N17658 0.01 N17658 0.01	0.05% 0.05%
INJECTABLE; INJECTION + COCENSYS * LISTERINE	EQ 50MG BASE/ML EQ 50MG BASE/ML	N18039 0.01 N18039 0.01	0.5% 0.5%
LOXAPINE SUCCINATE CAPSULE; ORAL LOXTANE COCENSYS	EQ 5MG BASE EQ 10MG BASE EQ 25MG BASE EQ 50MG BASE EQ 5MG BASE EQ 10MG BASE EQ 25MG BASE EQ 50MG BASE	> ADD > > ADD > > ADD > TABLET; ORAL MECLIZINE HCL KV PHARM AA AA	MANGAFODIPIR TRISODIUM INJECTABLE; INJECTION TESLASCAN + NYCOMED 3.7 .9MG/ML
LOXAPINE SUCCINATE CAPSULE; ORAL LOXTANE COCENSYS	EQ 5MG BASE EQ 10MG BASE EQ 25MG BASE EQ 50MG BASE EQ 5MG BASE EQ 10MG BASE EQ 25MG BASE EQ 50MG BASE	N17525 0.01 N17525 0.02 N17525 0.03 N17525 0.04 N17525 0.01 N17525 0.02 N17525 0.03 N17525 0.04	MECLIZINE HYDROCHLORIDE TABLET; ORAL MECLIZINE HCL KV PHARM AA AA
MAGNESIUM SULFATE INJECTABLE; INJECTION MAGNESIUM SULFATE PURISAWA	500MG/ML 500MG/ML	N19316 0.01 SEP 08, 1986 N19316 0.01 SEP 08, 1986	ZENITH GOLDLINE ZENITH LABS AA AA
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER ABBOTT	1GM/100ML 4GM/100ML	N20488 0.01 JUL 11, 1995 N20488 0.01 JUL 11, 1995	VINTAGE PHARMS ZENITH GOLDLINE ZENITH LABS AA
MAGNESIUM SULFATE IN PLASTIC CONTAINER ABBOTT	800MG/ML 4GM/100ML	JUL 11, 1995 N20309 0.01 JUN 24, 1994 N20309 0.02 JUN 24, 1994 N20309 0.01	TABLET, CHEWABLE; ORAL MECLIZINE HCL ZENITH GOLDLINE ZENITH LABS AA
			MENOTROPINS (FSH; LH) INJECTABLE; INJECTION HUMEGON + ORGANON
			75 IU/VIAL, 75 IU/VIAL N20328 001 SEP 01, 1994

MENOTROPINS (FSH,LH)

INJECTABLE; INJECTION

HOMEGON
ORGANON

<u>AB</u>	<u>*</u>	<u>75 IU/VIAL; 75 IU/VIAL</u>	N20328 001
<u>AB</u>		<u>150 IU/VIAL; 150 IU/VIAL</u>	SEP 01, 1994 N20328 002
<u>AB</u>		<u>150 IU/VIAL; 150 IU/VIAL</u>	SEP 01, 1994 N20328 002
<u>AB</u>		<u>150 IU/VIAL; 150 IU/VIAL</u>	SEP 01, 1994 N20328 002
<u>AB</u>		<u>150 IU/VIAL; 150 IU/VIAL</u>	SEP 01, 1994 N20328 002

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HCL
MALLINCKRODT

<u>AP</u>		<u>100MG/ML</u>	N40163 001
			MAY 12, 1997
<u>AA</u>		<u>50MG</u>	N40110 001
<u>AA</u>		<u>100MG</u>	MAR 12, 1997
<u>AA</u>		<u>50MG</u>	N40110 002
<u>AA</u>		<u>100MG</u>	MAR 12, 1997
<u>AA</u>		<u>50MG</u>	N40186 001
<u>AA</u>		<u>100MG</u>	JUN 30, 1997
<u>AA</u>		<u>50MG</u>	N40186 002
<u>AA</u>		<u>100MG</u>	JUN 30, 1997

<u>AA</u>	<u>MEPROBAMATE</u>	<u>TABLET; ORAL</u>	N84804 001
<u>AA</u>		<u>PUREPAC PHARM</u>	400MG
<u>AA</u>			200MG
<u>AA</u>			400MG
<u>AA</u>			600MG

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE
ZENITH LABS

<u>AA</u>	<u>600MG</u>	N84181 001
<u>AA</u>	<u>100MG/5ML</u>	N74702 001
<u>AA</u>	<u>100MG/5ML</u>	MAP 24, 1997
<u>AA</u>	<u>100MG/5ML</u>	N71656 001
<u>AA</u>	<u>100MG/5ML</u>	OCT 13, 1987
<u>AA</u>	<u>100MG/5ML</u>	N74702 001
<u>AA</u>	<u>100MG/5ML</u>	MAR 24, 1997
<u>AA</u>	<u>100MG/5ML</u>	N71656 001
<u>AA</u>	<u>100MG/5ML</u>	OCT 13, 1987

METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOPHAGE
BRISTOL MYERS SQUIBB

<u>AA</u>	<u>500MG</u>	N20357 001
<u>AA</u>	<u>850MG</u>	DEC 29, 1994
<u>AA</u>	<u>850MG</u>	N20357 002
<u>AA</u>	<u>850MG</u>	DEC 29, 1994
<u>AA</u>	<u>500MG</u>	N20357 001
<u>AA</u>	<u>850MG</u>	MAR 03, 1995
<u>AA</u>	<u>850MG</u>	N20357 002
<u>AA</u>	<u>850MG</u>	MAR 03, 1995
<u>AA</u>	<u>100MG/VIAL</u>	METHACHOLINE CHLORIDE
<u>AA</u>	<u>100MG/VIAL</u>	POWDER FOR RECONSTITUTION; INHALATION
<u>AA</u>	<u>100MG/VIAL</u>	METHAPHARM
<u>AA</u>	<u>100MG/VIAL</u>	PROVOCHOLINE
<u>AA</u>	<u>100MG/VIAL</u>	ROCHE

N19193 001
OCT 31, 1986
N19193 001
OCT 31, 1986

<u>METHAZOLAMIDE</u>							
<u>METHYCLOTHIAZIDE</u>							
<u>TABLET; ORAL</u>							
<u>METHAZOLAMIDE</u>	<u>25MG</u>						
<u>AB</u>	<u>APPLIED ANAL</u>						
<u>AB</u>	<u>50MG</u>						
<u>METHOCARBAMOL</u>							
<u>TABLET; ORAL</u>							
<u>METHOCARBAMOL</u>	<u>500MG</u>						
<u>AB</u>	<u>PUREPAC PHARM</u>	<u>750MG</u>					
<u>AB</u>		<u>500MG</u>					
		<u>750MG</u>					
<u>AB</u>			<u>500MG</u>				
			<u>750MG</u>				
<u>METHOTREXATE SODIUM</u>							
<u>INJECTABLE; INJECTION</u>							
<u>MEXATE AQ PRESERVED</u>							
<u>AP</u>	<u>BRISTOL MYERS</u>	<u>EQ 25MG BASE/ML</u>					
			<u>N89887 001</u>				
			<u>APR 14, 1989</u>				
			<u>N89887 001</u>				
			<u>APR 14, 1989</u>				
<u>METHOXSALLEN</u>							
<u>CAPSULE; ORAL</u>							
OXSORALEN-ULTRA							
+ ICN							
			<u>10MG</u>				
			<u>OCT 30, 1986</u>				
			<u>N19600 001</u>				
			<u>OCT 30, 1986</u>				
<u>CAPSULE, LIQUID FILLED; ORAL</u>							
OXSORALEN-ULTRA							
+ ICN							
			<u>10MG</u>				
			<u>OCT 30, 1986</u>				
<u>METHYLPHENIDATE HYDROCHLORIDE</u>							
<u>TABLET; ORAL</u>							
<u>METHYLPHENIDATE HCL</u>	<u>5MG</u>						
<u>AB</u>	<u>DANBURY PHARMA</u>						
<u>AB</u>	<u>10MG</u>						
			<u>OCT 30, 1986</u>				
			<u>N40220 001</u>				
			<u>AUG 29, 1997</u>				
			<u>N40220 002</u>				
			<u>AUG 29, 1997</u>				

METHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

METHYLPHENIDATE HCL
AB DANBURY PHARMA 20MGN19532 001
OCT 30, 1987
N19532 001
OCT 30, 1987METHYLPREDNISOLONE

TABLET; ORAL

MEDROL
AB PHARMacia AND UPJOHN 8MG
METHYLPREDNISOLONE
CHELSEA LABS 4MG
AB INVAMED 4MG
AB TRIGEN 4MG
AB 8MGN40220 003
AUG 29, 1997
+
N40232 001
OCT 16, 1997
N40194 001
OCT 31, 1997
N40189 001
OCT 31, 1997
N40189 002
OCT 31, 1997METOLAZONE

TABLET; ORAL

METRUX
MEDeva
0 . 5MG
0 . 5MGN19532 001
OCT 30, 1987
N19532 001
OCT 30, 1987METOPROLOL TARTRATEINJECTABLE; INJECTION
METOPROLOL TARTRATE
ABBOTT 1MG/ML
AP STERLING WINTHROP 1MG/MLN11153 004
N40232 001
OCT 16, 1997
N40194 001
OCT 31, 1997
N40189 001
OCT 31, 1997
N40189 002
OCT 31, 1997N20743 001
SEP 26, 1997CREAM; TOPICAL
NORITATE
+ DERMIC LABS 1%GEL; VAGINAL
METROGEL -VAGINAL
+ 3M
* CURATEK 0 . 75%
INJECTABLE; INJECTION
METRONIDAZOLE
STERIS 500MG/100MLN74147 001
AUG 02, 1996
N71990 001
JAN 18, 1989
N71990 001
JAN 18, 1989
N74147 001
AUG 02, 1996N20208 001
AUG 17, 1992
N20208 001
AUG 17, 1992N70042 001
DEC 26, 1984
N70042 001
DEC 20, 1984METOCLOPRAMIDEINJECTABLE; INJECTION
METOCLOPRAMIDE HCL
ABBOTT EQ 5MG BASE/ML
AP FAULDING EQ 5MG BASE/ML
@ EQ 5MG BASE/ML
AP SANOFI WINTHROP EQ 5MG BASE/MLN74703 001
OCT 31, 1997
> ADD >
> ADD >
> ADD >
> ADD >N20868 001
NOV 26, 1997METOCLOPRAMIDE

SOLUTION; ORAL

METOCLOPRAMIDE
JVL EQ 5MG BASE/5MLN71536 002
JAN 16, 1997
+ SEARLE 750MGTABLET, EXTENDED RELEASE; ORAL
FLAGYL ER
+ SEARLE 750MG

MEXILETINE HYDROCHLORIDE

<u>AB</u>	CAPSULE; ORAL MEXILETINE HCL WATSON LABS	<u>150MG</u>	N74711 001 FEB 26, 1997	TABLET; ORAL MINOCYCLINE HCL + LIEDERLE	<u>EQ 100MG BASE</u>	N50451 002 AUG 10, 1982
<u>AB</u>		<u>200MG</u>	N74711 002 FEB 26, 1997		<u>EQ 50MG BASE</u>	N50451 003 AUG 10, 1982
<u>AB</u>		<u>250MG</u>	N74711 003 FEB 26, 1997		<u>EQ 100MG BASE</u>	N50451 002 AUG 10, 1982
						AUG 10, 1982

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	TABLET; ORAL MINOCYCLINE HCL + LIEDERLE	<u>EQ 100MG BASE</u>	N50451 002 AUG 10, 1982
		<u>EQ 50MG BASE</u>	N50451 003 AUG 10, 1982
		<u>EQ 100MG BASE</u>	N50451 002 AUG 10, 1982
			AUG 10, 1982

MIBEFRADIL DIHYDROCHLORIDE

<u>AB</u>	TABLET; ORAL POSICOR ROCHE	<u>EQ 50MG BASE</u>	N20689 001 JUN 20, 1997	TABLET; ORAL REMERON * ORGANON	<u>LONG</u>	N20415 002 JUN 14, 1996
		<u>EQ 100MG BASE</u>	N20689 002 JUN 20, 1997		<u>30MG</u>	N20415 002 JUN 14, 1996
					<u>45MG</u>	N20415 003 MAR 17, 1997

<u>AB</u>	+		+ J AND J	TABLET; ORAL MINOCYCLINE HCL + LIEDERLE	<u>EQ 100MG BASE</u>	N50451 002 AUG 10, 1982

MIRTAZAPINE

<u>AB</u>	TABLET; ORAL REMERON * ORGANON	<u>LONG</u>	N20415 002 JUN 14, 1996
		<u>30MG</u>	N20415 002 JUN 14, 1996
		<u>45MG</u>	N20415 003 MAR 17, 1997

MOLINDONE HYDROCHLORIDE

<u>AB</u>	TABLET; ORAL MOBAN DUPONT MERCK	<u>CAPSULE; ORAL</u>	N17111 001
		<u>SMG</u>	N17111 001
		<u>10MG</u>	N17111 002
		<u>2.5MG</u>	N17111 003
		<u>5MG</u>	N17111 001
		<u>10MG</u>	N17111 002
		<u>2.5MG</u>	N17111 003
		<u>SMG</u>	N17111 004
		<u>20MG/ML</u>	N17938 001
		<u>20MG/ML</u>	N17938 001
		<u>TABLET; ORAL</u>	
		<u>MOBAN</u>	
		<u>DUPONT MERCK</u>	
		<u>* ENDO PHARMS</u>	
		<u>ENDO PHARMS</u>	
		<u>TABLET; ORAL</u>	
		<u>MOBAN</u>	
		<u>DUPONT MERCK</u>	
		<u>* ENDO PHARMS</u>	
		<u>ENDO PHARMS</u>	
		<u>TABLET; ORAL</u>	
		<u>MOBAN</u>	
		<u>DUPONT MERCK</u>	
		<u>* ENDO PHARMS</u>	
		<u>ENDO PHARMS</u>	
		<u>TABLET; ORAL</u>	
		<u>MINOCYCLINE HCL</u>	
		<u>MINOCYCLINE HCL</u>	
		<u>LIEDERLE</u>	
		<u>LIEDERLE</u>	

<u>MOLINDONE HYDROCHLORIDE</u>		<u>NALOXONE HYDROCHLORIDE</u>	
TABLET; ORAL MOBAN	25MG 50MG 100MG	N17111 006 N17111 007 N17111 008	AP * AP * AP *
+ ENDO PHARMS			DUPONT MERCK
+			AP * AP *
<u>MOMETASONE FUROATE MONOHYDRATE</u>		<u>TABLET; ORAL; INJECTION</u>	
SPRAY, METERED; NASAL NASONEX	EQ 0.05MG BASE/SPRAY	N20762 001	NARCAN
+ SCHERRING PLOUGH		OCT 01, 1997	DUPONT MERCK
<u>MYCOPHENOLATE MOFETIL</u>		<u>TABLET; ORAL; INJECTION</u>	
TABLET; ORAL CELLCEPT	500MG	N50723 001	ROYCE LABS
+ SYNTEX		JUN 19, 1997	EQ 0.5MG BASE; EQ 50MG BASE
<u>NALBUPHINE HYDROCHLORIDE</u>		<u>TABLET; ORAL; INJECTION</u>	
INJECTABLE; INJECTION NUBALIN	1.0MG/ML 2.0MG/ML	N18024 001 N18024 002	NANDROLONE DECANOATE
+ DUPONT MERCK		MAY 27, 1982	STERIS
AP *		N18024 001	5.0MG/ML
AP *		N18024 002	10.0MG/ML
AP *		MAY 27, 1982	
<u>NALOXONE HYDROCHLORIDE</u>		<u>INJECTABLE; INJECTION</u>	
INJECTABLE; INJECTION NALOXONE HCL	0.02MG/ML 0.4MG/ML	N70171 001 N70172 001	NEDOCROMIL SODIUM
AP ABBOTT		SEP 24, 1986	AEROSOL, METERED; TILADE
AP		N70172 001	* RIXONS
AP STERLING WINTHROP		SEP 24, 1986	1.75MG/INH
AP	0.4MG/ML	N70172 001	RHONE POULENC RORER
		SEP 24, 1986	1.75MG/INH

<u>NEDOCROMIL SODIUM</u>	<u>NIACIN</u>
SOLUTION; INHALATION TILADE + RHONE POULENC RORER 0.5%	TABLET; ORAL <u>NIACIN</u> 001 ④ PUREPAC PHARM ④ ZENITH GOLDLINE ④ ZENITH LABS
	500MG 500MG 500MG
	N83271 001 N83180 001 N83180 001
<u>NEFAZODONE HYDROCHLORIDE</u>	TABLET, EXTENDED RELEASE; ORAL NIASPAN 3.75MG + KOS
	500MG 750MG 1GM
	N20381 001 N20381 002 JUL 28, 1997 JUL 28, 1997 JUL 28, 1997
	N20152 001 DEC 22, 1994 N20152 001 DEC 22, 1994
	N20152 001 DEC 22, 1994
	NIASPAN TITRATION STARTER PACK 3.75MG; 500MG; 750MG
	+ KOS
	N20381 005 JUL 28, 1997
<u>NELFINAVIR MESYLATE</u>	
POWDER FOR RECONSTITUTION; ORAL VIRACEPT + AGOURON	EQ 500MG BASE/SCOOPFUL N20778 001 MAR 14, 1997
	<u>NICOTINE</u>
	FILM, EXTENDED RELEASE; TRANSDERMAL HABITROL 7MG/24HR AB + NOVARTIS
	1.4MG/24HR AB + 2.1MG/24HR AB +
	N20076 001 NOV 27, 1991 N20076 002 NOV 27, 1991 N20076 003 NOV 27, 1991
	N20779 001 MAR 14, 1997
<u>NEOMYCIN SULFATE</u>	
POWDER; FOR RX COMPOUNDING NEO-RX AA PHARMA TEK	EQ 2500MG BASE N61579 001 N61579 001
	100% 100% 100%
	N62385 001 JUN 01, 1982
	100% 21MG/24HR AB +
	INHALANT; INHALATION NICOTROL + PHARMACIA AND UPJOHN 4MG/CARTIDGE
	N20714 001 MAY 02, 1997
<u>NIACIN</u>	
	TABLET; ORAL <u>NIACIN</u> ④ HALSEY ④ PUREPAC PHARM
	500MG 500MG 500MG
	N83453 001 N83453 001 N83271 001

