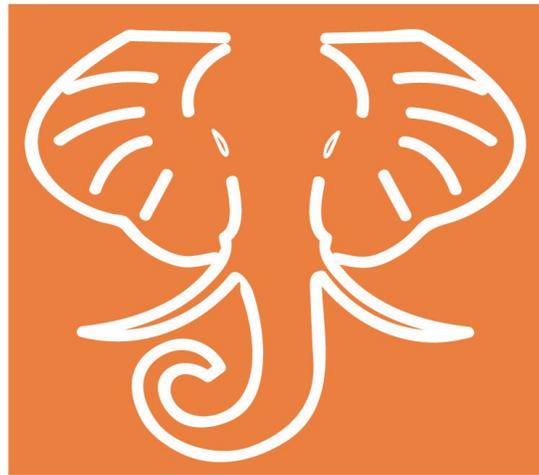


Approved prescription drug products with therapeutic equivalence evaluations.

[Washington, D.C.?] : U.S. Dept. of Health and Human Services, Public Health Service, Food and Drug Administration, Bureau of Drugs : 1980-

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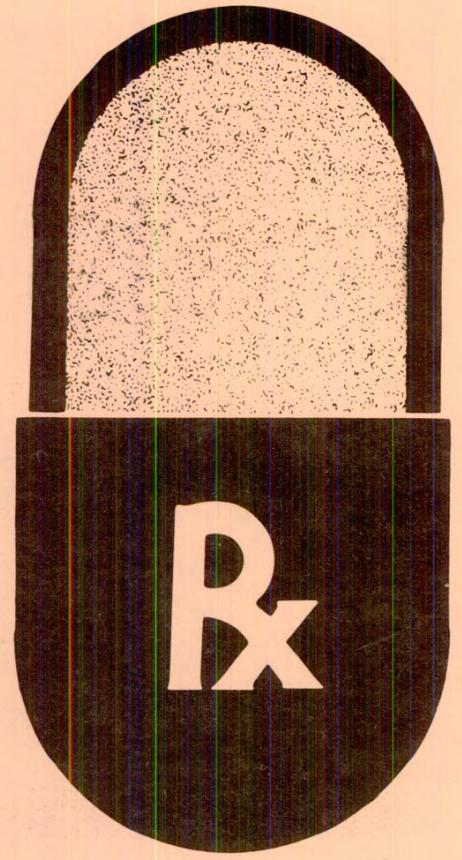
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APPROVED PRESCRIPTION DRUG PRODUCTS

SERIAL

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

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FOOD AND DRUG ADMINISTRATION
APPROVED PRESCRIPTION DRUG PRODUCTS
WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS
CUMULATIVE SUPPLEMENT

I. PREFACE

This cumulative supplement is one of a series of monthly updates to the Approved Prescription Drug Products with Therapeutic Equivalence Evaluations, 4th Edition, (the List) to cover interim revisions to the annual publication of the List in its entirety. The List is comprised of several parts and some by their nature are identified by the term "List". The cumulative supplements routinely provide updates to two of these lists: The Drug Product List and the DESI Addendum.

The List cannot be used effectively without the current cumulative supplement. Users may wish to place an asterisk (*) in the List to the left of the ingredient(s) in the Drug Product List and the product name in the Addendum to indicate that changes to that entry appear in the cumulative supplement. It is also suggested that earlier cumulative supplements be discarded to avoid possible confusion. In this way, only the List and current cumulative supplement need be referenced.

A. DRUG PRODUCT LIST

The Drug Product List cumulative supplements include the changes made since August 1, 1983. Each subsequent cumulative supplement replaces the previous month's cumulative supplement.

Information in this cumulative supplement follows the format of the Drug Product List. The presence of any therapeutic equivalence code indicates that the drug product is multिसource; 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.)

Context information on drug products is provided in each cumulative supplement for completeness to assist in locating the proper place in the Drug Product List for the revision. (Strength(s) which already exist in the publication will not be repeated for context.) A page number in parentheses referring to the Drug Product List is located to the right of the ingredient(s).

Additions to the Drug Product List are indicated by new information in the cumulative supplement. Additions new to the current cumulative supplement are indicated by the symbol >ADD> to the left of the line on which new information exists. The >ADD> symbol is dropped in subsequent cumulative supplements for that item.

Deletions from the Drug Product List are indicated by overstruck print in the cumulative supplement. Deletions new to the current cumulative supplement are indicated by the symbol >DLI> (DELETE) to the left of the line containing the overstruck print. The >DLI> symbol is dropped in subsequent cumulative supplements for that item.

A newly approved product is identified by the lozenge (◆) to the right of its strength. This identifier remains throughout all cumulative supplements for this edition.

B. ADDENDUM: DESI Pending List

Information in this cumulative supplement follows the format of the Addendum. Additions and deletions are indicated in the same manner as in the cumulative supplement to the Drug Product List. A change in Current Status of a DESI product is also indicated by an addition and a deletion.

II. SPECIAL NOTES

A. REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

Categories of counts derived from product information in the Drug Product List and from this cumulative supplement are presented. The report includes counts of new molecular entities approved by the agency during the current month.

