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Electronic Orange Book

Approved Drug Products
with
Therapeutic Equivalence Evaluations

Current through July 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: August 21, 2003

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

Appl No	TE Code	RLD	Active Ingredient	Dosage Form; Route	Strength	Proprietary Name	Applicant
020716	AB	Yes	HYDROCODONE BITARTRATE; IBUPROFEN	Tablet; Oral	7.5MG;200MG	VICOPROFEN	ABBOTT

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Search results from the "Rx" table for query on "020716."

Active Ingredient:	HYDROCODONE BITARTRATE; IBUPROFEN
Dosage Form;Route:	Tablet; Oral
Proprietary Name	VICOPROFEN
Applicant:	ABBOTT
Strength:	7.5MG;200MG
Application Number:	020716
Product Number:	001
Approval Date:	SEP 23, 1997
Reference Listed Drug:	Yes
RX/OTC/DISCN:	RX
TE Code:	AB
Patent and Exclusivity Info for this product:	Click Here

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