<u>NITROFURANTOIN</u>		<u>NYSTATIN</u>			
TABLET; ORAL		CREAM; TOPICAL			
NITROFURANTOIN ④ ZENITH GOLDLINE	5.0MG 1.00MG	MYKINAC NMC	AT	N62387 001	JUL 29, 1982
④ ZENITH LABS	5.0MG 1.00MG				
④	1.00MG				
NITROFURANTOIN, MACROCRYSTALLINE		OINTMENT; TOPICAL			
CAPSULE; ORAL		NYCOSTATIN			
NITROFURANTOIN AB MYLAN	1.00MG	* SQUIBB ④ WESTWOOD SQUIBB	AT	N60571 001	N60571 001
NITROFURANTOIN AB MYLAN	5.0MG	MYKINAC ④ ALPHARMA	AT	N62731 001	SEP 22, 1986
NOREpinephrine Bitartrate		NILSTAT		N62731 001	SEP 22, 1986
INJECTABLE; INJECTION		LEDERLE	AT	N6144 001	SEP 22, 1986
LEVOPHED + ABBOTT + SANOFI WINTHROP	EQ 1MG BASE/ML EQ 1MG BASE/MD	AT +		N6144 001	
NORTRIPTYLINE HYDROCHLORIDE		SUSPENSION; ORAL			
CAPSULE; ORAL		NYSTATIN			
NORTRIPTYLINE HCL AB INVAMED	EQ 10MG BASE	ALPHARMA	MA	100,000 UNITS/ML	OCT 29, 1985
AB	EQ 25MG BASE	ZOFRAN		N62571 001	OCT 29, 1985
AB	EQ 50MG BASE	+ GLAXO WELLCOME		N62571 001	OCT 29, 1985
AB	EQ 75MG BASE	OXACILLIN SODIUM			
NYSTATIN		SMITHKLINE BEECHAM		N61336 001	
CREAM; TOPICAL		BACTOCILL		N61336 001	
MYKINAC ④ ALPHARMA	100,000 UNITS/GM	AB + OXACILLIN SODIUM AB APOTHECON		N61450 002	
④		AB *		N61450 002	
④		AB @		N61450 001	
④		AB @		N61450 001	
N62387 001					JUL 29, 1982

<u>OXTRIPTYLLINE</u>		<u>OXYMORPHONE HYDROCHLORIDE</u>	
TABLET, EXTENDED RELEASE; ORAL CHOLEFDYL SA	N86742 001 600MG 600MG	INJECTABLE; INJECTION NUMORPHAN DUPOINT MERCK + ENDO PHARMS +	N11707 002 N11707 001 N11707 002 N11707 001
+ PARKE DAVIS + WARNER CHILCOTT			
<u>OXYBUTYNIN CHLORIDE</u>			
SYRUP; ORAL <u>OXYBUTYNIN CHLORIDE</u>	5MG/5ML 5MG/5ML	SUPPOSITORY; RECTAL NUMORPHAN DUPOINT MERCK + ENDO PHARMS +	N11738 004 N11738 004
<u>AB</u> MORTON GROVE	N74868 001 FEB 12, 1997		
<u>AB</u> NU PHARM	N74997 001 OCT 15, 1997	OXYTOCIN	
		INJECTABLE; INJECTION <u>OXYTOCIN</u> WYETH AYERST @	N18243 001 N18243 001
TABLET; ORAL <u>OXYBUTYNIN CHLORIDE</u>	5MG	SOLUTION; NASAL, SYNTOCINON * NOVARTIS @	N12285 001 N12285 001
<u>AB</u> VINTAGE PHARMS	N75079 001 OCT 31, 1997	> <u>DIL</u> > > <u>ADD</u> >	10 USP UNITS/ML 10 USP UNITS/ML
		AP	
<u>OXYCODONE HYDROCHLORIDE</u>			
TABLET, EXTENDED RELEASE; ORAL OXYCONTIN	10MG 20MG	PARMOMYCIN SULFATE	N20553 001 DEC 12, 1995
+ PURDUE FREDERICK		CAPSULE; ORAL N20553 002 DEC 12, 1995	N20553 002 DEC 12, 1995
*		HUMATIN AB + PARKE DAVIS AB	N20553 003 DEC 12, 1995
*	40MG	PAROMOMYCIN SULFATE	N20553 001 DEC 12, 1995
+ PURDUE PHARMA	10MG 20MG	CARACO	N20553 001 DEC 12, 1995
	40MG		N20553 003 DEC 12, 1995
	80MG		N20553 004 JAN 06, 1997
+ +			
		SUSPENSION; ORAL PAXIL + SMITHKLINE BECHAM	EQ 10MG BASE/5ML N20710 001 JUN 25, 1997
<u>OXYMETHOLONE</u>			
TABLET; ORAL ANADROL-50	50MG 50MG		N16848 001 N16848 001
+ SYNTEX + UNIMED			

PENICILLIN G POTASSIUM

TABLET; ORAL
PENICILLIN G POTASSIUM
 @ ZENITH GOLDLINE
 ZENITH LABS

400,000 UNITS
 400,000 UNITS

ABPENTOBARBITAL SODIUM

CAPSULE; ORAL
SODIUM PENTOBARBITAL
 @ ZENITH GOLDLINE
 ZENITH LABS

50MG
 100MG
 50MG
 100MG

AAAA

CAPSULE; ORAL
PENTOBARBITAL SODIUM
 @ ZENITH GOLDLINE
 ZENITH LABS

N83461 001
 N83461 002
 N83461 001
 N83461 002

AA

N16194 001
 N16194 001

N74877 001
 JUL 08, 1997
 N74425 001
 JUL 08, 1997

PENTOXIFYLLINE

TABLET; EXTENDED RELEASE; ORAL
PENTOXIFYLLINE
 PURÉPAC PHARM

AB

400MG
TRENTAL
 AB + HOECHST MARION RSSL 400MG
 AUG 30, 1984

N60073 004
N60073 004

N74878 001
 JUL 09, 1997N18631 001
 AUG 30, 1984PERINDOPRIL ERBUMINE

TABLET; ORAL
PERINDOPRIL ERBUMINE
 ACERON
 RHONE POULENC RORER

AB

2MG
 4MG
 8MG
 12MG
 24MG
 48MG
 *
 N60003 001
 N60003 001

DEC 30, 1993

N20184 002
 DEC 30, 1993

N20184 003

N20184 003
 DEC 30, 1993

N20184 004N20184 005N20184 006N20184 007N20184 008N20184 009N20184 010N20184 011N20184 012N20184 013N20184 014N20184 015N20184 016N20184 017N20184 018N20184 019N20184 020N20184 021N20184 022N20184 023N20184 024N20184 025PENTAZOCINE LACTATE

CAPSULE, EXTENDED RELEASE; ORAL
PENTAZOCINE LACTATE
 BONTRIL MALLINCKRODT 105MG

BC

PHENDIMETRAZINE TARTRATE
 GRAHAM DM 105MG
 BC
 BC
 BC
 BC
 BC
 BC

N68021 001SEP 21, 1982N88028 001AUG 16, 1982N88062 001SEP 13, 1982N88063 001SEP 10, 1982

TABLET, EXTENDED RELEASE; ORAL
PENTOXIFYLLINE
 ESI LEDERLE 400MG
MYLAN 400MG

N74877 001
 JUL 08, 1997
 N74425 001
 JUL 08, 1997

AB

N74878 001
 JUL 09, 1997
 N74425 001
 JUL 09, 1997

AB

PHENDIMETRAZINE TARTRATE

<u>BC</u>	CAPSULE, EXTENDED RELEASE; ORAL PHENDIMETRAZINE TARTRATE GRAHAM DM	105MG	<u>AA</u>	CAPSULE; ORAL <u>PHENTERMINE HCL</u>	37.5MG
@	OCT 18, 1982	N84111 001	<u>AA</u>	KING PHARMS	<u>3.0MG</u>
@	N87214 001	N87214 001	<u>AA</u>	⑥ ZENITH GOLDLINE	<u>3.0MG</u>
@	MAY 26, 1982	N88020 001	⑥ ZENITH LABS	3.0MG 3.0MG	
@	AUG 16, 1982	N88028 001	⑥ ZENITH LABS	3.0MG	
@	AUG 16, 1982	N88062 001	⑥ ZENITH LABS	3.0MG	
@	SEP 13, 1982	N88063 001	⑥ ZENITH LABS	3.0MG	
@	SEP 10, 1982	N88111 001	⑥ EON	3.0MG	
@	N88111 001	OCT 18, 1982	⑥ ZENITH GOLDLINE	3.0MG	
@	OCT 18, 1982	N88553 001	⑥ ZENITH LABS	3.0MG	

TABLET; ORAL
PHENDIMETRAZINE TARTRATE

<u>AA</u>	INWOOD LABS	<u>3.5MG</u>	<u>N84740</u>	CAPSULE; ORAL <u>PHENYLBUTAZONE</u>	100MG
<u>AA</u>	INWOOD LABS	<u>3.5MG</u>	<u>N84742</u>	⑥ ZENITH GOLDLINE	100MG
<u>AA</u>	INWOOD LABS	<u>3.5MG</u>	<u>N84743</u>	⑥ ZENITH LABS	100MG
@	N84740 001	N84740 001	⑥ ZENITH LABS	100MG	
@	N84741 001	N84741 001	⑥ ZENITH LABS	100MG	
@	N84742 001	N84742 001	⑥ ZENITH LABS	100MG	
@	N84743 001	N84743 001	⑥ ZENITH LABS	100MG	

<u>AA</u>	INWOOD LABS	<u>3.5MG</u>	<u>N84138</u>	CAPSULE; INJECTION <u>PHENYTOIN SODIUM</u>	50MG/ML
<u>AA</u>	INWOOD LABS	<u>3.5MG</u>	<u>N84141</u>	⑥ ZENITH GOLDLINE	50MG/ML
<u>AA</u>	INWOOD LABS	<u>3.5MG</u>	<u>N8525</u>	⑥ ZENITH GOLDLINE	50MG/ML
@	N84141 001	N84141 001	N8525 001	N85611 001	⑥ ZENITH GOLDLINE
@	N84141 001	N84141 001	N85612 001	N85612 001	⑥ ZENITH GOLDLINE
@	N85611 001	N85611 001	N85611 001	N85612 001	⑥ ZENITH GOLDLINE
@	N85612 001	N85612 001	N85612 001	N85612 001	⑥ ZENITH GOLDLINE

PHENYTOIN SODIUM, EXTENDED

<u>AA</u>	DAVIS	<u>3.0MG</u>	<u>N89744</u>	CAPSULE; ORAL <u>DILANTIN</u>	3.0MG
> <u>DLT</u>	+	+	N89744 001		
> <u>ADD</u>	+	+	N89744 001		
> <u>DLT</u>	+	+	DEC 18, 1987		
> <u>ADD</u>	+	+	N89744 001		
> <u>ADD</u>	+	+	DEC 18, 1987		

PHENTERMINE HYDROCHLORIDE

<u>AA</u>	CAPSULE; ORAL <u>PHENTERMINE HCL</u>	<u>3.0MG</u>	<u>N84349</u>	CAPSULE; ORAL <u>DILANTIN</u>	3.0MG
+			N84349 001		
JUN 18, 1997			N84349 001		

PIMOZIDE

TABLET; ORAL
ORAP
TEVA

1MG

PINDOLOL

N17473 003
AUG 27, 1997

TABLET; ORAL
PINDOLOL
LEXMIRON

5MG
AB *
10MG
AB *
TEVA

5MG
AB *
10MG
AB *

N74123 001
APR 17, 1997
N74123 002
APR 17, 1997
N74123 001
APR 17, 1997
N74123 002
APR 17, 1997

CAPSULE; ORAL
PIROXICAM
EGIS

10MG
AB *
20MG
AB *

N74808 001
JUL 08, 1997
N74808 002
JUL 08, 1997

PODOFILLOX

GEL; TOPICAL
CONDYLOX
+ OCCLASSEN

N20529 001
MAR 13, 1997

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC
POLYTRIM
AT + ALLERGAN

N50567 001
OCT 20, 1988

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE; ORAL
POTASSIUM CITRATE
MISSION PHARMA 5MEQ
1.0MEQ

N19071 001
AUG 30, 1985
N19071 002
AUG 31, 1992

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE
AT BAUSCH AND LOMB
EQ 1MG BASE/ML

N64120 001
FEB 14, 1997

POTASSIUM CHLORIDE

INJECTABLE; INJECTION
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER
245MG/100ML

AP * ABBOTT	AP *								
APR 17, 1997									
N74123 001	N74123 002								

INJECTABLE; INJECTION
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER
245MG/100ML

AP * ABBOTT	AP *								
APR 17, 1997									
N74123 001	N74123 002								

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL
POTASSIUM CITRATE
MISSION PHARMA 5MEQ
1.0MEQ

N19071 001
AUG 30, 1985
N19071 002
AUG 31, 1992

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL	
POTASSIUM CITRATE	
UNIV TX	
+ 10MEQ	

PREDNISOLE DIHYDROCHLORIDE

TABLET; ORAL	
MIRALEX	
+ PHARMACIA AND UPJOHN	0.125MG
0.25MG	
1MG	
1.25MG	
1.5MG	
+ @	

N19071.001 AUG 30, 1985	N19071.002 AUG 31, 1992
<u>PREDNISONE</u>	

PREDNISOLE DIHYDROCHLORIDE

TABLET; ORAL	
PREDNISONE	
HALSEY	
AB	

SOLUTION; ORAL	
PEDIAPIRED	
+ MEDEV'A	
EQ 5MG BASE/5ML	

PREDNISONE

TABLET; ORAL

PREDNISONE

HALSEY

ZENITH GOLDLINE

INWOOD LABS

WINTHROP

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL	
PEDIAPIRED	
+ MEDEV'A	
EQ 5MG BASE/5ML	

N19157.001
MAY 28, 1986

N80300.001
N80300.001

N80283.001
N80283.001

N84133.001
N84133.001

N84134.001
N84134.001

N80283.001
N80283.001

N84133.001
N84133.001

N84134.001
N84134.001

N80300.001
N80300.001

N80300.001
N80300.001

N80283.001
N80283.001

N84133.001
N84133.001

N84134.001
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N84133.001
N84133.001

N84134.001
N84134.001

N80300.001
N80300.001

N80300.001
N80300.001

N80283.001
N80283.001

N84133.001
N84133.001

N84134.001
N84134.001

N80300.001
N80300.001

N80300.001
N80300.001

N80283.001
N80283.001

N84133.001
N84133.001

PREDNISOLONE ACETATE

INJECTABLE; INJECTION	
PREDNISOLONE ACETATE	
STERIS	
(@)	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

INJECTABLE; INJECTION	
NOVOCAIN	
INNOVACIN	
ABOTT	

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL	
PEDIAPIRED	
+ FISON'S	
EQ 5MG BASE/5ML	

N19157.001
MAY 28, 1986

PROCAINE HYDROCHLORIDEINJECTABLE; INJECTION
PROCAINE HCL

<u>AP</u>	<u>STERIS</u>	<u>1%</u>
<u>AP</u>	<u>@</u>	<u>2%</u>
<u>AP</u>	<u>@</u>	<u>1%</u>
<u>AP</u>	<u>@</u>	<u>2%</u>

PROCaine HYDROCHLORIDE; TETRACYCLINE HYDROCHLORIDEINJECTABLE; INJECTION
TETRACYCN

<u>*</u>	<u>PFIZER</u>	<u>40MG/VIAL; 250MG/ML</u>
<u>*</u>	<u>PFIZER</u>	<u>40MG/VIAL; 250MG/ML</u>

PROCHLORPERAZINE EDISYLATEINJECTABLE; INJECTION
PROCHLORPERAZINE EDISYLATE

<u>AP</u>	<u>ABBOTT</u>	<u>EQ 5MG BASE/ML</u>
<u>AP</u>	<u>STERIS</u>	<u>EQ 5MG BASE/ML</u>
<u>AP</u>	<u>@</u>	<u>EQ 5MG BASE/ML</u>
<u>AP</u>	<u>STERLING WINTHROP</u>	<u>EQ 5MG BASE/ML</u>

PROCHLORPERAZINE MALEATETABLET; ORAL
PROCHLORPERAZINE MALEATE

<u>AB</u>	<u>DURAMED</u>	<u>EQ 5MG BASE</u>
<u>AB</u>	<u>@</u>	<u>EQ 10MG BASE</u>

PROGESTERONEGEL; VAGINAL
CRINONE
COLUMBIA RES LABS

4%

JUL 31, 1997

PROGESTERONEGEL; VAGINAL
CRINONE

N83535 001

N83535 002

N83535 001

N83535 002

N83838 002

N83552 001

N83532 002

N83532 001

N83532 002

N83838 002

N83838 002

N83535 001

N83535 002

N83535 001

N83535 002

N83089 001

N83089 002

N83089 001

N83089 002

PROMETHAZINE HYDROCHLORIDEINJECTABLE; INJECTION
PROMETHAZINE HCL

<u>AP</u>	<u>ABBOTT</u>	<u>50MG/ML</u>
<u>AP</u>	<u>STERIS</u>	<u>25MG/ML</u>
<u>AP</u>	<u>@</u>	<u>50MG/ML</u>
<u>AP</u>	<u>@</u>	<u>25MG/ML</u>
<u>AP</u>	<u>STERLING WINTHROP</u>	<u>50MG/ML</u>