B. PRODUCTS CONTAINING PHENACETIN

The October 5, 1983, Federal Register (48FR45466) provides the following Summary: "The Food and Drug Administration (FDA) is withdrawing approval of new drug applications or parts of new drug applications that provide for drug products containing phenacetin, except for those drug products that are the subject of a hearing request. The basis of the withdrawal is phenacetin's high potential for misuse and its unfavorable benefit-to-risk ratio when incorporated in analgesic combinations which are then subject to excessive chronic use." The effective date of this withdrawal order is November 4, 1983.

Because the subject products are no longer approved, the cumulative supplement has identified them by deleting the applicable active ingredient headers followed by a reference to this Special Note.

C. THEOPHYLLINE CONTROLLED RELEASE

Two controlled released theophylline tablets are listed as therapeutically equivalent (AB). Because one of these products was recently approved for once-daily dosing, it is important to be aware that the therapeutic equivalence rating was made on the basis of 12-hour dosing intervals. The rating does not apply to once-daily dosing. To date, no data have been submitted upon which the Agency can base a therapeutic equivalence determination among any of the approved theophylline controlled-release products when dosed once-daily.

D. APPLICANT (NAME) CHANGES

Because 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, the cumulation of these transfers and name changes will be identified in this Special Notes section only. Where only partial approved product lines are transferred between applicants, each approved product involved will appear as an applicant name change in the cumulative supplement. The current list of applicant holder changes follows.

APPLICANT (NAME) CHANGES

<u>Former Applicant (Name)</u>	<u>New Applicant (Name)</u>	<u>New Abbreviated Name</u>
STERI-MED INC SUB KETCHUM LABORATORIES INC	QUANTUM PHARMICS LTD	QUANTUM PHARMICS
KETCHUM LABORATORIES INC	QUANTUM PHARMICS LTD	QUANTUM PHARMICS
PROFESSIONAL PHARMACAL DIV STERI-MED INC	OPTOPICS LABORATORIES CORP	OPTOPICS LABS

The reader should consult the above cumulative list each month to become aware of such transfers and changes. By referring to the Applicant Index in the 4th Edition of the APPDP, the transferred products can be identified. The reader might wish to flag these products in the List as a reminder that the applicant has been changed.

III. REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

DESCRIPTION OF REPORT

The following report provides summary counts derived from product information in the Drug Product List and the current cumulative supplement. The counts appear in two sections. Section A. refers to the products in the List and Section B. to products in the current cumulative supplement. A new column of data will appear in Section A. each three-month period following July '83. Section A. therefore will provide baseline and quarterly data while Section B. provides monthly activity.

USE OF REPORT

From the data presented under Section B., users should be able to observe such things as (1) newly approved, DESI effective and remarketed drug products which are added to the List; (2) products that are being removed from the List as the result of withdrawal of approval, changes from prescription to over-the-counter status and discontinued marketing of products; and, (3) trends in approval of products as either multi-source or single source during each month within the quarter. The report does not reflect category changes from multi-source to single source and vice versa. However, the net gain that results from all additions, deletions and category changes is reflected in the quarterly counts for multi-source and single source products.

Drug Product Definition

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

New Molecular Entity

The active moiety 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 part of a combination.

Drug Product Count

This report provides counts in several categories from the list composed of domestically marketed drug products approved for both safety and effectiveness under sections 505 and 507 of the Federal Food, Drug, and Cosmetic Act. Counts of products still pending in the DESI review are not provided. Excluded also are those approved drug products marketed by distributors; those marketed solely abroad; and products now regarded as medical devices, biologics or foods. Also not included in the counts are those duplicate products of a given applicant whose only distinguishing characteristics are items such as package size, inactive ingredient(s), color and alternate manufacturing sites. These various counts are excluded because the Drug Product List itself excludes products from these categories.

REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

A. COUNTS CUMULATIVE BY QUARTERS

<u>CATEGORIES COUNTED</u>	<u>JULY '83 (BASELINE)</u>	<u>OCT '83</u>
DRUG PRODUCTS LISTED	6679	6783
SINGLE SOURCE	1908 (28.6%)	1915 (28.2%)
MULTISOURCE ⁽¹⁾	4771 (71.4%)	4868 (71.8%)
THERAPEUTICALLY EQUIVALENT	3804 (57.0%)	3891 (57.4%)
NOT THERAPEUTICALLY EQUIVALENT	957 (14.3%)	967 (14.3%)
EXCEPTIONS ⁽²⁾	10 (0.1%)	10 (0.1%)
NEW MOLECULAR ENTITIES APPROVED	-	2
NUMBER OF APPLICANTS	304	310

