EXHIBIT A

Electronic Orange Book Pages Showing
the Listing of the Listed Drug:
Merck's Mefoxin®

Electronic Orange Book

Approved Drug Products
with
Therapeutic Equivalence Evaluations

Current through March 2003

Preface

FAQ

Search by Active Ingredient Search by Applicant Holder

Search by Proprietary Name Search by Application Number

The products in this list have been approved under section 505 of the Federal Food, Drug, and Cosmetic Act.

Drug questions email: DRUGINFO@CDER.FDA.GOV

U.S Department of Health and Human Services
Food and Drug Administration
Center for Drug Evaluation and Research
Office of Pharmaceutical Science
Office of Generic Drugs

Updated: May 1, 2003

Proprietary Name Search Results from "Rx" table for query on "mefoxin."

Appl No	TE Code	RLD	Active Ingredient	Dosage Form; Route	Strength	Proprietary Name	Ap
050517	AP	Yes	CEFOXITIN SODIUM	Injectable; Injection	EQ 10GM BASE/VIAL	MEFOXIN	ME
050517	AP	Yes	CEFOXITIN SODIUM	Injectable; Injection	EQ 1GM BASE/VIAL	MEFOXIN	ME
062757	AP	No	CEFOXITIN SODIUM	Injectable; Injection	EQ 1GM BASE/VIAL	MEFOXIN	ME
050517	AP	Yes	CEFOXITIN SODIUM	Injectable; Injection	EQ 2GM BASE/VIAL	MEFOXIN	ME
062757	AP	No	CEFOXITIN SODIUM	Injectable; Injection	EQ 2GM BASE/VIAL	MEFOXIN	ME
063182		Yes	CEFOXITIN SODIUM	Injectable; Injection	EQ 20MG BASE/ML	MEFOXIN IN PLASTIC CONTAINER	ME
063182		Yes	CEFOXITIN SODIUM	Injectable; Injection	EQ 40MG BASE/ML	MEFOXIN IN PLASTIC CONTAINER	ME

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Search results from the "Rx" table for query on "050517."

Active Ingredient:	CEFOXITIN SODIUM
Dosage Form;Route:	Injectable; Injection
Proprietary Name	MEFOXIN
Applicant:	MERCK
Strength:	EQ 1GM BASE/VIAL
Application Number:	050517
Product Number:	001
Approval Date:	Approved prior to Jan 1, 1982
Reference Listed Drug:	Yes
RX/OTC/DISCN:	RX
TE Code:	AP
Patent and Exclusivity Info for this product:	Click Here

Active Ingredient:	CEFOXITIN SODIUM
Dosage Form;Route:	Injectable; Injection
Proprietary Name	MEFOXIN
Applicant:	MERCK
Strength:	EQ 2GM BASE/VIAL
Application Number:	050517
Product Number:	002
Approval Date:	Approved prior to Jan 1, 1982
Reference Listed Drug:	Yes
RX/OTC/DISCN:	RX
TE Code:	AP
Patent and Exclusivity Info for this product:	Click Here

Active Ingredient:	CEFOXITIN SODIUM
Dosage Form;Route:	Injectable; Injection
Proprietary Name	MEFOXIN
Applicant:	MERCK

Strength:	EQ 10GM BASE/VIAL
Application Number:	050517
Product Number:	003
Approval Date:	Approved prior to Jan 1, 1982
Reference Listed Drug:	Yes
RX/OTC/DISCN:	RX
TE Code:	AP

Patent and Exclusivity Info for this product: Click Here

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