PROPANTHELINE BROMIDETABLET; ORAL
PROPANTHELINE BROMIDE

<u>BP</u>	<u>ZENITH PHARM</u>	<u>1.25MG</u>
<u>BP</u>	<u>ZENITH LABS</u>	<u>2.5MG</u>
<u>BP</u>	<u>ZENITH LABS</u>	<u>5.0MG</u>
<u>BP</u>	<u>ZENITH LABS</u>	<u>12.5MG</u>
<u>BP</u>	<u>ZENITH LABS</u>	<u>25MG</u>
<u>BP</u>	<u>ZENITH LABS</u>	<u>50MG</u>

PROPOXYPHENE HYDROCHLORIDECAPSULE; ORAL
PROPOXYPHENE HCL

<u>MA</u>	<u>ROXANE</u>	<u>32MG</u>
<u>MA</u>	<u>@</u>	<u>65MG</u>
<u>MA</u>	<u>@</u>	<u>32MG</u>
<u>MA</u>	<u>@</u>	<u>65MG</u>

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL
PROPOXYPHENE HCL
 @ ZENITH GOLDLINE
AA

3.2MG
 3.2MG

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL
PROPRANOLOL HCL
 ROXANE

10MG

20MG

40MG

60MG

80MG

90MG

10MG

20MG

40MG

60MG

80MG

60MG

80MG

10MG

20MG

40MG

60MG

80MG

10MG

20MG

40MG

60MG

PROPRANOLOL HYDROCHLORIDE

CAPSULE; ORAL
PROPRANOLOL HCL
 ZENITH LABS

N83597 001
 N83597 001

AB

N72066 001
 JUL 29, 1988

N72067 001
 JUL 29, 1988

N72068 001
 JUL 29, 1988

N72069 001
 JUL 29, 1988

N72068 001
 JUL 29, 1988

N80215 001
 N80215 001

AB

N72069 001
 JUL 29, 1988

AB

N72068 001
 JUL 29, 1988

PROPYLTHIOURACIL

TABLET; ORAL
PROPYLTHIOURACIL
 ZENITH LABS

AB

AB

AB

AB

AB

AB

AB

AB

PROPYLENEDIHYDROCHEMICALINE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL
TRIPROLIDINE HCL AND PSEUDOEPHEDRINE HCL
 ZENITH GOLDLINE

AB

AB

AB

AB

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL
TRIPROLIDINE HCL AND PSEUDOEPHEDRINE HCL
 ZENITH GOLDLINE

AB

AB

AB

AB

PYRIDOSTIGMINE BROMIDE

TABLET; ORAL
PYRIDOSTIGMINE BROMIDE
 ROCHE

002

002

002

002

002

002

002

002

002

002

ZENITH LABS

TABLET; ORAL
PROPRANOLOL HCL
 ZENITH LABS

AB

ZENITH LABS

<u>PYRIDOSTIGMINE BROMIDE</u>		<u>QUINIDINE SULFATE</u>	
TABLET, EXTENDED RELEASE; ORAL + ION	NESTINON	N11665 001 N11665 001	TABLET, EXTENDED RELEASE; ORAL <u>QUINIDEX</u> AB + ROBINS AH
ROCHE	180MG 180MG	AB + WYETH AYERST	300MG 300MG
<u>PYRIDOXINE HYDROCHLORIDE</u>		<u>RANITIDINE HYDROCHLORIDE</u>	
<u>INJECTABLE; INJECTION</u>		<u>CAPSULE; ORAL</u>	
<u>PYRIDOXINE HCL</u>		<u>RANITIDINE HCL</u>	
AP STERIS @		GENEVA PHARMS	
100MG/ML	100MG/ML	EQ 150MG BASE	N74655 001
N83760 001 N83760 001	AB	EQ 300MG BASE	OCT 22, 1997
	AB	EQ 300MG BASE	N74655 002
	AB	EQ 300MG BASE	OCT 22, 1997
<u>QUETIAPINE FUMARATE</u>		<u>ZANTAC 150</u>	
TABLET; ORAL		GLAXO WELLCOME	
SERQUEL ZENECA		EQ 150MG BASE	N20095 001
EQ 25MG BASE	N20639 001 SEP 26, 1997	AB + ZANTAC 300	MAR 08, 1994
EQ 100MG BASE	N20639 002 SEP 26, 1997	GLAXO WELLCOME	N20095 002
+	EQ 200MG BASE	TABLET; ORAL	MAR 08, 1994
	N20639 003 SEP 26, 1997	RANITIDINE HCL BOEHRINGER INGELHEIM	N74662 001
	AB	EQ 150MG BASE	AUG 29, 1997
	AB	EQ 300MG BASE	N74662 002
	AB	EQ 300MG BASE	AUG 29, 1997
<u>QUINIDINE SULFATE</u>		<u>CHELSEA LABS</u>	
<u>CAPSULE; ORAL</u>		<u>EQ 150MG BASE</u>	
CIN-QIN SOLVAY		<u>EQ 300MG BASE</u>	
+ @ @		<u>EQ 150MG BASE</u>	
200MG	N85296 001 N85297 001 N85296 001	AB	N74864 001
300MG	N85297 001	GENEVA PHARMS	OCT 20, 1997
200MG	N85296 001	AB	N74864 002
300MG	N85297 001	AB	OCT 20, 1997
	N85297 001	AB	N74467 001
	AB	GENPHARM	AUG 29, 1997
	AB	EQ 150MG BASE	N74467 002
	AB	EQ 300MG BASE	AUG 29, 1997
	AB	EQ 300MG BASE	N74023 001
	AB	EQ 300MG BASE	AUG 22, 1997
	AB	EQ 300MG BASE	N74023 002
	AB	EQ 300MG BASE	AUG 22, 1997
<u>QUINIDINE SULFATE</u>		<u>GRANUTEC PHARMS</u>	
<u>TABLET; ORAL</u>		<u>EQ 150MG BASE</u>	
CIN-QIN SOLVAX		<u>EQ 300MG BASE</u>	
+ @ @		<u>EQ 150MG BASE</u>	
100MG	N85299 001 N84932 001 N85298 001	AB	N74488 001
200MG	N85299 001	AB	JUL 31, 1997
300MG	N85298 001	AB	N74488 002
100MG	N85299 001	AB	JUL 31, 1997
200MG	N84932 001	AB	N74488 002
300MG	N85298 001	AB	JUL 31, 1997
	N85299 001	TORPHARM	N74680 001
	N84932 001	AB	SEP 12, 1997
	N85298 001	AB	N74680 002
	N84549 001	AB	SEP 12, 1997
	N84549 001	AB	SEP 12, 1997

RANITIDINE HYDROCHLORIDE

TABLET; ORAL ZANTAC 150 <u>AB</u>	<u>EQ 150MG BASE</u>	N118703 001 JUN 09, 1983
<u>AB + GLAXO WELLCOME</u>	<u>EQ 300MG BASE</u>	N118703 002 DEC 09, 1985

RANITIDINE HYDROCHLORIDE

TABLET; ORAL ZANTAC 300 <u>AB</u>	<u>EQ 300MG BASE</u>	N118703 002 DEC 09, 1985
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RIFAMPIN

CAPSULE; ORAL <u>RIFAMPIN</u> AB	300MG EON	N64150 001 MAY 28, 1997
TABLET; ORAL REQUP + SMITHKLINE BEECHAM	EQ 0.25MG BASE EQ 0.5MG BASE	N20658 001 SEP 19, 1997

> DLT > > DLT > > DLT > > ADD > > ADD >	RAUWOLFIA SERPENTINA PAL PAK + @ @	50MG 100MG 50MG 100MG
> DLT > > ADD >	RAUWOLFIA SERPENTINA DARSBURY PHARMA GLOBAL PHARM BP	50MG 50MG 100MG 50MG
BP	GLOBAL PHARM	100MG
BP	DARSBURY PHARMA	50MG
③	BP	50MG
③	ZENITH GOLDLINE	100MG
③	ZENITH LABS	50MG
③	ZENITH LABS	100MG

RESERPINE

TABLET; ORAL RESERPINE PUR-E-PAC PHARM IP	0.1MG 0.25MG 0.1MG 0.25MG 0.25MG 0.1MG 0.25MG 0.25MG 0.1MG 0.25MG	N80753 002 N80753 003 N80753 002 N80753 001 N85207 001 N85207 001 N11185 001 N11185 002 N11185 001 N11185 002
③	TABLETCAPS	50mC ₁ /ML
③	ZENITH GOLDLINE	
③	ZENITH LABS	
③	ZENITH LABS	

SAMARIUM SM 153 LEXILDRONAM PENTASODIUM

INJECTABLE; INJECTION QUADRAMET CYTOGEN	N20570 001 MAR 28, 1997
SAQUINAVIR MESYLATE	

CAPSULE; ORAL FORTOVASE ROCHE

N20692 001
NOV 07, 1997

SCOPOLAMINE

FILM, EXTENDED RELEASE; TRANSDERMAL
 > DLT >
 * CIBA
 + NOVARTIS
 > ADD >

N17874 001
 N17874 001
 0.5MG/24HR
 1MG/72HR
 SEP 26, 1997

SECOBARBITAL SODIUM

CAPSULE; ORAL
 SECOBARBITAL SODIUM
 ® ZENITH GOLDLINE
 ® ZENITH LABS
SODIUM SECOCARBITAL
 HALSEY
 ®
 AA
 > ADD >

N85869 001
 N85869 001
 100MG
 100MG
100MG
 100MG
 N84676 001
 N84676 001
 100MG
 100MG
 > ADD >

SELEGILINE HYDROCHLORIDECAPSULE; ORALELDEPRYLSMGSMG+TABLET; ORALSELEGILINE HCLALPHAPHARMSMGABAPOTEXSMGABAPOTHECONSMGABENDO LABSSMGABENDO PHARMSSMGABTEVAAA> ADD >> ADD >AAAAAAAAAAAAAAAAAASERMORELIN ACETATE

INJECTABLE; INJECTION
 GEREF
 + SERONO

EQ 1MG BASE/VIAL

N20443 001

SEP 26, 1997

SIBUTRAMINE HYDROCHLORIDE

> ADD >				
CAPSULE; ORAL MERIDIA KNOLL PHARM				
N85869 001 N85869 001 100MG 100MG	N85869 001 N85869 001 100MG 100MG	N84676 001 N84676 001 100MG 100MG	N84676 001 N84676 001 100MG 100MG	N84676 001 N84676 001 100MG 100MG

SODIUM CHLORIDE

INJECTABLE; INJECTION
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER
 ABBOTT
 AP
 9MG/ML

N19465 002

JUL 15, 1985

SODIUM IODIDE, I-123

CAPSULE; ORAL GOLDEN PHARMS				
AA AA AA AA AA	AA AA AA AA AA	AA AA AA AA AA	AA AA AA AA AA	AA AA AA AA AA
N74866 001 N74871 001 JUN 06, 1997 N74672 001 APR 01, 1997 N74565 001 AUG 02, 1996 N74565 001 AUG 02, 1996 N74744 001 JAN 27, 1997	N74866 001 N74871 001 JUN 06, 1997 N74672 001 APR 01, 1997 N74565 001 AUG 02, 1996 N74565 001 AUG 02, 1996 N74744 001 JAN 27, 1997	N74866 001 N74871 001 JUN 06, 1997 N74672 001 APR 01, 1997 N74565 001 AUG 02, 1996 N74565 001 AUG 02, 1996 N74744 001 JAN 27, 1997	N74866 001 N74871 001 JUN 06, 1997 N74672 001 APR 01, 1997 N74565 001 AUG 02, 1996 N74565 001 AUG 02, 1996 N74744 001 JAN 27, 1997	N74866 001 N74871 001 JUN 06, 1997 N74672 001 APR 01, 1997 N74565 001 AUG 02, 1996 N74565 001 AUG 02, 1996 N74744 001 JAN 27, 1997

SERMORELIN ACETATE

INJECTABLE; INJECTION
 GEREF
 + SERONO

EQ 0.5MG BASE/VIAL

N20443 001

SEP 26, 1997

400 uCl
 N18671 003
 MAY 27, 1982

400 uCl
 N18671 003
 MAY 27, 1982

<u>SODIUM IODIDE, I-131</u>		<u>SPIRONOLACTONE</u>	
CAPSULE; ORAL IODOTOPIC BRACCO	8-100 uCi 1-50mCi 8-100 uCi 1-130mCi	N10929 001 N10929 003 N10929 001 N10929 003	AB <u>ZENITH LABS</u> <u>25MG</u>
<u>SODIUM NITROPRUSSIDE</u>			
INJECTABLE; INJECTION <u>NITROPRESS</u> AP ABBOTT ④		N71555 001 NOV 16, 1987 N71555 001 NOV 16, 1987	> DLT > > DLT > > ADD > > ADD >
<u>SOMATOTROPIN, BIOSYNTHETIC</u>			
INJECTABLE; INJECTION NORDITROPIN BX + NOVO NORDISK BX SERONI		N19721 001 MAY 08, 1995 N20604 003 JUL 25, 1997	
<u>SOTALOL HYDROCHLORIDE</u>			
TABLET; ORAL BETAPACE BERLEX		N19865 001 OCT 30, 1992 N19865 003 OCT 30, 1992 N19865 001 OCT 30, 1992 N19865 003 OCT 30, 1992	> DLT > > DLT > > DLT > > ADD > > ADD > > ADD > > ADD >
<u>SPIRONOLACTONE</u>			
TABLET; ORAL <u>SPIRONOLACTONE</u> ④ ZENITH GOLDLINE		N87108 001 25MG	
<u>SULFINPYRAZONE</u>			
TABLET; ORAL <u>SULFINPYRAZONE</u> ④ ZENITH GOLDLINE		N87769 001 100MG	

SULFINPYRAZONE

TABLET; ORAL
SULFINPYRAZONE
 AB ZENITH LABS

100MG

N87769 001
 JUN 01, 1992

SULFISOXAZOLE

TABLET; ORAL
SULFISOXAZOLE
 AB [®]UREPAC PHARM

500MG
 500MG
 500MG
 500MG

N80087 001
 N80087 001
 N80142 001
 N80142 001

AB + ZENITH GOLDLINE
 AB * ZENITH LABS

SULFISOXAZOLE DIOLAMINE

SOLUTION/DROPS; OPHTHALMIC
GANTIVISIN
 + ROCHE

EQ 4% BASE
 EQ 4% BASE

SUMATRIPTAN

SPRAY; NASAL
IMITREX
 + GLAXO WELLCOME

5MG/SPRAY
 10MG/SPRAY

20MG/SPRAY

TAMOXIFEN CITRATE

TABLET; ORAL
NOLVADEX
 + ZENECA

EQ 10MG BASE
 EQ 10MG BASE

N20579 001
 APR 15, 1997

N20600 001
 JUN 13, 1997

N20600 002
 JUN 13, 1997

TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL
FLOMAX

+ BOEHRINGER INGELHEIM

0.4MG
 JUN 01, 1992

TAZAROTENE

GEL; TOPICAL
TAZORAC
 ALLERGAN

0.05%
 N80087 001
 N80142 001
 N80142 001

+
 N80142 001

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION
AN-MDA
 @ CIS
 SORIN

N/A
 N/A
 N/A

TECHNETIUM TC-99M ALBUMIN KIT

INJECTABLE; INJECTION
TECHNETIUM TC 99M HSA
 MEDI PHYSICS
 NYCOMED AMERSHAM

N/A
 N/A
 N/A

TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION
CERETEC
 AMERSHAM

N19829 001
 DEC 30, 1988

N19829 001
 DEC 30, 1988

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION
MDP-BRACCO
 AB
 BRACCO

N/A

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION
MDP-SQUIBB N/A
BRACCIO N/A
TECHNETIUM TC 99M MPI MDP N/A
MEDI PHYSICS N/A
NYCOMED AMERSHAM N/A
AP >
AP >
AP >

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION
MPI DTPA KIT - CHELATE N/A
MEDI PHYSICS N/A
NYCOMED AMERSHAM N/A
TECHNETIUM TC-99M PENTETATE KIT
MEDI PHYSICS N/A
NYCOMED AMERSHAM N/A
AP >
AP >
AP >
AP >

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION
MIRALUMA N/A
DUPONT N/A
AP >
AP >

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL
TECHNETIUM TC 99M GENERATOR N17693 001
MEDI PHYSICS 830-16, 600mCi/GENERATOR
NYCOMED AMERSHAM 830-16, 600mCi/GENERATOR
AP >
AP >

TECHNETIUM TC-99M SUCCIMER KIT

INJECTABLE; INJECTION
MPI DMSA KIDNEY REAGENT N/A
MEDI PHYSICS N/A
NYCOMED AMERSHAM N/A
AP >
AP >
AP >

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION
MYOVIEW N/A
MEDI PHYSICS N/A
NYCOMED AMERSHAM N/A
AP >
DLT >
DLT >
ADD >
ADD >

TERBINAFINE HYDROCHLORIDE

SOLUTION; TOPICAL
LAMISIL 1%
+ NOVARTIS

N17255 001
N17255 001

N17264 002
N17264 002

TERFENADINE

TABLET; ORAL
SELDANE 60MG
AB + HOECHST MARION RSSL 60MG
TERFENADINE

N18949 001
MAY 08, 1985

N74475 001
JAN 03, 1997

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL
ANDRODERM
+ THERATECH 5MG/24HR

N20489 002
MAY 02, 1997

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL
TETRACYCLINE HCL
WARNER CHILCOTT