B. ACTIVITY FOR SUPPLEMENT NUMBER 4

	<u>NOV '83</u>	<u>DEC '83</u>	<u>CUMULATIVE</u>
DRUG PRODUCTS ADDED:	43	39	82
NEWLY APPROVED	43	39	82
DESI EFFECTIVE	0	0	0
REMARKETED	0	0	0
DRUG PRODUCTS REMOVED:	36	6	42
WITHDRAWN APPROVAL	27	0	27
RX TO OTC SWITCH	0	0	0
DISCONTINUED MARKETING	9	6	15
NET GAIN IN DRUG PRODUCTS	7	33	40
SINGLE SOURCE PRODUCTS APPROVED	10	9	19
MULTISOURCE DRUG PRODUCTS APPROVED	33	30	63
NEW MOLECULAR ENTITIES APPROVED:	3	2	5
AS THE ENTITY	3	1	4
AS A SALT, ESTER OR DERIVATIVE OF THE ENTITY	0	1	1

(1) THERAPEUTIC EQUIVALENCE EVALUATIONS PROVIDED ONLY FOR MULTISOURCE PRODUCTS (I.E., AVAILABLE FROM MORE THAN ONE APPLICANT)

(2) AMINO ACID-CONTAINING PRODUCTS OF VARYING COMPOSITION (SEE PAGE 1-5 OF THE LIST)

APPROVED PRESCRIPTION DRUG PRODUCTS
 DRUG PRODUCT LIST
 CUMULATIVE SUPPLEMENT NUMBER 4 / AUGUST '83 - DECEMBER '83

ACETAMINOPHEN; CODEINE PHOSPHATE (PAGE 3-1)

CAPSULE; ORAL
 > ADD > AA ACETAMINOPHEN W/ CODEINE #3
 > ADD > AA LEMMON 300MG;30MG*
TYLENOL W/ CODEINE NO. 3
 > ADD > AA MCNEIL PHARM 300MG;30MG

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE (PAGE 3-2)

TABLET; ORAL
 > ADD > AA CODACET
 > ADD > AA HALSEY DRUG 325MG;5MG*
/PERCOCET-S/
PERCOCET

/ACETAMINOPHEN; PHENACETIN; PHENYLPROPANOLAMINE HYDROCHLORIDE;/
/PHENYLTOLOXAMINE CITRATE/ (PAGE 3-2)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ALBUMIN, IODINATED, I-131, SERUM (PAGE 3-4)

INJECTABLE; INJECTION
/RADIOIODINATED SERUM ALBUMIN (HUMAN) I-131/
/MALLINCKRODT/ 16.7-250 UCI/ML/

AMINOPHYLLINE (PAGE 3-8)

/LIQUID; ORAL/
 SOLUTION; ORAL
AMINOPHYLLINE
 > ADD > AA BAY LABORATORIES 105MG/5ML*
AA ROXANE LABORATORIES 105MG/5ML*
 TABLET; ORAL
AMINOPHYLLINE
 BD BARR LABORATORIES 100MG*
 BD 200MG*
 AB VANGARD LABS/MM 100MG*
 AB 200MG*

/AMPHETAMINE SULFATE/ (PAGE 3-13)

/CAPSULE; CONTROLLED RELEASE; ORAL/
/BENZEDRINE/
/SKF LABORATORIES/ 15MG/

/AMPHETAMINE SULFATE/ (PAGE 3-13)

/TABLET; ORAL/
/BENZEDRINE/
/SKF LABORATORIES/ 5MG/
10MG/

AMPICILLIN TRIHYDRATE; PROBENECID (PAGE 3-13)

POWDER FOR RECONSTITUTION; ORAL
POLYGILLIN-PRB
 AB BRISTOL LABS/B-M EQ 3.5GM BASE/BOT;1GM/BOT
PROBAMPACIN
 AB BIOCRRAFT LABS EQ 3.5GM BASE/BOT;1GM/BOT

ASPIRIN; BUTALBITAL (PAGE 3-15)

TABLET; ORAL
 AXOTAL
 ADRIA LABORATORIES 650MG;50MG*

ASPIRIN; BUTALBITAL; CAFFEINE (PAGE 3-15)

TABLET; ORAL
 ASPIRIN AND CAFFEINE W/ BUTALBITAL
 PUREPAC/KALIPHARMA 325MG;50MG;40MG*

/ASPIRIN; BUTALBITAL; CAFFEINE; PHENACETIN/ (PAGE 3-15)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE (PAGE 3-15)

CAPSULE; ORAL
 SYNALGOS-DC
 IVES LABS/AMHO 356.4MG;30MG;16MG*

/ASPIRIN; CAFFEINE; PHENACETIN; PROPOXYPHENE HYDROCHLORIDE/
 (PAGE 3-15)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE (PAGE 3-16)

CAPSULE; ORAL
DARVON COMPOUND-65
 AA ELI LILLY INDSTRS/PR 389MG;32.4MG;65MG

BUTABARBITAL SODIUM (PAGE 3-24)

ELIXIR; ORAL
BUTABARBITAL SODIUM
 /AA/ /WEST-HARD/ /33MG/5ML/
 BUTALAN
 /AA/ LANNETT 33.3MG/5ML

/CAFFEINE; CARISOPRODOL; CODEINE PHOSPHATE; PHENACETIN/
 (PAGE 3-24)

/TABLET; ORAL/
 /SOMA COMPOUND W/ CODEINE/
 /WALLACE PHARMS/C-W/ /32MG;200MG;16MG;160MG/

/CAFFEINE; CARISOPRODOL; PHENACETIN/ (PAGE 3-25)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