N62300 001
N62300 002
N62300 001
N62300 002

ACHROMYCIN

AP * LEDERLE

N50273 002
250MG/VIAL

TETRACYCLINE HYDROCHLORIDEINJECTABLE; INJECTION
ACHROMYCIN

AP + LEADERLE 500MG/VI_L
250MG/VI_L
500MG/VI_L

AP TETRACYCIN
PFIZER

AP AP 250MG/VI_L
500KG/VI_L
250MG/VI_L
500MG/VI_L

SUSPENSION; ORAL
TETRACYCLINE HCL

AB ALPHARMA 125MG/5ML
125MG/5ML
125MG/5ML
125MG/5ML

AB PUREPAC PHARM 125MG/5ML

AB TETRACYCIN
PEPPHARMECS 125MG/5ML
125MG/5ML

AB TETRAMED
ZENITH GOLDLINE

AB ZENITH LABS 125MG/5ML

THALLOUS CHLORIDE, TL-201

INJECTABLE; INJECTION
THALLOUS CHLORIDE TL 201
DUPONT

AP + MALLINCKRODT 1mCi/ML
1mCi/ML
1mCi/ML
1mCi/ML

AP + MEDIA PHYSICS 1mCi/ML

AP + NYCOMED AMERSHAM 1mCi/ML

THEOPHYLLINECAPSULE, ORAL
THEOPHYLLINE

N50273 003 BP NB5263 001
N50273 002 BP NB5263 002
N50273 003 @ NB5263 001

N60096 001 BP NB7763 001
N60096 002 BP NB7763 001
N60096 001 NB7763 001
N60096 002 NB7763 001

N60096 002 BC NB7763 001
N60096 001 BC NB7763 001
N60096 002 BC NB7763 001

N60633 001 BC NB7763 001
N60633 001 BC NB7763 001
N60291 001 BC NB7763 001
N60291 001 BC NB7763 001

N60095 001 BC NB7763 001
N60095 001 BC NB7763 001

N61468 001 BC NB7763 001
N61468 001 BC NB7763 001

THEO-DUR
KEY PHARMS BC NB7763 001

THEO-DUR
KEY PHARMS BC NB7763 001

N17806 001 BC NB8015 001
N17806 001 BC NB8015 001
N18150 001 BC NB8015 001
N18150 001 BC NB8015 001

N18110 002 BC NB8015 001
FEB 27, 1996 BC NB8015 001
N18110 002 BC NB8015 001

N18110 002 BC NB8015 001

THEOPHYLLINE BC NB8015 001

CAPSULE; ORAL
BRONKODYL BP NB8016 001
STERLING WINTHROP BP NB8016 001

100MG BP NB8016 001
200MG BP NB8016 001
100MG BP NB8016 001
200MG BP NB8016 001

N85264 001 BP NB8016 001
N85264 002 BP NB8016 002
N85264 001 BP NB8016 001
N85264 002 BP NB8016 002

THIAMINE HYDROCHLORIDE

<u>INJECTABLE; INJECTION</u>	
<u>THIAMINE HCL</u>	
<u>AB</u>	<u>1.00MG/ML</u>
<u>AP</u>	<u>1.00MG/ML</u>
<u>AP</u>	<u>SANOFI WINTHROP</u>
<u>AP</u>	<u>STERIS</u>
<u>AP</u>	<u>(@)</u>
	<u>2.00MG/ML</u>
	<u>1.00MG/ML</u>
	<u>2.00MG/ML</u>

TIMOLOL

<u>SOLUTION/DROPS; OPHTHALMIC</u>	
<u>BETIMOL</u>	
<u>* LIBRAS</u>	
	<u>EQ 0.25% BASE</u>
<u>N40079 001</u>	
<u>MAY 03, 1996</u>	
<u>N40079 001</u>	
<u>MAY 03, 1996</u>	
<u>N83534 001</u>	
<u>N83534 002</u>	
<u>N83534 001</u>	
<u>N83534 002</u>	
<u>EQ 0.5% BASE</u>	
<u>EQ 0.25% BASE</u>	
<u>OY STAR</u>	
<u>+</u>	
<u>EQ 0.5% BASE</u>	
<u>N20439 001</u>	
<u>MAR 31, 1995</u>	
<u>N20439 002</u>	
<u>MAR 31, 1995</u>	
<u>N20439 001</u>	
<u>MAR 31, 1995</u>	
<u>N20439 002</u>	
<u>MAR 31, 1995</u>	

THIORDIAZINE HYDROCHLORIDE

<u>CONCENTRATE; ORAL</u>	
<u>THIORDIAZINE HCL</u>	
<u>AA PHARM ASSOC</u>	<u>3.0MG/ML</u>

<u>THIOTROPIN</u>	
<u>INJECTABLE; INJECTION</u>	
<u>THYTROPAR</u>	<u>10 IU/VIAL</u>
<u>* ARMOUR PHARM</u>	<u>10 IU/VIAL</u>
<u>@ RHONE POULENC</u>	<u>10 IU/VIAL</u>

TIGAGABINE HYDROCHLORIDE

<u>TABLET; ORAL</u>	
<u>GABITRIAL</u>	
<u>ABBOTT</u>	
<u>4MG</u>	
<u>1.2MG</u>	
<u>1.6MG</u>	
<u>2.0MG</u>	
<u>+</u>	
<u>N20646 001</u>	
<u>SEP 30, 1997</u>	
<u>N20646 002</u>	
<u>SEP 30, 1997</u>	
<u>N20646 003</u>	
<u>SEP 30, 1997</u>	
<u>N20646 004</u>	
<u>SEP 30, 1997</u>	

<u>TILDUDRONATE DISODIUM</u>	
<u>TABLET; ORAL</u>	
<u>SKELID</u>	
<u>+ SANOFI</u>	

<u>TABLET; ORAL</u>	
<u>EQ 200MG BASE</u>	
<u>N20707 001</u>	

MAR 07, 1997

N19355 001
DEC 30, 1986

6.5%
* BRISTOL MYERS

TIOPRONIN

TABLET; ORAL
TIOPRONIN
+ MISSION PHARMA
④ UNITV TX

100MG
N19569 001
AUG 11, 1988
N19569 001
AUG 11, 1988

N74357 001
APR 30, 1997

TRAZODONE HYDROCHLORIDE

TABLET; ORAL
TRAZODONE HCL
AB TEVA

150MG

TOPIRAMATE

TABLET; ORAL
TOPRAMATE
AB PUREPAC PHARM
④ TEVA

400MG
EQ 600MG BASE

400MG
N88950 001
JUN 17, 1985
N88950 001
JUN 17, 1985

N20404 003
JAN 14, 1997

TABLET; ORAL
TOPRAMATE
AB ZENITH GOLDLINE
AB ZENITH GOLDLINE

400MG
EQ 600MG BASE
EQ 600MG BASE
EQ 600MG BASE
EQ 600MG BASE

400MG
N74399 001
MAR 28, 1996
N74399 001
FEB 27, 1997
N74399 001
FEB 27, 1997
N74399 001
MAR 28, 1996

N19049 001
SEP 16, 1988

TABLET; ORAL
TOPRAMAX
④ JOHNSON RW

400MG
N20505 006
DEC 24, 1996

N4039 001
NOV 26, 1997

TABLET; ORAL
TOPRAMAX
④ JOHNSON RW

400MG
N20497 001
MAY 29, 1997

N87192 001
SEP 08, 1982

TABLET; ORAL
TOPRAMAX
④ JOHNSON RW

400MG
N20497 001
MAY 29, 1997

N87192 001
SEP 08, 1982

TRETINOIN

CREAM; TOPICAL
AVITA
AB PENEDERM

0.025%

N20404 003
JAN 14, 1997

0.025%

N19049 001
SEP 16, 1988

0.025%

N20475 001
FEB 07, 1997

0.1%

TRIACINOLONE

TABLET; ORAL
TRIACINOLONE
BP PUREPAC PHARM

200G

N84020 002
N84020 003
N84020 002
N84020 003
N83750 001
N83750 001

TABLET; TOPICAL
RETIN-A MICRO
+ ADV POLYMER
BP TARO

0.1%

N20475 001
FEB 07, 1997

TRIACINOLONE ACETONIDE

CREAM; TOPICAL
TRIACINOLONE ACETONIDE
AT TARO

0.1%

N4039 001
NOV 26, 1997

LOTION; TOPICAL
TRIACINOLONE ACETONIDE
AT ALPHARMA

0.1%

N87192 001
SEP 08, 1982

TABLET; ORAL

TOPRAMAX

AB ZENITH GOLDLINE

N87192 001
SEP 08, 1982

TABLET; ORAL

TOPRAMAX

AB ZENITH GOLDLINE

N87192 001
SEP 08, 1982

<u>TRIACINOLONE DIACETATE</u>		<u>TRIMETHOPRIM</u>	
INJECTABLE; INJECTION		TABLET; ORAL	
TRIACINOLONE DIACETATE		PROLOPRIM	
BP STERIS @	4.0MG/ML	GLAXO WELLCOME	2.00MG
	4.0MG/ML		
		AB +	2.00MG
<u>TRIFLUOPERAZINE HYDROCHLORIDE</u>		<u>TRIMPEx 200</u>	2.00MG
		AB + ROCHE	
<u>TABLET; ORAL</u>		(@)	2.00MG
<u>TRILOUOPERAZINE HCL</u>			
AB MYLAN	EQ 1MG BASE	N40209 001	
	EQ 2MG BASE	JUL 07, 1997	
AB	EQ 5MG BASE	N40209 002	
	EQ 10MG BASE	JUL 07, 1997	
		N40209 003	
		JUL 07, 1997	
		N40209 004	
		JUL 07, 1997	
<u>TRIHEXYPHENIDYL HYDROCHLORIDE</u>			
<u>CAPSULE, EXTENDED RELEASE; ORAL</u>			
ARTANE + LEDERLE @	5MG 5MG	N12947 001	
		N12947 001	
ELIXIR; ORAL			
<u>TRIHEXYPHENIDYL HCL</u>	2MG/5ML	N40177 001	
MA PHARM ASSOC		APR 17, 1997	
<u>TRIMETHOBENZAMIDE HYDROCHLORIDE</u>		<u>UREA, C-14</u>	
INJECTABLE; INJECTION		CAPSULE; ORAL	
<u>TRIMETHOBENZAMIDE HCL</u>	100MG/ML	PYTEST	
AB Abbott		+ TRI MED SPECLTS	1 uCi
AP STERLING WINTHROP	100MG/ML	PYTEST KIT	
		+ TRI MED SPECLTS	1 uCi

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION
LYPHOCIN

FUJISAWA

EQ 500MG BASE/VIAL

AP

N62663 001

MAR 17, 1987

N20699 004
OCT 20, 1997

EQ 1GM BASE/VIAL

AP

N62663 002

JUL 31, 1987

N20699 002
OCT 20, 1997

EQ 5GM BASE/VIAL

AP

N62663 003

JUN 03, 1988

N20699 004
OCT 20, 1997

+ VANCOCIN HCL IN PLASTIC CONTAINER
+ BAXTER HLTHCARE

EQ 500MG BASE/100ML

AP

N50671 001

APR 29, 1993

N70577 001
FEB 02, 1987

EQ 500MG BASE/100ML

AP

N50671 001

APR 29, 1993

N70577 001
FEB 02, 1987

EQ 500MG BASE/VIAL

AP

N62682 001

JUL 22, 1986

N19152 003
MAR 06, 1991

EQ 500MG BASE/VIAL

AP

N62682 001

JUL 22, 1986

N74330 001
JAN 31, 1994

EQ 1GM BASE/VIAL

AP

N62682 002

MAR 30, 1988

N73568 001
JUL 31, 1992

EQ 1GM BASE/VIAL

AP

N62682 002

MAR 30, 1988

N74587 002
FEB 21, 1997

EQ 10GM BASE/VIAL

AP

N62682 005

MAY 11, 1988

N74587 003
SEP 09, 1997

EQ 10GM BASE/VIAL

AP

N62682 005

MAY 11, 1988

N73568 002
OCT 10, 1997

EQ 10GM BASE/VIAL

AP

N62682 005

MAY 11, 1988

N74330 001
JAN 31, 1994

EQ 500MG BASE/VIAL

AP

N62663 001

MAR 17, 1987

N73568 001
JUL 31, 1992

EQ 1GM BASE/VIAL

AP

N62663 002

JUL 31, 1987

N73568 002
OCT 10, 1997

EQ 5GM BASE/VIAL

AP

N62663 003

JUN 03, 1988

N73568 001
JAN 31, 1994

EQ 10GM BASE/VIAL

AP

N62663 004

NOV 28, 1997

N73568 001
JUL 31, 1992

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
EFFEXOR XR

+ WYETH AYERST

EQ 37.5MG BASE

AP

N20699 001

OCT 20, 1997

N20699 002
OCT 20, 1997

EQ 75MG BASE

AP

N20699 002

OCT 20, 1997

N20699 003
OCT 20, 1997

EQ 100MG BASE

AP

N20699 003

OCT 20, 1997

N20699 003
OCT 20, 1997

VINCRISTINE SULFATE

INJECTABLE; INJECTION
VINCRITEX

AP

* BRISTOL MYERS

5MG/VIAL

EQ 5MG/VIAL

@ BRISTOL MYERS SQUIBB

5MG/VIAL

VINCRISTINE SULFATE

FULDING

5MG/VIAL

EQ 5MG/VIAL

@

N70867 001
JUL 12, 1988

N70867 001
JUL 12, 1988

N71561 001
APR 11, 1988

VINCRISTINE SULFATE

INJECTABLE; INJECTION
VINCRITEX

AP

* BRISTOL MYERS

5MG/VIAL

EQ 5MG/VIAL

@ BRISTOL MYERS SQUIBB

5MG/VIAL

VINCRISTINE SULFATE

FULDING

5MG/VIAL

EQ 5MG/VIAL

@

N70867 001
JUL 12, 1988

N70867 001
JUL 12, 1988

N71561 001
APR 11, 1988

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL
DYNAX-HEX
XTTRIUM

0.75%

N20111 001
SEP 11, 1997

TINCTURE; TOPICAL
HIBITANE
© ZENECA

0.5%

N18049 001

CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL
CHLORPHENIRAMINE MALEATE
GENEVA PHARMS

1.2MG

N70797 001
AUG 12, 1988

+
TELDRIN
* SMITHKLINE

1.2MG

N70797 001
AUG 12, 1988

1.2MG
1.2MG

N17369 002
N17369 002

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
COLD CAPSULE IV
+ GRAHAM DM

1.2MG; 75MG

N18793 001
APR 25, 1985

1.2MG; 75MG
COLD CAPSULE V
GRAHAM DM

8MG; 75MG

N18794 001
APR 23, 1985

8MG; 75MG
+ APR 23, 1985

N18794 001
APR 23, 1985

TABLET, EXTENDED RELEASE; ORAL
CONTAC
NOVARTIS

1.2MG; 75MG

N19613 001
JUN 13, 1986

PHENYLPROPANOLAMINE HCL/CHLORPHENIRAMINE
SANDOZ

1.2MG; 75MG

N19613 001
JUN 13, 1986

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
CODIMAL-L.A. 1.2
* GENT PHARMS

1.2MG; 120MG

N18935 001
APR 15, 1985

@ SCHWARZ PHARMA
KV PHARM

1.2MG; 120MG

N18935 001
APR 15, 1985

PSEUDOEPHEDRINE HCL AND CHLORPHENIRAMINE MALEATE
@

1.2MG; 120MG

N18935 001
APR 15, 1985

CIMETIDINE

CIMETIDINE
TABLET; ORAL
TAGAMET HB

120MG

SMITHKLINE BEPHCIAH

100MG

@

N17369 002
JUN 19, 1995

100MG
N17369 002
JUN 19, 1995

CLEMASTINE FUMARATE

CLEMASTINE FUMARATE
LEMMON
TEVA

1.34MG

N73282 002
DEC 03, 1992

CLOTRIMAZOLE
+
LOTRIMIN AF
+ SCHERING PLough

1.34MG

N73282 002
DEC 03, 1992

CREAM; TOPICAL
LOTRIMIN AF
+ SCHERING PLough

1%

N17619 002
OCT 27, 1989

GYNE-LOTRIMIN 3 COMBINATION PACK
+ SCHERING PLough

1%; 200MG
N20526 002
JUL 29, 1996

CLOTRIMAZOLECLOTRIMAZOLE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

GYNE-LOTRIMIN COMBINATION PACK
* SCHERING PLOUGH
1%, 100MGN20289 002
APR 26, 1993
N20389 002
JUN 23, 1994CREAM, TABLET, TOPICAL, VAGINAL
GYNE-LOTRIMIN 3 COMBINATION PACK
+ SCHERING PLOUGH 1%, 200MGGYNE-LOTRIMIN COMBINATION PACK
+ SCHERING PLOUGH 1%, 100MGMYCELEX-7 COMBINATION PACK
BAKER 1%, 100MGTABLET; VAGINAL
GYNE-LOTRIMIN 3
+ SCHERING PLOUGH 200MG
N20526 002
JUL 29, 1996
N20289 002
APR 26, 1993
N20389 002
JUN 23, 1994TABLET; VAGINAL
GYNE-LOTRIMIN 3
+ SCHERING PLOUGH 200MG
N20525 001
DEC 26, 1991TABLET; VAGINAL
GYNE-LOTRIMIN 3
+ SCHERING PLOUGH 100MG
N18182 002
DEC 26, 1991CROMOLYN SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDESOLUTION; ORAL
BENTYLIN
+ PARKE DAVIS
12.5MG / 5ML; 30MG / 5ML
N19014 001
JUN 11, 1985SOLUTION; ORAL
PROVEL
@ SANDOZ
12.5MG / 5ML; 30MG / 5ML
N19014 001
JUN 11, 1985IBUPROFEN
N17613 002
OCT 27, 1989
N20890 001
OCT 27, 1989
> DLT >
> DLT >
> DLT >
> ADD >
> ADD >IBUPROFEN
CAPSULE; ORAL
PROVEL
@ SANDOZ
200MG
N20402 001
APR 20, 1995SUPPOSITORY; VAGINAL
CHILDREN'S ADVIL-FLAVORED
WHITEHALL ROBINS 100MG / 5ML
N20589 002
NOV 07, 1997SUPPOSITORY; VAGINAL
GYNE-LOTRIMIN
+ SCHERING PLOUGH
100MG
GYNE-LOTRIMIN 3
+ SCHERING PLOUGH
200MG
N17717 002
NOV 30, 1990
N20525 001
JUL 29, 1996
> ADD >
> ADD >
> ADD >TABLET; ORAL
IBUPROFEN
PUREPAC PHARM
200MG
N71122 001
OCT 03, 1986TABLET; VAGINAL
IBUPROFEN
PUREPAC PHARM
200MG
N711664 001
FEB 03, 1987
N71122 001
OCT 03, 1986TABLET; VAGINAL
GYNE-LOTRIMIN
+ SCHERING PLOUGH
100MG
N17717 002
NOV 30, 1990
@