CAFFEINE; ERGOTAMINE TARTRATE (PAGE 3-25)

SUPPOSITORY; RECTAL
 CAFERGOT
 BR SANDOZ PHARMS/SANDOZ 100MG;2MG
 WIGRAINE
 BR ORGANON/AKZONA 100MG;2MGx

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE (PAGE 3-26)

SOLUTION; INTRAPERITONEAL
 DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER
 AM MCGAW/AM HOSP 510MG/100ML;30GM/100ML;
 200MG/100ML;9.2GM/100ML;
 9.6GM/100MLx
 510MG/100ML;30GM/100ML;
 200MG/100ML;9.4GM/100ML;
 11GM/100MLx
 DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER
 AM MCGAW/AM HOSP 510MG/100ML;50GM/100ML
 200MG/100ML;9.2GM/100ML;
 9.6GM/100MLx
 510MG/100ML;50GM/100ML;
 200MG/100ML;9.4GM/100ML;
 11GM/100MLx

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE (PAGE 3-26)

SOLUTION; INTRAPERITONEAL
 DIALYTE W/ DEXTROSE 2.5% IN PLASTIC CONTAINER
 AM MCGAW/AM HOSP 26MG/100ML;2.5GM/100ML;
 15MG/100ML;560MG/100ML;390MG/100MLx

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE (PAGE 3-27)

INJECTABLE; INJECTION
 ISOLYTE E IN PLASTIC CONTAINER
 AM MCGAW/AM HOSP 35MG/100ML;30MG/100ML;
 74MG/100ML;640MG/100ML;
 500MG/100ML;74MG/100MLx

CEFTIZOXIME SODIUM (PAGE 3-30)

INJECTABLE; INJECTION
 CEFIZOX
 SK&F LABORATORIES EQ 1GM BASE/VIALx
 EQ 2GM BASE/VIALx

CEFUROXIME SODIUM (PAGE 3-30)

INJECTABLE; INJECTION
 ZINACEF
 GLAXO EQ 750MG BASE/VIALx
 EQ 1.5GM BASE/VIALx

CEPHALOTHIN SODIUM (PAGE 3-31)

INJECTABLE; INJECTION
KEFLIN
 AP ELI LILLY EQ 1GM BASE/VIAL
 AP EQ 2GM BASE/VIAL
SEFFIN
 AP GLAXO EQ 1GM BASE/VIALx
 AP EQ 2GM BASE/VIALx
 EQ 10GM BASE/VIALx

CHLORDIAZEPOXIDE (PAGE 3-33)

CAPSULE, CONTROLLED RELEASE; ORAL
 LIBRELEASE
 HOFFMANN-LA ROCHE 30MGx

DEXAMETHASONE (PAGE 3-49)

SOLUTION; ORAL
 DEXAMETHASONE
 ROXANE LABORATORIES 0.5MG/5MLX
 DEXAMETHASONE INTENSOL
 ROXANE LABORATORIES 0.5MG/0.5MLX

TABLET; ORAL
 DECADRON
 BP MS&D/MERCK 6MG
 DEXAMETHASONE
 BP PAR PHARMACEUTICAL 6MGX
 BP ROXANE LABORATORIES 6MGX
 1MGX

DEXAMETHASONE SODIUM PHOSPHATE (PAGE 3-50)

INJECTABLE; INJECTION
/HEXADROL PHOSPHATE/
HEXADROL
 AP ORGANON/AKZONA EQ 4MG PHOSPHATE/ML
EQ 20MG PHOSPHATE/ML

SOLUTION/DROPS; OPHTHALMIC

> ADD > DEXAIR
 > ADD > AT PHARMAFAIR EQ 0.1% PHOSPHATEX
DEXAMETHASONE SODIUM PHOSPHATE
 > ADD > AT BARNES-HIND PHARMS EQ 0.1% PHOSPHATE

DEXTROSE; DOPAMINE HYDROCHLORIDE (PAGE 3-53)

INJECTABLE; INJECTION
/DOPAMINE HCL IN DEXTROSE 5%/
DOPAMINE HCL
 AP ABBOTT LABORATORIES 5GM/100ML;80MG/100ML
 AP 5GM/100ML;160MG/100ML
DOPAMINE HCL IN PLASTIC CONTAINER
 AP ABBOTT LABORATORIES 5GM/100ML;80MG/100MLX
 AP 5GM/100ML;160MG/100MLX
5GM/100ML;320MG/100MLX

DEXTROSE; HEPARIN SODIUM (PAGE 3-53)

INJECTABLE; INJECTION
 HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER
 TRAVENOL LABS 5GM/100ML;4,000 UNITS/100MLX

DIATRIZOATE MEGGLUMINE; DIATRIZOATE SODIUM (PAGE 3-57)

INJECTABLE; INJECTION
/DIATRIZOATE MEGGLUMINE AND DIATRIZOATE SODIUM/
DIATRIZOATE-60

DIETHYLPROPION HYDROCHLORIDE (PAGE 3-59)

TABLET; ORAL
DIETHYLPROPION HCL
 AA CAMALL 25MGX

DIMENHYDRINATE (PAGE 3-60)

INJECTABLE; INJECTION
DIMENHYDRINATE
 /AA/ /INTL MEDICATION SYS//50MG/ML/

DIPHENHYDRAMINE HYDROCHLORIDE (PAGE 3-61)

CAPSULE; ORAL
SK-DIPHENHYDRAMINE
 /AA/ SK&F LABORATORIES /25MG/

ELIXIR; ORAL

BELIX
 AA HALSEY DRUG 12.5MG/5MLX
 > ADD > DIBENIL
 > ADD > AA HR CENCI LABS 12.5MG/5MLX
 /AA/ /SK-DIPHENHYDRAMINE/
/SK&F LABORATORIES/ /12.5MG/5ML/

DISULFIRAM (PAGE 3-63)

TABLET; ORAL
 DISULFIRAM
 BX CHELSEA LABORATORIES 250MGX
 BX 500MGX
 > ADD > BX SIDMAK LABORATORIES 250MGX
 > ADD > BX 500MGX

DOXYCYCLINE HYCLATE (PAGE 3-64)

CAPSULE; ORAL
DOXYCYCLINE HYCLATE
 AB CHELSEA LABORATORIES EQ 50MG BASE
 > ADD > AB HEATHER DRUG EQ 50MG BASEX
 > ADD > AB EQ 100MG BASEX
 > ADD > AB PAR PHARMACEUTICAL EQ 100MG BASEX
 > ADD > AB PUREPAC/KALIPharma EQ 50MG BASEX
 > ADD > AB EQ 100MG BASEX

INJECTABLE; INJECTION
 > ADD > DOXY 100
 > ADD > AP LYPHO-MED EQ 100MG BASE/VIALX
 > ADD > DOXY 200
 > ADD > AP LYPHO-MED EQ 200MG BASE/VIALX

GENTAMICIN SULFATE (PAGE 3-79)

INJECTABLE; INJECTION

<u>GENTAMICIN SULFATE</u>		
AP	ABBOTT LABORATORIES	<u>EQ 60MG BASE/100ML^x</u>
AP		<u>EQ 70MG BASE/100ML^x</u>
AP		<u>EQ 80MG BASE/100ML^x</u>
AP		<u>EQ 90MG BASE/100ML^x</u>
AP		<u>EQ 100MG BASE/100ML^x</u>
AP		<u>EQ 1.2MG BASE/ML^x</u>
AP		<u>EQ 1.4MG BASE/ML^x</u>
AP		<u>EQ 1.6MG BASE/ML^x</u>
AP		<u>EQ 1.8MG BASE/ML^x</u>
AP		<u>EQ 2MG BASE/ML^x</u>
AP		<u>EQ 10MG BASE/ML^x</u>
AP		<u>EQ 40MG BASE/ML^x</u>
AP	LYPHO-MED	<u>EQ 40MG BASE/ML^x</u>
<u>GENTAMICIN SULFATE IN PLASTIC CONTAINER</u>		
AP	ABBOTT LABORATORIES	<u>EQ 60MG BASE/100ML^x</u>
AP		<u>EQ 70MG BASE/100ML^x</u>
AP		<u>EQ 80MG BASE/100ML^x</u>
AP		<u>EQ 90MG BASE/100ML^x</u>
AP		<u>EQ 100MG BASE/100ML^x</u>
AP		<u>EQ 1.2MG BASE/ML^x</u>
AP		<u>EQ 1.4MG BASE/ML^x</u>
AP		<u>EQ 1.6MG BASE/ML^x</u>
AP		<u>EQ 1.8MG BASE/ML^x</u>
AP		<u>EQ 2MG BASE/ML^x</u>

OINTMENT; TOPICAL

> ADD >	<u>GENTAMICIN SULFATE</u>	
> ADD >	AT THAMES PHARMACAL	<u>EQ 1MG BASE/GM^x</u>

GONADOTROPIN, CHORIONIC (PAGE 3-81)

INJECTABLE; INJECTION

<u>/AA/</u>	<u>/ANTUITRIN'S/</u>	<u>/5,000 UNITS/VIAL/</u>
	<u>/PARKE-DAVIS/M-L/</u>	

GRISEOFULVIN, ULTRAMICROCRYSTALLINE (PAGE 3-82)

TABLET; ORAL

AB	<u>FULVICIN P/G 165</u>	<u>165MG</u>
	SCHERING	
AB	<u>FULVICIN P/G 330</u>	<u>330MG</u>
	SCHERING	
AB	<u>GRISACTIN ULTRA</u>	<u>165MG^x</u>
	AYERST LABS/AMHO	
AB		<u>330MG^x</u>

HEPARIN SODIUM (PAGE 3-83)

INJECTABLE; INJECTION

AP	<u>HEPARIN SODIUM</u>	
	NATCON CHEMICAL	<u>1,000 UNITS/ML^x</u>

HOMATROPINE METHYLBROMIDE (PAGE 3-86)