IBUPROFEN

TABLET; ORAL

IBUPROFEN
⑧ PUREPAC PHARM

200MG

@ ZENITH GOLDLINE

200MG

⑧

200MG

>

PHARMACIA AND UPJOHN

200MG

&

JUNIOR STRENGTH MOTRIN
MCNEIL

100MG

+

100MG

>

NUPRIN
⑧ MCNEIL

200MG

>

VELOSULIN HUMAN
⑧ NOVO NORDISK

200MG

>

IBUPROFEN POTASSIUMCAPSULE; ORAL
PROVEL
⑧ NOVARTIS

200MG

>

KETOCONAZOLESHAMPOO; TOPICAL
NIZORAL A-D

J AND J

1%

N20310 001
OCT 10, 1997N71154 001
OCT 27, 1987LOPERAMIDE HYDROCHLORIDEN72040 001
APR 29, 1988N71154 001
OCT 27, 1987N72040 001
APR 29, 1988N72040 001
APR 29, 1988N20602 001
JUN 10, 1996N20602 001
JUN 10, 1996N19012 001
MAY 18, 1984N19012 003
JUL 29, 1987N19012 001
MAY 18, 1984N19012 003
JUL 29, 1987N20402 001
APR 20, 1995N20402 001
APR 20, 1995N19450 001
MAY 30, 1986N19450 001
MAY 30, 1986N20606 001
JUN 26, 1996N20448 001
JUL 24, 1997MICONAZOLE NITRATECREAM; VAGINAL
MICONAZOLE NITRATE
PERRIGO

2%

TARO

2%

N74760 001
MAY 15, 1997N7444 001
JAN 13, 1997CREAM, SUPPOSITORY; TOPICAL, VAGINAL
M-ZOLE 7 DUAL PACK
ALPHARMA

2%, 100MG

N74586 001
JUL 17, 1997N74414 001
APR 30, 1997
N74395 001
MAR 20, 1997INSULIN SEMISYNTHETIC PURIFIED HUMANINJECTABLE; INJECTION
VELOSULIN BR HUMAN
+ NOVO NORDISK

100 UNITS/ML

100 UNITS/ML

VELOSULIN HUMAN
+ NOVO NORDISK

100 UNITS/ML

100 UNITS/ML

<u>MINOXIDIL</u>		<u>SODIUM FLUORIDE; TRICLOSAN</u>	
SOLUTION; TOPICAL MINOXIDIL (FOR MEN) MORTON GROVE	2%	N74767 001 FEB 28, 1997	PASTE; DENTAL COLGATE TOTAL + COLGATE PALMOLIVE 0.24% + 0.3%
> ADD > > ADD > > ADD >	ROGAINE EXTRA STRENGTH (FOR MEN) + PHARMACIA AND UPJOHN 5%	N20834 001 NOV 14, 1997	JUL 11, 1997
			N20231 001 JUL 11, 1997
<u>NAPROXEN SODIUM</u>		<u>OINTMENT; VAGINAL</u>	
TABLET; ORAL NAPROXEN SODIUM GRANUTEC	EQ 200MG BASE	N74635 001 JAN 13, 1997	VAGISTAT-1 + BRISTOL MYERS SQUIBB 6.5%
INVAMED	EQ 200MG BASE	N74646 001 JAN 13, 1997	
NOVOPHARM	EQ 200MG BASE	N74635 001 JAN 13, 1997	
PERRIGO	EQ 200MG BASE	N74661 001 JAN 13, 1997	
PVT™ FORM	EQ 200MG BASE	N74789 001 FEB 27, 1997	
<u>PERMETHRIN</u>		<u>PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE</u>	
LOTION; TOPICAL NIX + WARNER LAMBERT	1%	N19918 001 MAY 02, 1990	CAPSULE, EXTENDED RELEASE, ORAL TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES KV PHARM 120MG, 5MG
* WARNER WELLCOME	1%	N19918 001 MAY 02, 1990	N71798 001 MAR 16, 1989
			N71798 001 MAR 16, 1989
			②

DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST
CUMULATIVE SUPPLEMENT NUMBER 11 / NOVEMBER '97

NO NOVEMBER 1997 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 Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
15AU81 TN=	Treatment of primary pulmonary hypertension.	Lung Rx, Inc. 2 Davis Drive P.O. Box 13169 Research Triangle Park, NC 27709 DD=06/04/1997
8 Cyclopentyl 1,3-dipropylxanthine TN=	Treatment of cystic fibrosis.	SciClone Pharmaceuticals, Inc. 901 Mariner's Island Boulevard Suite 315 San Mateo, CA 94404 DD=03/24/1997
9-cis-retinoic acid TN=	Prevention of retinal detachment due to proliferative vitreoretinopathy.	Allergan, Inc. 2525 Dupont Drive P.O. Box 19534 Irvine, CA 92623 DD=01/02/1997
Allogeneic peripheral blood mononuclear cells sensitized against patient alloantigens by mixed lymphocyte culture TN= CYTOIMPLANT	Treatment of pancreatic cancer.	Applied Immunotherapeutics, LLC 14132 E. Firestone Boulevard Santa Fe Springs, CA 90670 DD=06/13/1997
Alpha-melanocyte stimulating hormone TN=	Prevention and treatment of intrinsic acute renal failure due to ischemia.	Star, Robert A., M.D. UT-Southwestern Medical School 5323 Harry Hines Dallas, TX 75235 DD=08/19/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Anagrelide TN= Agrylin	Treatment of essential thrombocythemia.	Roberts Pharmaceutical Corp. Meridian Center III 6 Industrial Way West Eatonontown, NJ 07724 DD=01/27/1988 MA=03/14/1997
Ancrod TN=	To establish and maintain anticoagulation in heparin-intolerant patients undergoing cardiopulmonary bypass.	Knoll Pharmaceuticals 30 North Jefferson Road Whippany, NJ 07981 DD=11/12/1997
Apomorphine TN= Zydis	For use as rescue treatment for early morning motor dysfunction in late-stage Parkinson's disease.	Scherer DDS Frankland Road Swindon Wiltshire UK SN5 8RU, DD=10/20/1997
B2036-PEG TN= Trovert	Treatment of acromegaly.	Sensus Corporation Suite 430, 98 San Jacinto Boulevard Austin, TX 78701 DD=06/24/1997
Beta alethine TN= Betathine	Treatment of multiple myeloma.	Dovetail Technologies, Inc. Bldg. 337, Paint Branch Drive College Park, MD 20742 DD=03/24/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Beta alethine TN= Betathine	Treatment of metastatic melanoma.	Dovetail Technologies, Inc. Bldg. 337, Paint Branch Drive College Park, MD 20742 DD=03/24/1997
Busulfan TN= Spartaject	Treatment of primary brain malignancies.	Sparta Pharmaceuticals, Inc. 111 Rock Road Horsham, PA 19044 DD=07/07/1997
CAMPATH-1H TN=	Treatment of chronic lymphocytic leukemia.	L&I Partners, LP 11550 IH 10 West, Suite 300 San Antonio, TX 78230 DD=10/20/1997
CY-1503 TN= Cylexin	Treatment of neonates and infants undergoing cardiopulmonary bypass during surgical repair of congenital heart lesions.	Cytel Corporation 3525 John Hopkins Court San Diego, CA 92121 DD=07/18/1997
Calcium gluconate TN= Calgonate	For use as a wash for hydrofluoric acid spills on human skin.	Calgonate Corp. 190 Commerce Drive Warwick, RI 02886 DD=11/20/1997
Coagulation Factor IX (recombinant) TN= BeneFix	Treatment of hemophilia B.	Genetics Institute, Inc. 87 Cambridge Park Drive Cambridge, MA 02140 DD=10/03/1994 MA=02/11/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Cysteamine hydrochloride TN=	Treatment of corneal cystine crystal accumulation in cystinosis patients.	Sigma-Tau Pharmaceuticals, Inc. 800 South Frederick Avenue Gaithersburg, MD 20877 DD=08/19/1997
Dehydroepiandrosterone sulfate sodium TN=	To accelerate the re-epithelialization of donor sites in those hospitalized burn patients who must undergo autologous skin grafting.	Pharmadigm, Inc. 2401 Foothill Drive Salt Lake City, UT 84109 DD=01/28/1997
Dehydroepiandrosterone sulfate sodium TN=	Treatment of serious burns requiring hospitalization.	Pharmadigm, Inc. 2401 Foothill Drive Salt Lake City, UT 84109 DD=01/29/1997
Diazepam viscous solution for rectal administration TN=	For the management of selected, refractory, patients with epilepsy, on stable regimens of antiepileptic drugs (AEDs), who require intermittent use of diazepam to control bouts of increased seizure activity.	Athena Neurosciences, Inc. 800f Gateway Boulevard South San Francisco, CA 94080 DD=02/25/1992 MA=07/29/1997
Dimethylsulfoxide e TN=	Topical treatment for the prevention of soft tissue injury following extravastion of cytotoxic drugs.	Cancer Technologies, Inc. 7301 East 22nd Street Suite 10E Tucson, AZ 85710 DD=04/15/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Enadoline hydrochloride TN=	Treatment of severe head injury.	Warner-Lambert Company Parke-Davis Pharmaceutical Research Division 2800 Plymouth Road Ann Arbor, MI 48105 DD=01/28/1997
Fampridine TN=	Treatment of chronic, incomplete spinal cord injury.	Acorda Therapeutics, Inc. 145 West 58th Street Suite 8J New York, NY 10019 DD=06/02/1997
Gp100 adenoviral gene therapy TN=	Treatment of metastatic melanoma.	Genzyme Corporation P.O. Box 9322 One Mountain Road Framingham, MA 01701 DD=03/25/1997
Human retinal pigmented epithelial cells on collagen microcarriers TN= Spheramine	Treatment of Hoehn and Yahr stage 3 and 4 Parkinson's disease.	Theracell, Inc. 50 Division Street, Suite 503 Somerville, NJ 08876 DD=07/18/1997
Icodextrin 7.5% with Electrolytes Peritoneal Dialysis Solution TN= Extraneal (with 7.5% Icodextrin) Peritoneal Dialysis Solutio	Treatment of those patients having end stage renal disease and requiring peritoneal dialysis treatment.	Baxter Healthcare Corporation Renal Division 1620 Waukegan Road Waukegan, IL 60085 DD=07/18/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Lepirudin TN= Refluden	Treatment of heparin-associated thrombocytopenia Type II.	Behringwerke AG P.O. Box 1140 D-35001 Marburg Germany, DD=02/13/1997
Levocarnitine TN= Carnitor	Treatment of zidovudine-induced mitochondrial myopathy.	Sigma-Tau Pharmaceuticals, Inc. 800 S. Frederick Avenue, Suite 300 Gaithersburg, MD 20877 DD=04/07/1997
MART-1 adenoviral gene therapy for malignant melanoma TN=	Treatment of metastatic melanoma.	Genzyme Corporation One Kendall Square Cambridge, MA 02139 DD=03/28/1997
Oxandrolone TN= Oxandrin	Treatment of patients with Duchenne's muscular dystrophy and Becker's muscular dystrophy.	Bio-Technology General Corp. 70 Wood Avenue South Iselin, NJ 08830 DD=04/22/1997
Paclitaxel TN= Taxol	Treatment of AIDS-related Kaposi's sarcoma.	Bristol-Myers Squibb Pharmaceutical Research Institute 5 Research Parkway P.O. Box 5100 Wallingford, CT 06492 DD=03/25/1997 MA=08/04/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Paclitaxel TN= Paxene	Treatment of AIDS-related Kaposi's sarcoma.	Baker Norton Pharmaceuticals, Inc. 4400 Biscayne Boulevard Miami, FL 33137 DD=04/15/1997
Patul-end TN=	Treatment of patulous eustachian tube.	Ear Foundation 24209 Castillo Street, Suite 100 Santa Barbara, CA 93105 DD=02/18/1997
Pegylated recombinant human megakaryocyte growth and development factor TN= MEGAGEN	For reducing the period of thrombocytopenia in patients undergoing hematopoietic stem cell transplantation.	Amgen, Inc. 1840 DeHavilland Drive Thousand Oaks, CA 91320 DD=10/20/1997
Pergolide TN= Permax	Treatment of Tourette's syndrome.	Sallee, Floyd R. M.D., Ph.D. Medical University of South Carolina 171 Ashley Avenue, Room PH246 Charleston, SC 29425 DD=11/20/1997
Poloxamer 188 TN=	Treatment of vasospasm in subarachnoid hemorrhage patients following surgical repair of a ruptured cerebral aneurysm.	CytRx Corporation 154 Technology Parkway Norcross, GA 30092 DD=08/05/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Poly-ICLC TN=	Treatment of primary brain tumors.	Salazar, Andres M. M.D. and Levy, Hilton B. Ph.D. 3202 Cleveland Avenue N.W. Washington, DC 20008 DD=03/17/1997
Porcine Sertoli cells aseptically prepared for intracerebral co-implantation with fetal neural tissue TN= N-Graft	Treatment of Hoehn and Yahr stage four and five Parkinson's disease.	Theracell, Inc. 50 Division Street Suite 503 Somerville, NJ 08876 DD=06/24/1997
Porfiromycin TN= Promycin	Treatment of cervical cancer.	Vion Pharmaceuticals, Inc. Four Science Park New Haven, CT 06511 DD=03/13/1997
Purified extract of Pseudomonas aeruginosa TN= ImmuDyn	Treatment of immune thrombocytopenia purpura where it is required to increase platelet counts.	DynaGen, Inc. 99 Erie Street Cambridge, MA 02139 DD=09/22/1997
RGG0853, E1A lipid complex TN=	Treatment of ovarian cancer.	Targeted Genetics Corporation 1100 Olive way, Suite 100 Seattle, WA 98101 DD=10/27/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Recombinant human acid alpha-glucosidas e TN=	Treatment of glycogen storage disease type II.	Chen, Y.T., M.D., Ph.D. Department of Pediatrics Duke University Medical Center Durham, NC 27710 DD=08/19/1997
Recombinant human alpha-L-iduronid ase TN=	Treatment of patients with mucopolysaccharidosis-I.	BioMarin Pharmaceuticals, Inc. 11 Pimental Court Novato, CA 94949 DD=09/24/1997
Recombinant human interleukin-12 TN=	Treatment of renal cell carcinoma.	Genetics Institute 87 CambridgePark Drive Cambridge, MA 02140 DD=10/20/1997
Recombinant human thrombopoietin TN=	For use in accelerating platelet recovery in patients undergoing hematopoietic stem cell transplantation.	Genentech, Inc. 1 DNA Way South San Francisco, CA 94080 DD=09/29/1997
Retroviral vector, R-GC and GC gene 1750 TN=	Treatment of Gaucher disease.	Genzyme Corporation One Kendall Square Cambridge, MA 02139 DD=05/06/1997
Sermorelin acetate TN= Geref	Treatment of idiopathic or organic growth hormone deficiency in children with growth failure.	Serono Laboratories, Inc. 100 Longwater Circle Norwell, MA 02061 DD=09/14/1988 MA=09/26/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Short chain fatty acid enema TN= Colomed	Treatment of chronic radiation proctitis.	Orphan Medical, Inc. 13911 Ridgedale Drive Suite 475 Minnetonka, MN 55305 DD=08/19/1997
Somatropin TN= Genotropin	Treatment of adults with growth hormone deficiency.	Pharmacia & Upjohn 7000 Portage Road Kalamazoo, MI 49001 DD=09/06/1994 MA=10/31/1997
Suramin TN=	Treatment of metastatic hormone-refractory prostate cancer.	Warner-Lambert Company Parke-Davis Pharmaceutical Research Division 2800 Plymouth Road Ann Arbor, MI 48105 DD=05/06/1997
Testosterone TN= TheraDerm Testosterone Transdermal System	For use as physiologic testosterone replacement in androgen deficient HIV+ patients with an associated weight loss.	TheraTech, Inc. 417 Wakara Way Salt Lake City, UT 84108 DD=09/22/1997
Toremifene TN= Fareston	Hormonal therapy of metastatic carcinoma of the breast.	Orion Corporation P.O. Box 65 02101 ESPOO Finland, DD=09/19/1991 MA=05/29/1997

Orphan Product Designations and Approvals List
January 1997 through November 1997

Name
Generic Name
TN=Trade Name

Indication Designated

Sponsor & Address
DD= Date Designated
MA=Marketing Approval

Zinc acetate Treatment of Wilson's disease.
TN= Galzin

Lemmon Company
1510 Delp Drive
Kulpsville, PA 19443
DD=11/06/1985
MA=01/28/1997

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

NO NOVEMBER 1997 ADDITIONS

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, 17TH EDITION FOR A FULL LISTING OF PATENT AND EXCLUSIVITY TERMS (ABBREVIATIONS, NEW DOSING SCHEDULE, NEW INDICATIONS AND PATENT USE CODES). ONLY NEW CODES WILL BE ADDED TO THE CUMULATIVE SUPPLEMENT.