TABLET; ORAL

<u>/AA/</u>	<u>HOMAPIN-10</u>	<u>10MG</u>
	MISSION PHARMACAL	
<u>/AA/</u>	<u>/SEUS-TENS'SE/</u>	<u>/10MG/</u>
	/LEMON/	

HYDRALAZINE HYDROCHLORIDE (PAGE 3-86)

TABLET; ORAL

AA	<u>HYDRALAZINE HCL</u>	<u>100MG^x</u>
	PAR PHARMACEUTICAL	
AA	PUREPAC/KALIPHARMA	<u>50MG^x</u>

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE (PAGE 3-87)

TABLET; ORAL

<u>/R-HCTZ-H/</u>		
	RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HCL	
BP	RESERPINE, HYDRALAZINE HCL, AND HYDROCHLOROTHIAZIDE	
	REID-PROVIDENT LABS	<u>25MG;15MG;0.1MG^x</u>

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE (PAGE 3-89)

TABLET; ORAL

AB	<u>SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE</u>	<u>25MG;25MG^x</u>
	PUREPAC/KALIPHARMA	

HYDROCORTISONE (PAGE 3-90)

CREAM; TOPICAL

<u>/AT/</u>	<u>ELDECORT</u>	<u>/0.5%/</u>
	ELDER PHARMS	
AT	<u>HYDROCORTISONE</u>	<u>1%^x</u>
	BAY LABORATORIES	
AT		<u>2.5%^x</u>

LOTION; TOPICAL

<u>/AT/</u>	<u>/dELAcort/</u>	<u>/0.5%/</u>
	/MERICON INDUSTRIES/	
AT	<u>GLY-CORT</u>	<u>1%^x</u>
	HERAN PHARMACEUTICAL	
AT	<u>HYDROCORTISONE</u>	<u>0.5%</u>
	MERICON INDUSTRIES	
<u>/AT/</u>	<u>/TOWNE PAULSEN/</u>	<u>/0.5%/</u>

OINTMENT; TOPICAL

AT	<u>HYDROCORTISONE</u>	<u>1%^x</u>
	BAY LABORATORIES	
AT		<u>2.5%^x</u>

LIDOCAINE HYDROCHLORIDE (PAGE 3-104)

INJECTABLE; INJECTION
LIDOCATON
 AP PHARMATON/SZ 22M

METHYLTESTOSTERONE (PAGE 3-121)

TABLET; BUCCAL/SUBLINGUAL
 METHYLTESTOSTERONE
 /b/ /f'OHNE'PAULSEN/ /10M/

METRONIDAZOLE (PAGE 3-122)

INJECTABLE; INJECTION
METRONIDAZOLE
 AP ABBOTT LABORATORIES 500MG/100MLM
METRONIDAZOLE IN PLASTIC CONTAINER
 AP ABBOTT LABORATORIES 500MG/100MLM
METRO I.V. IN PLASTIC CONTAINER
 AP AM MCGAW/AM HOSP 500MG/100MLM

TABLET; ORAL
METRONIDAZOLE
 AB PAR PHARMACEUTICAL 250MGM
 AB 500MGM

NANDROLONE DECANOATE (PAGE 3-125)

INJECTABLE; INJECTION
NANDROLONE DECANOATE
 > ADD > AO CARTER-GLOGAU 200MG/MLM
 AO LEMMON 100MG/ML
 50MG/ML
 AO LYPHO-MED 100MG/MLM
 AO 200MG/MLM
 AO MAURRY BIOLOGICAL 100MG/MLM

NITROGLYCERIN (PAGE 3-128)

INJECTABLE; INJECTION
NITRONAL
 AP G POHL-BOSKAMP 5MG/MLM
 1MG/MLM
TRIDIL
 AM CRITICAL CARE/AHS 0.5MG/MLM

NYSTATIN (PAGE 3-129)

> ADD > POWDER; ORAL
 > ADD > NILSTAT
 > ADD > LEDERLE LABS/AM CYAN 100M

TABLET; ORAL

NYSTATIN
 > ADD > AA PAR PHARMACEUTICAL 500,000 UNITSM

TABLET; VAGINAL

NYSTATIN
 AT E FOUGERA/BYK-GLDN 100,000 UNITSM
 AT PHARMADERM/BYK-GLDN 100,000 UNITSM

> ADD > OPRENOLOL HYDROCHLORIDE (PAGE 3-131)

> ADD > CAPSULE; ORAL
 > ADD > TRASICOR
 > ADD > CIBA/CIBA-GEIGY 20MGM
 > ADD > 40MGM
 > ADD > 80MGM
 > ADD > 160MGM

OXTRIPHYLLINE (PAGE 3-131)

> ADD > ELIXIR; ORAL
 > ADD > OXTRIPHYLLINE
 > ADD > BAY LABORATORIES 100MG/5MLM

SYRUP; ORAL

CHOLEDYL
 > ADD > AA PARKE-DAVIS/W-L 50MG/5ML
 > ADD > OXTRIPHYLLINE PEDIATRIC
 > ADD > AA BAY LABORATORIES 50MG/5MLM

TABLET, ENTERIC COATED; ORAL

CHOLEDYL
 AB PARKE-DAVIS/W-L 100MG
 AB 200MG
OXTRIPHYLLINE
 AB BOLAR PHARMACEUTICAL 100MGM
 AB 200MGM

OXYTETRACYCLINE HYDROCHLORIDE (PAGE 3-132)

CAPSULE; ORAL
 /AB/ /OXLOPAR/ /e'250MG'BASE/

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE
(PAGE 3-155)

TABLET; ORAL
TRIPROLIDINE HCL AND PSEUDOEPHEDRINE HCL
AA CHELSEA LABORATORIES 60MG;2.5MG

QUINIDINE SULFATE (PAGE 3-157)

TABLET; ORAL
QUINIDINE SULFATE
AB VITARINE/WEST CHEM 300MG

RESERPINE (PAGE 3-158)

TABLET; ORAL
/BP/ /RAURINE/
/BP/ /ONEAL, JONES & FELDMAN/ /0.25MG/
/BP/ /VIO-SERPINE/
/BP/ /ROWELL LABORATORIES/ /0.1MG/
/BP/ /0.25MG/

SECOBARBITAL SODIUM (PAGE 3-160)

CAPSULE; ORAL
SECOBARBITAL SODIUM
AA LANNETT 50MG
AA SECONAL SODIUM
AA ELI LILLY 50MG
AA 100MG

SELENIUM SULFIDE (PAGE 3-161)

LOTION/SHAMPOO; TOPICAL
SELENIUM SULFIDE
AT BAY LABORATORIES 2.5%

SILVER SULFADIAZINE (PAGE 3-161)

CREAM; TOPICAL
SILVER SULFADIAZINE
SSD

SITOSTEROL, ALPHA (PAGE 3-161)

/SUSPENSION; ORAL/
/CYTELLIN/
/ELI LILLY/ /30M/15ML/

SODIUM CHLORIDE (PAGE 3-162)

INJECTABLE; INJECTION
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER
TRAVENOL LABS 3GM/100ML \times
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER
TRAVENOL LABS 5GM/100ML \times

SODIUM POLYSTYRENE SULFONATE (PAGE 3-165)

SUSPENSION; ORAL, RECTAL
SODIUM POLYSTYRENE SULFONATE
AA ROXANE LABORATORIES 15GM/60ML
SPS
AA CAROLINA MED PRODS 15GM/60ML

SOMATROPIN (PAGE 3-165)

INJECTABLE; INJECTION
ASELLACRIN 2 2 IU/VIAL \times
SERONO LABS
/ASELLACRIN/
ASELLACRIN 10

SPIRONOLACTONE (PAGE 3-166)

TABLET; ORAL
SPIRONOLACTONE
AB PUREPAC/KALIPHARMA 25MG

SULFACETAMIDE SODIUM (PAGE 3-167)

SOLUTION/DROPS; OPHTHALMIC
SODIUM SULFACETAMIDE
> DLT > /At/ /KETCHUM LABORATORIES/ /10%/
> DLT > /At/ /30%/
> ADD > AT OPTOPICS LABS 10%
> ADD > AT 30%
SULFACEL-15
> DLT > /At/ /STERI-MED/KETCHUM/ /15%/
> ADD > AT OPTOPICS LABS 15%
SULFAIR FORTE
AT PHARMAFAIR 30%

TESTOSTERONE (PAGE 3-173)

INJECTABLE; INJECTION
TESTOSTERONE
CARTER-GLOGAU LABS 50MG/ML \times

THIORIDAZINE HYDROCHLORIDE (PAGE 3-176)

AB	TABLET; ORAL	10MG
AB	THIORIDAZINE HCL	10MG
AB	BARR LABORATORIES	15MG
AB		25MG
AB		50MG
AB		100MG
AB	BOLAR PHARMACEUTICAL	10MG
AB		100MG
AB	CORD LABORATORIES	10MG
AB		15MG
AB		25MG
AB		50MG
AB	DANBURY PHARMACAL	10MG
AB		15MG
AB		25MG
AB		50MG
AB	MYLAN PHARMS	100MG
AB		10MG
AB	PAR PHARMACEUTICAL	10MG
AB		15MG
AB		25MG
AB		50MG
AB		100MG
AB	ZENITH LABORATORIES	100MG

TIMOLOL MALEATE (PAGE 3-179)

TABLET; ORAL	BLOCDAREN	5MG
	MS&D/MERCK	

TRIAMCINOLONE ACETONIDE (PAGE 3-180)

< DLT	AEROSOL; INHALATION	
< DLT	/ARISTOCORT/	
< DLT	/FEDERLE LABS/AM, CYAN/D, ZENITH/INH/	
< ADD	AZMACORT	
< ADD	WILLIAM H RORER	0.25MG/INH

AT	CREAM; TOPICAL	0.025%
AT	TRIAMCINOLONE ACETONIDE	0.1%
AT	BAY LABORATORIES	0.5%

AT	OINTMENT; TOPICAL	0.025%
AT	TRIAMCINOLONE ACETONIDE	0.1%
AT	BAY LABORATORIES	0.5%

TETRACYCLINE HYDROCHLORIDE (PAGE 3-174)