REFERENCES NEW DOSING SCHEDULE

- D-33 ONCE DAILY DOSING FOR PLAQUE PSORIASIS
- D-34 EVERY FOUR MONTHS DOSAGE REGIMEN
- D-35 FOR A ONE WEEK DOSING PERIOD FOR INTERDIGITAL TINEA PEDIS
- D-36 FOR A SINGLE 2MG DOSE AS AN ALTERNATIVE TO THE 1MG DOSE GIVE TWICE DAILY

NEW INDICATION

- I-177 TREATMENT OF MODERATE ACNE VULGARIS IN FEMALES, GREATER OR EQUAL TO 15 YEARS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, DESIRE CONTRACEPTION, HAVE ACHIEVED MENARCHE AND ARE UNRESPONSIVE TO TOPICAL ANTI-ACNE MEDICATIONS
- I-178 TREATMENT OF ONCHOMYCOSIS OF THE FINGERNAIL WITHOUT CONCOMITANT ONCHOMYCOSIS OF THE TOENAIL WITH A PULSE DOSING REGIMEN
- I-179 NOSOCOMIAL PNEUMONIA-MILD TO MODERATE AND SEVERE CAUSED BY HAEMOPHILUS INFLUENZAE OR KLEBSIELLA PNEUMONIAE
- I-180 TREATMENT OF PLANTAR TINEA PEDIS (MOCCASIN TYPE)
- I-181 TREATMENT OF PATIENTS WITH COMPLEX PARTIAL SEIZURES WITH AND WITHOUT SECONDARY GENERALIZATION
- I-182 TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME
- I-183 MAINTENANCE THERAPY IN THE MANAGEMENT OF MILD TO MODERATE ASTHMA IN PEDIATRIC PATIENTS AGES 6-11
- I-184 TREATMENT OF PANIC DISORDER AT A RECOMMENDED DOSE RANGE OF 1 TO 2 MG/DAY (MAXIMUM OF 4MG)
- I-185 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-186 TREATMENT OF TINEA (PITYRIASIS) VERSCOLOR CAUSED BY OR PRESUMED TO BE CAUSED BY PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE)
- I-187 PREVENTION OF FRACTURES IN THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-188 TREATMENT OF ACUTE SINUSITIS AND ACUTE EXACERBATION OF CHRONIC SINUSITIS
- I-189 TREATMENT OF ACUTE OTITIS MEDIA IN PEDIATRIC PATIENTS
- I-190 PLANAR IMAGING AS A SECOND LINE DIAGNOSTIC DRUG AFTER MAMMOGRAPHY TO ASSIST IN THE EVALUATION OF BREAST LESIONS IN PATIENTS WITH AN ABNORMAL MAMMOGRAM OR A PALPABLE BREAST MASS
- I-191 ENDOMETRIAL THINNING AGENT PRIOR TO ENDOMETRIAL ABLATION FOR DYSFUNCTIONAL UTERINE BLEEDING
- I-192 THE PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS AND A NEW DOSAGE REGIMEN, 40MG ONCE DAILY, FOR THIS INDICATION
- I-193 TREATMENT OF PANIC DISORDER IN A RECOMMENDED DOSE RANGE OF 50 TO 200MG/DAY
- I-194 CONGESTIVE HEART FAILURE
- I-195 USE OF LANSOPRAZOLE IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF DUODENAL ULCER

**PATENT AND EXCLUSIVITY TERMS
NEW INDICATION**

- I-196 ACUTE TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- I-197 MAINTENANCE OF HEALING OF DUODENAL ULCER
- I-198 USE OF LANSOPRAZOLE IN COMBINATION WITH AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OF A ONE-YEAR HISTORY OF A DUODENAL ULCER
- I-199 MONOTHERAPY AND COMBINATION THERAPY WITH SULFONYLUREAS IN THE TREATMENT OF TYPE II DIABETES
- I-200 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR
- I-201 EMPIRICAL THERAPY FOR FEBRILE NEUTROPENIC PATIENTS
- I-202 SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA
- I-203 MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
- I-204 USE IN PEDIATRIC PATIENTS BETWEEN THE AGES OF 6 AND 11 FOR THE TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- I-205 INITIAL ANTICONVULSANT TREATMENT OF STATUS EPILEPTICUS
- I-206 TREATMENT OF EDEMA ASSOCIATED WITH CHRONIC RENAL FAILURE
- I-207 FOR THE SUPPRESSION OF RECURRENT EPISODES OF GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
- I-208 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER IN THE PEDIATRIC POPULATION
- I-209 PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA (PSVT)

PATENT USE CODE

- U-161 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS IN A PATIENT
- U-162 METHOD OF USE TO INHIBIT CHOLESTEROL SYNTHESIS IN A HUMAN SUFFERING FROM HYPERCHOLESTEROLEMIA
- U-163 METHOD OF USING TROGLITAZONE TO TREAT IMPAIRED GLUCOSE TOLERANCE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS
- U-164 METHOD OF USING TROGLITAZONE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS IN A DEFINED POPULATION OF PATIENTS
- U-165 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA
- U-166 TREATMENT OF H. PYLORI ASSOCIATED DUODENAL ULCER
- U-167 METHOD FOR TREATING HIV-1 INFECTION
- U-168 METHOD OF INHIBITING LIPOXYGENASE ACTIVITY IN A MAMMAL WHICH IS THE MODE OF ACTION IN THE TREATMENT OF ASTHMA
- U-169 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS A CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING
- U-170 METHOD OF OBTAINING AN MR IMAGE USING THE COMPOSITION/DRUG PRODUCT AS A CONTRAST AGENT
- U-171 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS AN ORAL CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING OF THE GASTROINTESTINAL TRACT
- U-172 TREATMENT OF GENITAL WARTS
- U-173 ADMINISTRATION TO A HOST SUFFERING FROM GESTATIONAL DIABETES
- U-174 USE AS AN ANTIHISTAMINE AGENT
- U-175 METHOD OF TREATING MALIGNANT TUMORS
- U-176 METHOD OF TREATING A PATIENT SUFFERING FROM LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSIS
- U-177 FUNGICIDE

**PATENT AND EXCLUSIVITY TERMS
PATENT USE CODE**

- U-178 FACILITATED ADHERENCE OF AGENTS TO SKIN
- U-179 ENHANCED CUTANEOUS PENETRATION OF A DERMALLY-APPLIED PHARMACOLOGICALLY ACTIVE AGENT
- U-180 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER SIX MONTHS OF AGE) WITH ADVANCED HIV INFECTION
- U-181 PRODUCING ALPHA ADRENERGIC ANTAGONISTIC ACTION IN A HOST
- U-182 USE OF SALMETEROL IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
- U-183 TREATMENT OF CONDITIONS CAUSED BY DISTURBANCE OF NEURONAL 5HT FUNCTION
- U-184 TREATING ALLERGIC EYE DISEASES IN HUMANS
- U-185 METHOD OF TREATING HYPERTENSION
- U-186 METHOD FOR TREATING GI DISORDERS CAUSED BY H. PYLORI WHICH COMPRISES ADMINISTRATION OF RANITIDINE BISMUTH CITRATE AND CLARITHROMYCIN FOR A GREATER THAN ADDITIVE EFFECT
- U-187 THERAPEUTIC TREATMENT OF CALCIFIC TUMORS
- U-188 TREATMENT OF H. PYLORI ASSOCIATED DUODENAL ULCER
- U-189 ENHANCEMENT OF THE BIOAVAILABILITY OF THE DRUG SUBSTANCE
- U-190 USE OF RITONAVIR IN COMBINATION WITH ANY REVERSE TRANSCRIPTASE INHIBITOR
- U-191 METHOD OF TREATMENT FOR CONTROLLING AND LOWERING INTRAOCULAR PRESSURE IN A HUMAN
- U-192 USE IN TREATING ALLERGIC REACTIONS
- U-193 PSORIASIS
- U-194 TREATING AGINA PECTORIS AND HIGH BLOOD PRESSURE
- U-195 METHOD FOR THE DIAGNOSIS OF GASTROINTESTINAL DISORDERS BY UREA ISOTOAC OR NITROGEN LABELED CARBON
- U-196 TREATMENT OF METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH ESTROGEN RECEPTOR POSITIVE TUMORS
- U-197 USE IN COMBINATION WITH CERTAIN LHRH ANALOGUES FOR THE TREATMENT OF ADVANCED PROSTATE CANCER
- U-198 TREATMENT METASTATIC CARCINOMA OF OVARY AFTER FIRST-LINE FAILURE OR SUBSEQUENT CHEMOTHERAPY, TREATMENT OF BREAST CANCER AFTER FAILURE OF COMBINATION CHEMOTHERAPY FOR METASTATIC DISEASE AND SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA
- U-199 METHOD OF TREATING INFECTIOUS UPPER GI TRACT DISORDERS CAUSED BY CAMPYLOBACTER PYLORIS INFECTION COMPRISING ADMINISTRATION OF A BISMUTH AGENT AND AN ANTIMICROBIAL AGENT
- U-200 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF A BISMUTH-CONTAINING AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
- U-201 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF CAMPYLOBACTER-INHIBITING ANTIMICROBIAL AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
- U-202 METHOD OF TREATING PEPTIC ULCER DISEASE CAUSED BY CAMPYLOBACTER PYLORIS COMPRISING ORAL ADMINISTRATION OF 50 TO 5,000MG BISMUTH DAILY FOR 3-56 DAYS
- U-203 TREATMENT OF ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-204 USE OF TAXOL IN COMBINATION WITH G-CSF FOR TREATMENT OF PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-205 METHOD OF TREATING HEARTBURN
- U-206 METHOD OF USING FSH ALONE, WITHOUT THE PRESENCE OF EXOGENOUS LH, IN IN VITRO FERTILIZATION
- U-207 USE AS NASAL SPRAY
- U-208 VAGINAL ADMINISTRATION USING SPECIFIED FORMULATION
- U-209 VAGINAL ADMINISTRATION OF PROGESTERONE USING SPECIFIED FORMULATION

PATENT AND EXCLUSIVITY TERMS
PATENT USE CODE

U-210 METHOD OF TREATING CONGESTIVE HEART FAILURE
U-211 USE IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
U-212 METHOD OF TREATMENT OF PARKINSON'S DISEASE
U-213 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS AND TREATING HYPERCHOLESTEROLEMIA
 AND METHOD FOR TREATING HYPERLIPIDEMIA

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
>ADD>						
020059 001	ADENOSINE; ADENOSCAN	5070877	MAY 18, 2009	U-116	NCE	DEC 18, 2002
020760 001	ALATROFLOXACIN MESYLATE; TROYAN	5603918	JUN 09, 2015	NC	OCT 24, 1999	
020291 001	ALBUTEROL SULFATE; COMBIVENT	5225183	JUL 06, 2010	NP	AUG 15, 1999	
020503 001	ALBUTEROL SULFATE; PROVENTIL-HFA	5459670	JUL 06, 2010			
020560 001	ALENDRONATE SODIUM; FOSAMAX	5605674	FEB 25, 2014			
020560 002	ALENDRONATE SODIUM; FOSAMAX	4621077	AUG 06, 2007	U-114	I-185	APR 25, 2000
020560 003	ALENDRONATE SODIUM; FOSAMAX	5681590	DEC 02, 2012	I-187	APR 25, 2000	
		4621077	AUG 06, 2007	U-114	I-185	APR 25, 2000
		5681590	DEC 02, 2012	I-187	APR 25, 2000	
020649 001	ALPROSTADIL; EDEX	5424471	JUL 31, 2012	NP	JUL 06, 1998	
020649 002	ALPROSTADIL; EDEX	5591731	JUL 31, 2012	NP	JUL 06, 1998	
020649 003	ALPROSTADIL; EDEX			NP	JUL 06, 1998	
020649 004	ALPROSTADIL; EDEX			NP	JUL 06, 1998	
020221 001	AMIFOSTINE; ETHYOL			NP	JUL 06, 1998	
020333 001	ANAGRELIDE HYDROCHLORIDE; AGRYLIN			NP	JUL 06, 1998	
020333 002	ANAGRELIDE HYDROCHLORIDE; AGRYLIN			NP	JUL 06, 1998	
020561 001	ANASTROZOLE; ARIMIDEX	4935437	DEC 27, 2009	ODE	MAR 14, 2004	
020420 001	ARDEPARIN HYDROCHLORIDE; GENESA			NCE	MAR 14, 2002	
020227 002	ARDEPARIN SODIUM; NORNIFLO			NCE	MAR 14, 2004	
020702 001	ATORVASTATIN CALCIUM; LIPITOR			NCE	MAR 14, 2002	
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020702 002	ATORVASTATIN CALCIUM; LIPITOR	5686104	NOV 11, 2014	U-213		
		4681893	MAY 30, 2006	U-161		
		5273995	DEC 28, 2010	U-162		
		5385929	MAY 04, 2014	U-59		
		5686104	NOV 11, 2014	U-213	DEC 17, 2001	
		4681893	MAY 30, 2006	U-161		
		5273995	DEC 28, 2010	U-162		
		5385929	MAY 04, 2014	U-59		
		5686104	NOV 11, 2014	U-213	DEC 17, 2001	
		4681893	MAY 30, 2006	U-161		
		5273995	DEC 28, 2010	U-162		
		5385929	MAY 04, 2014	U-59		
		5164194	NOV 17, 2009	U-207		
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020114 001	AZELASTINE HYDROCHLORIDE; ASTELIN	4397839	JUL 01, 2005	NP	DEC 24, 1999	
020486 001	BECLOMETHASONE DIPROPIONATE; VANCERIL DOUBLE STRENGTH	5655172	JUN 03, 2014	NC	APR 17, 2000	
020032 001	BERACTANT; SURVANTA	4656505	OCT 01, 2008			
020619 001	BETAXOLOL HYDROCHLORIDE; BETOPTIC PILO	4472382	SEP 18, 2001	U-197		
020498 001	BICALUTAMIDE; CASODEX	5389813	SEP 18, 2001	U-197		
020490 001	BRIMONIDINE TARTRATE; ALPHAGAN			NP	MAR 13, 2000	
020335 002	BROMfenac Sodium; Duract			NCE	SEP 06, 2002	
				NCE	JUL 15, 2002	

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

APPL/PROD NUMBER		INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020441 002		BUDESONIDE;PULMICORT	4907583	MAR 13, 2007			
020441 003		BUDESONIDE;PULMICORT	4524769	JUN 25, 2002			
018644 001		BUPROPION HYDROCHLORIDE; WELLBUTRIN	4668218	APR 11, 2006			
018644 002		BUPROPION HYDROCHLORIDE; WELLBUTRIN	4907583	MAR 13, 2007			
018644 003		BUPROPION HYDROCHLORIDE; WELLBUTRIN	4524769	JUN 25, 2002			
020711 002		BUPROPION HYDROCHLORIDE; ZYBAN	4668218	APR 11, 2006			
020711 003		BUPROPION HYDROCHLORIDE; ZYBAN	5358970	AUG 12, 2013			
			5328970	AUG 12, 2013			
			5427798	AUG 12, 2013			
			RE33994	AUG 18, 2004			
			5358970	AUG 12, 2013			
			5427798	AUG 12, 2013			
			RE33994	AUG 18, 2004			
018731 003		BUSPIRONE HYDROCHLORIDE;BUSPAR	4215104	MAR 26, 1999			
018731 004		BUSPIRONE HYDROCHLORIDE;BUSPAR	4258027	MAR 26, 1999			
>ADD>	020524 001	BUTENATE HYDROCHLORIDE; MENTAX	4215104	MAR 26, 1999			
	019881 001	BUTOCONAZOLE NITRATE; FENSTAT ONE	4258027	MAR 26, 1999			
	020664 001	CABERGOLINE;DOSTINEX	4258027	MAR 26, 1999			
	020273 001	CALCIPOTRIENE;DOVONEX	4078071	MAR 07, 1997			
	020554 001	CALCIPOTRIENE;DOVONEX	45266892	JUL 02, 2002			
	020611 001	CALCIPOTRIENE;DOVONEX	48666048	DEC 29, 2007			
			48666048	SEP 12, 2006			
			4657927	APR 14, 2004	U-175		
			4657927	APR 14, 2004	U-175		
			4657927	APR 14, 2004	U-175		
			4657927	APR 14, 2004	U-175		
019880 001		CARBOPLATIN;PARAPLATIN	4503067	MAR 05, 2002	U-3		
019880 002		CARBOPLATIN;PARAPLATIN	5006530	JAN 17, 2009			
019880 003		CARBOPLATIN;PARAPLATIN	5177080	NOV 26, 2011			
020297 001		CARVEDILOL;COREG	5006530	JAN 17, 2009			
020297 002		CARVEDILOL;COREG	5177080	NOV 26, 2011			
020297 003		CARVEDILOL;COREG	5006530	JAN 17, 2009			
020297 004		CARVEDILOL;COREG	5177080	NOV 26, 2011			
020740 001		CERIVASTATIN SODIUM;BAYCOL	5006530	JAN 17, 2009			
020740 002		CERIVASTATIN SODIUM;BAYCOL	5006530	JAN 17, 2009			
020740 003		CERIVASTATIN SODIUM;BAYCOL	5177080	NOV 26, 2011			
020740 004		CERIVASTATIN SODIUM;BAYCOL	4525558	JUN 25, 2007			
			4525558	JUN 25, 2007			
			5177080	NOV 26, 2011			
			5006530	JAN 17, 2009			
			5177080	NOV 26, 2011			
			4525558	JUN 25, 2007			
			4525558	JUN 25, 2007			
019835 001		CETIRIZINE HYDROCHLORIDE;ZYRTEC	5286754	FEB 15, 2011			
019835 002		CETIRIZINE HYDROCHLORIDE;ZYRTEC	4670444	DEC 09, 2003	U-36		
020346 001		CETIRIZINE HYDROCHLORIDE;ZYRTEC			I-188		
020519 001		CICLOPIROX-LOPROX					
019537 001		CIPROFLOXACIN HYDROCHLORIDE;CIPRO					
019537 002		CIPROFLOXACIN HYDROCHLORIDE;CIPRO			I-188		

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019537 003 019537 004 019847 001	CIPROFLOXACIN HYDROCHLORIDE;CIPRO CIPROFLOXACIN HYDROCHLORIDE;CIPRO CIPROFLOXACIN;CIPRO	4705789	NOV 10, 2004		1-188 1-188 1-201	JUN 03, 2000 JUN 03, 2000 AUG 07, 2000
>ADD> >ADD> 020780 001 020780 002 019857 001	CIPROFLOXACIN;CIPRO CIPROFLOXACIN;CIPRO CIPROFLOXACIN;CIPRO IN DEXTROSE 5%	5695784 5695784 4808583	DEC 09, 2014 DEC 09, 2014 FEB 28, 2006		1-179 1-201	OCT 21, 1999 AUG 07, 2000
019858 001	CIPROFLOXACIN;CIPRO IN SODIUM CHLORIDE 0.9%	4705789	NOV 10, 2004		1-179 1-201	OCT 21, 1999 AUG 07, 2000
>ADD> 020767 001 017533 001 017533 002 017533 003 017533 005 017533 006 020813 001	CISAPRIDE MONOHYDRATE;PROPULSID QUICKSOLV CLONAZEPAM;KLONOPIN CLONAZEPAM;KLONOPIN CLONAZEPAM;KLONOPIN CLONAZEPAM;KLONOPIN CLONAZEPAM;KLONOPIN CLONAZEPAM;KLONOPIN	4962115	OCT 09, 2007	U-79	NCE	JUL 29, 1998 APR 09, 2000 APR 09, 2000 APR 09, 2000 APR 09, 2000 APR 09, 2000 APR 09, 2000
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>ADD> >ADD> >ADD> >ADD> >ADD> >ADD> 020813 003	CLONAZEPAM;KLONOPIN	5695777	OCT 03, 2010		NCE	NOV 17, 2002
>ADD> >ADD> >ADD> >ADD> >ADD> >ADD> 020813 004	CLONAZEPAM;KLONOPIN	5563142 548293 5500413 5674850 5498598 5500413 548293 5500413	OCT 08, 2013 JUN 29, 2013 JUN 29, 2013 DEC 23, 2013 JUN 29, 2013 DEC 23, 2013 JUN 29, 2013 DEC 23, 2013		NCE	JAN 03, 2000 APR 04, 2002
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>ADD> >ADD> >ADD> >ADD> >ADD> >ADD> 020839 001 020463 001 020430 001 020705 001 017922 001	CLOPIDOGREL BISULFATE;PLAVIX CROMOLYN SODIUM;NASALCROM DANAPAROID SODIUM ORGARAN DELAVIDINE MESYLATE;RESCRIPTOR DESMOPRESSIN ACETATE;DDAVP	5164377 5563142 548293 5500413 5674850 5498598 5500413 548293 5500413	OCT 03, 2010 OCT 08, 2013 JUN 29, 2013 JUN 29, 2013 DEC 23, 2013 JUN 29, 2013 DEC 23, 2013 JUN 29, 2013 DEC 23, 2013			
>DLT> 017922 002	DESMOPRESSIN ACETATE;DDAVP					
>ADD> >ADD> 017922 003	DESMOPRESSIN ACETATE;DDAVP					
>ADD> 018938 001 018938 002	DESMOPRESSIN ACETATE;DDAVP DESMOPRESSIN ACETATE;DDAVP					

PREScription AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019955 001	DESMOPRESSIN ACETATE;DDAVP	5500413 5674850 5500413 5674850 5462740	JUN 29, 2013 DEC 23, 2013 JUN 29, 2013 DEC 23, 2013 OCT 31, 2012		JUL 29, ND F OOE ND F OOE	2000 2004 2000 2004 2004
019955 002	DESMOPRESSIN ACETATE;DDAVP				JUL 29, ND F	2004
020648 001	DIAZEPAM;DIASSTAT				JUL 29, ND F	2004
020648 002	DIAZEPAM;DIASSTAT				JUL 29, ND F	2004
020648 003	DIAZEPAM;DIASSTAT				JUL 29, ND F	2004
020648 004	DIAZEPAM;DIASSTAT				JUL 29, ND F	2004
020648 005	DIAZEPAM;DIASSTAT				JUL 29, ND F	2004
020037 001	DICLOFENAC SODIUM;VOLTAREN				JUL 29, ND F	2004
020154 002	DIDANOSINE;VIDEX	4966799	OCT 03, 2007	00 E		
020154 003	DIDANOSINE;VIDEX	4820988	APR 14, 2007			
020154 004	DIDANOSINE;VIDEX	5616566	AUG 29, 2006			
020154 005	DIDANOSINE;VIDEX	5616566	AUG 29, 2006			
020155 003	DIDANOSINE;VIDEX	5616566	AUG 29, 2006			
020155 004	DIDANOSINE;VIDEX	5616566	AUG 29, 2006			
020155 005	DIDANOSINE;VIDEX	5616566	AUG 29, 2006			
020156 001	DIDANOSINE;VIDEX	5616566	AUG 29, 2006			
>ADD> 020148 001	DIHYDROERGOTAMINE MESYLATE;MIGRANAL					
018723 001	DIVALPROEX SODIUM;DEPAKOTE					
018723 002	DIVALPROEX SODIUM;DEPAKOTE					
018723 003	DIVALPROEX SODIUM;DEPAKOTE					
019680 001	DIVALPROEX SODIUM;DEPAKOTE					
020623 001	DOLASETRON MESYLATE MONOHYDRATE;ANZEMET					
020623 002	DOLASETRON MESYLATE MONOHYDRATE;ANZEMET					
020624 001	DOLASETRON MESYLATE MONOHYDRATE;ANZEMET					
020624 002	DOLASETRON MESYLATE MONOHYDRATE;ANZEMET					
>ADD> 020706 001	EMEDASTINE DIFUMARATE;EMADINE					
020668 001	ENALAPRIL MALEATE;LEXEL					
020164 001	ENOKAPARIN SODIUM;LOVENOX	4472380	SEP 18, 2001			
020444 001	EPOPROSTENOL SODIUM;FLOLAN	4374829 4703038	DEC 30, OCT 07,	U-3 2005		
020444 002	EPOPROSTENOL SODIUM;FLOLAN	4335139	JUN 15, 1999		1-192	MAY 06, 2000
>ADD> 020738 004	EPROSARTAN MESYLATE;TEVETEN	4539333	SEP 03,			
>ADD> 020738 005	EPROSARTAN MESYLATE;TEVETEN	4883812	MAY 12,	U-185		
020417 001	ESTRADIOL;FEMPATCH	4338325	SEP 03, JUL 06,	2002		
>ADD>		4335139	MAY 12, JUN 15,	1999	U-185	
>ADD>		4906463 5006542	MAR 06, APR 09,	2007 2008	NCE NP	DEC 22, DEC 22, DEC 03, 1999

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APL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020683 001	ETHINYL ESTRADIOL; ALESSE				NP	MAR 27, 2000
020683 002	ETHINYL ESTRADIOL; ORTHO TRI-CYCLEN	4544554	SEP 26, 2003	U-66	NP	MAR 27, 2000
019697 001	ETHINYL ESTRADIOL; ORTHO TRI-CYCLEN	4544554	SEP 26, 2003	U-66	I-177	DEC 31, 1999
019697 002	ETODOOLAC; IODINE	RE35524	AUG 04, 2007		I-177	DEC 31, 1999
018922 005	ETOPOSIDE PHOSPHATE; ETOPHOS				I-24	JUN 28, 1999
020457 001	FAMCICLOVIR; FAMVIR					
020363 001	FAMCICLOVIR; FAMVIR					
020363 002	FAMCICLOVIR; FAMVIR					
020363 003	FAMOTIDINE; PEPCID AC					
020325 001	FENOLODOPAM MESYLATE; CORLOPAM	4283408	OCT 15, 2000		I-207	SEP 16, 2000
019922 001	FERRIC AMMONIUM CITRATE; FERRIT SELTZ	5667794	MAY 02, 2015	U-205	I-207	SEP 16, 2000
020292 001	FERUMOXSIL; GASTROMARK					
020410 001						
020625 001	FEXOFENADINE HYDROCHLORIDE; ALLEGRA-D	5069216	MAY 09, 2006	U-171	NCE	SEP 23, 2002
>ADD> 020786 001	FLUDARABINE PHOSPHATE; FLUDARA	5219554	JUN 15, 2010		NP	OCT 14, 2000
>ADD> 020038 001	FLUTICASONE PROPIONATE; FLOVENT	4827945	MAY 09, 2006		NP	DEC 06, 2001
>ADD> 020549 001	FLUTICASONE PROPIONATE; FLOVENT	4951675	SEP 13, 2005		U-170	
>ADD> 020549 002	FLUTICASONE PROPIONATE; FLOVENT				U-169	
>ADD> 020549 003	FLUTICASONE PROPIONATE; FLOVENT					
>ADD> 020582 001	FOLLITROPIN ALFA/BETA; FOLLISTIM	5055288	OCT 08, 2008			
020582 002	FOLLITROPIN ALFA/BETA; FOLLISTIM	4695392	SEP 22, 2004			
020378 001	FOLLITROPIN ALFA/BETA; GONAL-F	4695393	SEP 22, 2004			
020378 002	FOMEPOZOLE; ANTIZOL	4770183	SEP 13, 2005		U-169	
>ADD> 020696 001	GABAPENTIN; NEURONTIN	5578610	NOV 26, 2013		U-192	
>ADD> 020255 001	GABAPENTIN; NEURONTIN					
020255 002	GABAPENTIN; NEURONTIN					
020255 003	GABAPENTIN; NEURONTIN					
020622 001	GLATIRMER ACETATE; COPAXONE	4357324	FEB 24, 2003			
020329 001	GLIPIZIDE; GLUCOTROL XL					
020329 002	GLIPIZIDE; GLUCOTROL XL					
020329 003	GOSERELIN ACETATE; ZOLADERX					
019726 001	GRANSETRON HYDROCHLORIDE; KYTRIL					
020305 001	GREPAFLOXACIN HYDROCHLORIDE; RAXAR					
>ADD> 020695 001	HYDROCHLOROTHIAZIDE; IRBESARTAN-HYDROCHLOROTHIAZID	5270317	MAR 20, 2011		D-36	OCT 06, 2002
020758 001	HYDROCHLOROTHIAZIDE; IRBESARTAN-HYDROCHLOROTHIAZID	5270317	MAR 20, 2011		NCE	NOV 06, 2002
020758 002	HYDROCHLOROTHIAZIDE; MICROZIDE				SEP 30, 2002	SEP 30, 2002
020504 001	HYDROCHLOROTHIAZIDE; UNIRETIC	4344949	OCT 03, 2000		NCE	DEC 27, 1999
020729 001	HYDROCHLOROTHIAZIDE; UNIRETIC	4743450	FEB 24, 2007		NS	JUN 27, 2000

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020729 002	HYDROCHLOROTHIAZIDE;UNI RETIC	4344949 4743450	OCT 03, 2000 FEB 24, 2007	NC	JUN 27, 2000	
020716 001	HYDROCODONE BITARTRATE;VICOPROFEN					
020267 002	IBUPROFEN JUNIOR STRENGTH ADVIL					
020723 001	IMIQUIMOD; ALDARA					
020808 001	IODIXANOL;VISIPAQUE 270					
020808 002	IODIXANOL;VISIPAQUE 320					
020757 001	IRBESARTAN;AVAPRO					
020757 002	IRBESARTAN;AVAPRO					
020757 003	IRBESARTAN;AVAPRO					
040024 001	IRON DEXTRAN;DEXFERRUM					
020083 001	ITRACONAZOLE;SPORANOX					
020657 001	ITRACONAZOLE;SPORANOX					
019927 001	KETOCONAZOLE;NIZORAL A-D					
020310 001	KETOCONAZOLE;NIZORAL A-D					
>ADD>	KETOROLAC TROMETHAMINE;ACULAR PRESERVATIVE FREE					
>ADD>						
>ADD>						
>ADD>						
020811 001	LAMIVUDINE;COMBIVIR					
020857 001	LANSOPRAZOLE;PREVACID					
020406 001	LANSOPRAZOLE;PREVACID					
020406 002	LANSOPRAZOLE;PREVACID					
020726 001	LETROZOLE;FEMARA					
019732 001	LEUPROLIDE ACETATE;LUPRON DEPOT					
020011 001	LEUPROLIDE ACETATE;LUPRON DEPOT					
020517 001	LEUPROLIDE ACETATE;LUPRON DEPOT					
		4978672	DEC 18, 2007	U-203		
		5631020	MAY 20, 2014			
		5631021	MAY 20, 2014			
		4954298	NOV 01, 2004			
		5480656	JAN 02, 2013			
		5643607	JUL 01, 2014			
		5631020	MAY 20, 2014			
		5631021	MAY 20, 2014			
		4954298	NOV 01, 2004			
		5480656	JAN 02, 2013			
		5643607	JUL 01, 2014			
		5631020	MAY 20, 2014			
		5631021	MAY 20, 2014			
		5643607	JUL 01, 2014			

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020263 002	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5480656 5631020 5631021	JAN 02, 2013 MAY 20, 2014 MAY 20, 2014			
020263 003	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	4954298 5643607 5643607	NOV 01, 2004 JUL 01, 2014 JUL 01, 2014			
020263 004	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5480656 5631020 5631020	JAN 02, 2013 MAY 20, 2014 MAY 20, 2014			
020263 005	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5631021 4954298 5643607	MAY 20, 2014 NOV 01, 2004 JUL 01, 2014			
020263 006	LEUPROLIDE ACETATE;LUPRON DEPOT-PED	5480656 5631020 5631021	JAN 02, 2013 MAY 20, 2014 MAY 20, 2014			
020708 001	LEUPROLIDE ACETATE;LUPRON DEPOT-3	4954298 5643607 4652411	NOV 01, 2004 JUL 01, 2014 NOV 01, 2004			
020517 002	LEUPROLIDE ACETATE;LUPRON DEPOT-4	4677191 4728721 4849228 4917893 4954298 5330767 5476663 5480656 5575987 5631020 5631021 4652441 5643607 4677191 4728721 4849228 5631020 5631021	JUL 03, 2005 MAY 01, 2006 JUL 18, 2006 NOV 01, 2004 NOV 01, 2004 NOV 01, 2004 APR 17, 2007 JAN 02, 2013 NOV 19, 2013 MAY 20, 2014 MAY 20, 2014 NOV 01, 2004 NOV 01, 2014 JUL 01, 2014 MAY 01, 2006 JUL 03, 2005 MAY 01, 2006 JUL 18, 2006 MAY 20, 2014 MAY 20, 2014 MAY 20, 2014 MAY 20, 2014 NOV 01, 2004 APR 17, 2007 JAN 02, 2013 NOV 19, 2013 NOV 01, 2004 NOV 01, 2004 JUL 01, 2014	NP	MAR 07, 2000	
					NS D-34	MAY 30, 2000 MAY 30, 2000

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020580 001	LIPASE;COTAZYM	5489436	FEB 06, 2013	NP	DEC 09, 1999	
020448 001	LOPERAMIDE HYDROCHLORIDE;IMODIUM A-D	5248505	JUL 28, 2010	NC	JUN 26, 2000	
020606 001	LOPERAMIDE HYDROCHLORIDE;IMODIUM ADVANCED	5612054	SEP 28, 2010			
020641 001	LORATADINE;CLARITIN	5489436	FEB 06, 2013			
020704 001	LORATADINE;CLARITIN REDITABS	5679376	OCT 21, 2014	U-142		
018140 001	LORAZEPAM;ATIVAN	4659716	APR 21, 2004	U-77	NC	APR 12, 1998
018140 002	LORAZEPAM;ATIVAN	428233	JUN 19, 2002	U-142		
020652 001	MANGAFODIPIR TRISODIUM;TESLASCAN	4659716	APR 21, 2004	U-77		
020264 001	MEGESTROL ACETATE;MEGACE	4833931	APR 21, 2004	U-142		
019651 001	MESALAMINE;ASACOL	4833931	SEP 15, 2008			
020357 001	METFORMIN HYDROCHLORIDE;GLUCOPHAGE	1-205	SEP 05, 2000			
020357 002	METFORMIN HYDROCHLORIDE;GLUCOPHAGE	1-205	SEP 05, 2000			
020868 001	METRONIDAZOLE;FLAGYL ER	NCE	MAR 03, 2000			
020743 001	METRONIDAZOLE;NITRATATE	NDF	MAR 03, 2000			
020689 001	MIBERFADIL DIHYDROCHLORIDE;POSICOR	NOV 26,	NOV 26, 2000			
020689 002	MIBERFADIL DIHYDROCHLORIDE;POSICOR	NP	SEP 26, 2000			
020762 001	MOMETASONE FURATE MONOHYDRATE;MASONEX	4808605	NOV 10, 2007	U-194	NCE	JUN 20, 2002
020353 001	NAPROXEN SODIUM;NAPRELAN	4808605	NOV 10, 2007	U-194	NDF	JUN 20, 2002
020353 002	NAPROXEN SODIUM;NAPRELAN	5637320	JUN 10, 2014			
020353 003	NAPROXEN SODIUM;NAPRELAN	5637320	JUN 10, 2014			
019660 001	NEDOCROMIL SODIUM;TILADE	5637320	JUN 10, 2014			
020750 001	NEDOCROMIL SODIUM;TILADE					
020778 001	NELFINAVIR MESYLATE;VIRACEPT	1-183	MAR 06, 2000			
020779 001	NELFINAVIR MESYLATE;VIRACEPT	NDF	OCT 01, 2000			
020636 001	NEVIRAPINE;VIRAMUNE	NCE	MAR 14, 2002			
020381 001	NIACIN;NIASPAN	MAR	MAR 14, 2002			
020381 002	NIACIN;NIASPAN	NCE	MAR 14, 2002			
020381 003	NIACIN;NIASPAN	JUN	JUN 21, 2001			
020381 004	NIACIN;NIASPAN	NDF	JUL 28, 2000			
020381 005	NIACIN;NIASPAN TITRATION STARTER PAC	NDF	JUL 28, 2000			
020714 001	NICOTINE;NICOTROL	NDF	JUL 28, 2000			
020592 001	OLANZAPINE;ZYPREXA	NP	MAY 02, 2000			
020592 002	OLANZAPINE;ZYPREXA	5605897	FEB 25, 2014	U-176		
020592 003	OLANZAPINE;ZYPREXA	5605897	FEB 25, 2014	U-176		
020592 004	OLANZAPINE;ZYPREXA	5605897	FEB 25, 2014	U-176		
020688 001	OLOPATADINE HYDROCHLORIDE;PATANOL	4871865	OCT 03, 2006			
019810 001	OMEPRAZOLE;PRILOSEC	5605897	FEB 25, 2014	U-174		
019810 003	OMEPRAZOLE;PRILOSEC	5116863	MAY 26, 2009			
		5605897	JUN 24, 2014	U-184		
		4636499	MAY 30, 2005			
		5093342	FEB 02, 2010			
		5599794	FEB 04, 2014	U-166		
		5629305	FEB 04, 2014	U-166		
		4636499	MAY 30, 2005	U-188		
		5093342	FEB 02, 2010	U-166		
		5599794	FEB 04, 2014	U-166		
		5629305	FEB 04, 2014	U-188		