AB	CAPSULE; ORAL	10MG
AB	TETRACYCLINE HCL	10MG
AB	/LEHMAN/	15MG
AB		25MG
AB		50MG
AB		100MG
AB	BARR LABORATORIES	10MG
AB		100MG
AB	BOLAR PHARMACEUTICAL	10MG
AB		100MG
AB	CORD LABORATORIES	10MG
AB		15MG
AB		25MG
AB		50MG
AB		100MG
AB	DANBURY PHARMACAL	10MG
AB		15MG
AB		25MG
AB		50MG
AB	MYLAN PHARMS	100MG
AB		10MG
AB	PAR PHARMACEUTICAL	10MG
AB		15MG
AB		25MG
AB		50MG
AB		100MG
AB	ZENITH LABORATORIES	100MG

THALLOUS CHLORIDE, TL-201 (PAGE 3-175)

INJECTABLE; INJECTION	THALLOUS CHLORIDE TL 201	2MCI/ML
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THEOPHYLLINE (PAGE 3-176)

BC	SEARLE/SEARLE PHARMS	100MG
BC	THEO-24	100MG
BC	FISONS	100MG
BC	SOMOPHYLLIN-CRI	100MG
BC	CAPSULE, CONTROLLED RELEASE; ORAL	
	(FOR CONTROLLED RELEASE PRODUCTS - SEE SPECIAL NOTE C.)	

THEOPHYLLINE (PAGE 3-176)

BC	SEARLE/SEARLE PHARMS	100MG
BC	THEO-24	100MG
BC	FISONS	100MG
BC	SOMOPHYLLIN-CRI	100MG

THEOPHYLLINE (PAGE 3-176)

AB	SOLUTION; ORAL	300MG
AB	THEOLAIR	300MG
AB	RIKER LABS/3M	300MG
AB	THEOPHYLLINE	300MG
AB	ROXANE LABORATORIES	300MG

THEOPHYLLINE (PAGE 3-176)

BC	THEOCONTIN	200MG
BC	PURDUE FREDERICK	200MG
BC	KEY PHARMACEUTICALS	200MG

THIORIDAZINE HYDROCHLORIDE (PAGE 3-176)

AA	CONCENTRATE; ORAL	30MG/ML
AA	THIORIDAZINE HCL	30MG/ML
AA	CORD LABORATORIES	30MG/ML
AA	NATL PHARM MFG/BARRE	100MG/ML
AA	MELLARIL	100MG/ML
AB	TABLET; ORAL	100MG
AB	SANDOZ PHARMS/SANDOZ	100MG

TRISULFAPYRIMIDINES (PAGE 3-185)

SUSPENSION; ORAL

/QUAD-RAMID/
/AB/ /ELDER PHARMS/ /500MG/5ML/
/TRISEN/
/AB/ /BEECHAM LABS/BEECHAM/500MG/5ML/
/TRISUREID/
/AB/ /REID-PROVIDENT LABS//500MG/5ML/

TABLET; ORAL

/QUADETS/
/AB/ /ELDER PHARMS/ /500MG/
/TRIPLE SULFA #2/
/AB/ /ZENITH LABORATORIES//500MG/

TROPICAMIDE (PAGE 3-186)

SOLUTION/DROPS; OPHTHALMIC

MYDRIAFAIR
AT PHARMAFAIR 0.5%N
AT 1%N

TYBAMATE (PAGE 3-186)

/CAPSULE; ORAL/
/TYBATRAN/
/AH. ROBINS/ /250MG/
/350MG/

VITAMIN A PALMITATE (PAGE 3-188)

CAPSULE; ORAL

/SOLVITSYN A/
/AA/ /TOWNE PAULSEN/ /EQ 50,000 UNITS BASE/

CURRENT STATUS - INEFFECTIVE

/CARBRITAL/ /PARKE-DAVIS/W-L/
/CARBRONAL; PENTOBARBITAL SODIUM/

/CLISTIN RA/ /MCNEIL PHARM/
/CARBINOXAMINE MALEATE/

> ADD > DIMETAPP AH ROBINS
> ADD > BROMPHENIRAMINE MALEATE; PHENYLEPHRINE HYDROCHLORIDE;
> ADD > PHENYLPROPANOLAMINE HYDROCHLORIDE

> ADD > ELIXIR DIMETAPP AH ROBINS
> ADD > BROMPHENIRAMINE MALEATE; PHENYLEPHRINE HYDROCHLORIDE;
> ADD > PHENYLPROPANOLAMINE HYDROCHLORIDE

/FORHISTAL/ /CIBA/CIBA-GEIGY/
/DIMETHINDENE MALEATE/

METHANDROSTENOLONE PAR PHARMACEUTICAL
METHANDROSTENOLONE

TUSS-ORNADE SK&F LABORATORIES
CARAMIPHEN EDISYLATE; CHLORPHENIRAMINE MALEATE;
ISOPROPAMIDE IODIDE; PHENYLPROPANOLAMINE HYDROCHLORIDE

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