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020605 001		ONDANSETRON HYDROCHLORIDE; ZOFRAN	4753789 4695578 5578628	JUN 24, 2006 JUN 25, 2005 JUN 24, 2006	U-44 U-183 U-44	I-200 0DE I-202	AUG 18, 2000 AUG 04, 2004 AUG 04, 2000
019828 001		OXICONAZOLE NITRATE; OXISTAT	5641803 5670537	AUG 03, 2012 AUG 03, 2012	U-198 U-198		
020262 001		PACLITAXEL; TAXOL	5496804 4721723 4721723 4721723 4721723 4721723 4721723 4721723	MAR 09, 2013 DEC 29, 2006 DEC 29, 2006 DEC 29, 2006 DEC 29, 2006 DEC 29, 2006 DEC 29, 2006 DEC 29, 2006	U-204 U-12 U-12 U-12 U-12 U-12 U-12 U-12	I-150 I-150 I-150 I-150 I-150 I-150 I-150 I-150	MAY 07, 1999 MAY 07, 1999
020031 001		PAROXETINE HYDROCHLORIDE; PAXIL	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
020031 002		PAROXETINE HYDROCHLORIDE; PAXIL	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
020031 003		PAROXETINE HYDROCHLORIDE; PAXIL	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
020031 004		PAROXETINE HYDROCHLORIDE; PAXIL	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
020031 005		PAROXETINE HYDROCHLORIDE; PAXIL	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
020710 001		PAROXETINE HYDROCHLORIDE; PAXIL	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
019385 001		PERGOLIDE MESYLATE; PERMAX	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
019385 002		PERGOLIDE MESYLATE; PERMAX	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
019385 003		PERGOLIDE MESYLATE; PERMAX	5114948 4180582	OCT 19, 2009 FEB 08, 1998	U-20		
018796 001		PILOCARPINE HYDROCHLORIDE; PILOPINE HS	4271143 4680399	MAY 09, 1999 JUL 14, 2004	NDF	MAR 13,	2000
020529 001		POOFILUX; CONDYLOX	5057616 4866812	OCT 15, 2008 DEC 12, 2006			
020667 001		PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4866812	DEC 12, 2006	NCE	JUL 01,	2002
020667 002		PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4866812	DEC 12, 2006	NCE	JUL 01,	2002
020667 003		PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4866812	DEC 12, 2006	NCE	JUL 01,	2002
020667 004		PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4866812	DEC 12, 2006	NCE	JUL 01,	2002
020667 005		PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4866812	DEC 12, 2006	NCE	JUL 01,	2002
019568 001		PREDNICKARBATE; DERMATOP	4242334	AUG 02, 2000	U-50		
020279 001		PREDNICKARBATE; DERMATOP	4242334	AUG 02, 2000	U-50		
020545 001		PROCANAMIDE HYDROCHLORIDE; PROCANBD	565696	AUG 12, 2014			
020545 002		PROCANAMIDE HYDROCHLORIDE; PROCANBD	5656296	AUG 12, 2014			
020701 001		PROGESTERONE; CRINONE	4615697	OCT 07, 2003	NP	JUL 31,	2000
020701 002		PROGESTERONE; CRINONE	5543150	SEP 15, 2013	NP	JUL 31,	2000
>ADD>	019151 001	PROGESTERONE; CRINONE	4615697	OCT 07, 2003	NP	JUL 31,	2000
>ADD>	019151 002	PROPafenone HYDROCHLORIDE; RYTHMOL	5543150	SEP 15, 2013	NP	JUL 31,	2000
>ADD>	019151 003	PROPafenone HYDROCHLORIDE; RYTHMOL	4879288	MAR 20, 2007			
	019627 002	PROPOFOL; DIPRIVAN	4879288	MAR 20, 2007			
	020639 001	QUETiapine FUMARATE; SERQUEL	4879288	MAR 20, 2007			
	020639 002	QUETiapine FUMARATE; SERQUEL	4879288	MAR 20, 2007			
	020639 003	QUINAPRIL HYDROCHLORIDE; ACCUPRIL	5684016	NOV 04, 2014	U-210		
>ADD>	019885 001	QUINAPRIL HYDROCHLORIDE; ACCUPRIL	5684016	NOV 04, 2014	U-210		
>ADD>	019885 002						

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
>ADD> 019885 003 QUINAPRIL HYDROCHLORIDE;ACCUPRIL >ADD> 019885 004 QUINAPRIL HYDROCHLORIDE;ACCUPRIL >ADD> 020815 001 RALOXIFENE HYDROCHLORIDE;EVISTA >ADD> 020559 001 RANITIDINE BISMUTH CITRATE;TRITEC		5684016 5684016	NOV 04, 2014 NOV 04, 2014	U-210 U-210	NCE	DEC 09, 2002
>ADD> 020741 001 REPAGLINIDE;PRANDIN >ADD> 020741 002 REPAGLINIDE;PRANDIN >ADD> 020741 003 REPAGLINIDE;PRANDIN >ADD> 020474 001 RIMEXOLONE;VEKOL 020272 001 RISPERIDONE;RISPERDAL 020272 002 RISPERIDONE;RISPERDAL 020272 003 RISPERIDONE;RISPERDAL 020272 004 RISPERIDONE;RISPERDAL 020588 001 RISPERIDONE;RISPERDAL		46886214 5158952 5158952 5158952 5158952 5158952 5158952 5158952	JUL 23, 2008 OCT 27, 2009 OCT 27, 2009 OCT 27, 2009 OCT 27, 2009 OCT 27, 2009 JUN 11, 2014	U-100 U-90 U-90 U-90 U-90 U-90 U-100	NCE NCE NCE NCE NCE NCE NCE	DEC 22, 2002 DEC 22, 2002 DEC 22, 2002
RITONAVIR;NORVIR		5616587 5635523 5635523 5635523 5635523 5648497 5648497	JUN 11, 2014 JUN 03, 2014 JUN 03, 2014 JUN 03, 2014 JUN 03, 2014 JUL 15, 2014 JUL 15, 2014	U-190 U-190 U-190 U-190 U-190 U-190		
RITONAVIR;NORVIR		4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 5380922	DEC 07, 2002 MAY 19, 2008 DEC 07, 2002 MAY 19, 2008 JAN 10, 2012	U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212	NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE	SEP 19, 2002 SEP 19, 2002
ROPINIROLE HYDROCHLORIDE;REQUIP		4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 4452808 4824860 5380922	DEC 07, 2002 MAY 19, 2008 DEC 07, 2002 MAY 19, 2008 JAN 10, 2012	U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212 U-212	NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE NCE	SEP 19, 2002 SEP 19, 2002
SALMETEROL XINAFOATE;SEREVENT		5225645 4992474 5225645 5380922 4898724	FEB 12, 2008 FEB 12, 2008 FEB 19, 2012 MAY 14, 2013 FEB 06, 2007	U-182 U-182 U-211 U-187 U-187		
SALMETEROL XINAFOATE;SEREVENT		4898724 4898724	FEB 06, 2007			
SAMARIUM SM 153 LEXIDRONAM PENTASODIUM;QUADRAMET SERMORELIN ACETATE;GEREF		5380922 4898724	MAR 28, 2002 MAR 28, 2002 NP SEP 26, 2004 NP SEP 26, 2000 NP SEP 26, 2004 NP SEP 26, 2000 NP SEP 26, 2000 NP SEP 26, 2000 NP SEP 26, 2000			
SERTRALINE HYDROCHLORIDE;ZOLOFT		019839 001	I-193			
SERTRALINE HYDROCHLORIDE;ZOLOFT		019839 002	I-208	OCT 10,	2000	
SERTRALINE HYDROCHLORIDE;ZOLOFT		019839 003	I-193	JUL 08,	2000	
			I-208	OCT 10,	2000	
			I-193	JUL 08,	2000	
			I-208	OCT 10,	2000	
			I-193	JUL 08,	2000	
			I-208	OCT 10,	2000	

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019839 004	SERTRALINE HYDROCHLORIDE; ZOLOFT				1-193	JUL 08, 2000
019839 005	SERTRALINE HYDROCHLORIDE; ZOLOFT				1-208	OCT 10, 2000
>ADD>	SIBUTRAMINE HYDROCHLORIDE; MERIDIA	47466680	JUN 11, 2002	NCE	1-193	JUL 08, 2000
>ADD>	SIBUTRAMINE HYDROCHLORIDE; MERIDIA	4929629	MAY 29, 2007	NCE	1-208	OCT 10, 2000
>ADD>	SIBUTRAMINE HYDROCHLORIDE; MERIDIA	47466680	JUN 11, 2002	NCE	NOV 22, 2002	NOV 22, 2002
>ADD>	SIBUTRAMINE HYDROCHLORIDE; MERIDIA	4929629	MAY 29, 2007	NCE	NOV 22, 2002	NOV 22, 2002
>ADD>	SIBUTRAMINE HYDROCHLORIDE; MERIDIA	47466680	JUN 11, 2002	NCE	NOV 22, 2002	NOV 22, 2002
020231 001	SODIUM FLUORIDE; COLGATE TOTAL SOMATROPIN, BIOSYNTHETIC; GENOTROPIN SOMATROPIN, BIOSYNTHETIC; GENOTROPIN PRESERVATIVE FREE	4929629	MAY 29, 2007	NCE	JUL 11, 2000	JUL 08, 2000
020280 006	SOMATROPIN, BIOSYNTHETIC; GENOTROPIN SOMATROPIN, BIOSYNTHETIC; HUMATROPE	4816470	DEC 28, 2006	ODE	OCT 31, 2004	OCT 31, 2004
020280 004	SOMATROPIN, BIOSYNTHETIC; HUMATROPE	4816470	DEC 28, 2006	ODE	OCT 31, 2004	OCT 31, 2004
019640 001	SOMATROPIN, BIOSYNTHETIC; HUMATROPE	4816470	DEC 28, 2006	ODE	DEC 30, 1999	DEC 30, 1999
019640 004	SOMATROPIN, BIOSYNTHETIC; HUMATROPE	4816470	DEC 28, 2006	ODE	DEC 30, 2003	DEC 30, 2003
020168 001	SOMATROPIN, BIOSYNTHETIC; NUTROPIN SOMATROPIN, BIOSYNTHETIC; NUTROPIN	4816470	DEC 28, 2006	ODE	DEC 30, 1999	DEC 30, 1999
020168 002	SOMATROPIN, BIOSYNTHETIC; NUTROPIN	4816470	DEC 28, 2006	ODE	DEC 30, 2003	DEC 30, 2003
020604 003	SUMATRIPTAN, BIOSYNTHETIC; SEROSTIM SUMATRIPTAN, IMITREX SUMATRIPTAN, IMITREX	4816470	DEC 28, 2006	U-72	NP	AUG 23, 1999
020626 001	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	NDF	AUG 26, 2000	AUG 26, 2000
020626 002	TACRINE HYDROCHLORIDE; COGNEX	4816456	DEC 01, 2006	NDF	AUG 26, 2000	AUG 26, 2000
020626 003	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	U-82	U-82	U-82
>ADD>	TACRINE HYDROCHLORIDE; COGNEX	4816456	DEC 01, 2006	U-82	U-82	U-82
>DLT>	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	U-82	U-82	U-82
>ADD>	TACRINE HYDROCHLORIDE; COGNEX	4816456	DEC 01, 2006	U-82	U-82	U-82
>DLT>	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	U-82	U-82	U-82
>ADD>	TACRINE HYDROCHLORIDE; COGNEX	4816456	DEC 01, 2006	U-82	U-82	U-82
>DLT>	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	U-82	U-82	U-82
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>DLT>	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	U-82	U-82	U-82
>ADD>	TACRINE HYDROCHLORIDE; COGNEX	4816456	DEC 01, 2006	U-82	U-82	U-82
>ADD>	TACRINE HYDROCHLORIDE; COGNEX	4816456	SEP 09, 2007	U-82	U-82	U-82
>ADD>	TACRINE HYDROCHLORIDE; COGNEX	4816456	DEC 01, 2006	U-82	U-82	U-82
020579 001	TAMSOLOIN HYDROCHLORIDE; FLOMAX	4868216	SEP 19, 2006	U-181	NCE	DEC 24, 2002
		4731478	OCT 27, 2004			APR 15, 2002
		4703063	OCT 27, 2004			
		4772475	FEB 27, 2006			
020600 001	TAZAROTENE; TAZORAC	5089509	FEB 18, 2009	U-193	NCE	JUN 13, 2002
020600 002	TAZAROTENE; TAZORAC	5089509	FEB 18, 2009	U-193	NCE	JUN 13, 2002
019785 003	TECHNETIUM TC-99M SESTAMBI KIT; MIRALUMA	5045302	AUG 14, 2008			MAY 23, 2000
020372 001	TECHNETIUM TC-99M TROFOSMIN KIT; MYOVIEW	5504207	APR 29, 2013			
019057 001	TERAZOSIN HYDROCHLORIDE; HYTRIN	5294615	APR 29, 2013	U-3		
019057 002	TERAZOSIN HYDROCHLORIDE; HYTRIN	5504207	APR 29, 2013	U-165		
019057 003	TERAZOSIN HYDROCHLORIDE; HYTRIN	5294615	APR 29, 2013	U-3		

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT EXPRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019057 004	TERAZOSIN HYDROCHLORIDE; HYTRIN	5504207 5294615	APR 29, 2013 APR 29, 2013	U-165		
020347 001	TERAZOSIN HYDROCHLORIDE; HYTRIN	5294615	APR 29, 2013	U-3		
020347 002	TERAZOSIN HYDROCHLORIDE; HYTRIN	5294615	APR 29, 2013	U-3		
020347 003	TERAZOSIN HYDROCHLORIDE; HYTRIN	5294615	APR 29, 2013	U-3		
020347 004	TERAZOSIN HYDROCHLORIDE; HYTRIN	5294615	APR 29, 2013	U-3		
020192 001	TERBINAINE HYDROCHLORIDE; LAMISIL	4755534 4755534	DEC 30, 2006 DEC 30, 2006	U-73		
020539 001	TERBINAINE HYDROCHLORIDE; LAMISIL	4755534	DEC 30, 2006	NDF	OCT 17, 2000	
020749 001	TERBINAINE HYDROCHLORIDE; LAMISIL	5010090	OCT 07, 2008	NCE	SEP 30, 2002	
020646 001	TIAGABINE HYDROCHLORIDE; GABITRIL	5354760	MAR 24, 2012			
020646 002	TIAGABINE HYDROCHLORIDE; GABITRIL	5010090	OCT 07, 2008	NCE	SEP 30, 2002	
020646 003	TIAGABINE HYDROCHLORIDE; GABITRIL	5354760	MAR 24, 2012			
020646 004	TIAGABINE HYDROCHLORIDE; GABITRIL	5010090	OCT 07, 2008	NCE	SEP 30, 2002	
020707 001	TILDURONATE DISODIUM; SKELID	5354760	MAR 24, 2012			
020676 001	TILOCONAZOLE; VAGISTAT - 1	4872488	OCT 24, 2006	NCE	MAR 07, 2002	
050753 001	TOBRAMYCIN; TOBI	4980171	APR 06, 2009			
020497 001	TOREMIFENE CITRATE; FARESTON	4696949	SEP 29, 2004	U-196		
>ADD>	TORSEMIDE; DEMADEX	4971800 5043317 4690825	NOV 20, 2007 SEP 03, 2008 OCT 04, 2005	U-178 U-179 U-134	NP I-206 I-206	FEB 11, 2000 DEC 22, 2004 MAY 29, 2002 MAY 29, 2004 I-206 I-206 I-206 I-206 I-206
	TORSEMIDE; DEMADEX				SEP 09,	SEP 09, 2000
	TORSEMIDE; DEMADEX				SEP 09,	SEP 09, 2000
	TORSEMIDE; DEMADEX				SEP 09,	SEP 09, 2000
	TORSEMIDE; DEMADEX				SEP 09,	SEP 09, 2000
	TRETINOIN; AVITA				SEP 09,	SEP 09, 2000
	TRETINOIN; RETIN-A MICRO				I-206	
	TRIAMCINOLONE ACETONIDE; NASACORT AQ				I-206	
	TRIMETHOPRIM HYDROCHLORIDE; PRIMSOLO				I-206	
	TRIMETREXATE GLUCURONATE; NEUTREXIN				I-206	
	TROGLITAZONE; PRELAY				I-206	
020719 002	TROGLITAZONE; PRELAY	5104888 5478852 5457109 5602133	AUG 28, 2004 SEP 15, 2013 SEP 15, 2013 SEP 15, 2013	NCE U-163 U-164 U-173	I-189 I-199	JUN 17, 2000 AUG 04, 2000 JAN 29, 2002 JAN 29, 2002
		5104888 5478852 5457109 5602133	AUG 28, 2004 SEP 15, 2013 SEP 15, 2013 SEP 15, 2013	NCE U-163 U-164 U-173	I-199	AUG 04, 2000

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