Electronic Orange Book

Approved Drug Products
with
Therapeutic Equivalence Evaluations

Current through September 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: October 20, 2003

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

Appl No	TE Code	RLD	Active Ingredient	Dosage Form; Route	Strength	Proprietary Name	Applicant
021046		Yes	CITALOPRAM HYDROBROMIDE	Solution; Oral	EQ 10MG BASE/5ML	CELEXA	FOREST LABS
020822		No	CITALOPRAM HYDROBROMIDE	Tablet; Oral	EQ 10MG BASE	CELEXA	FOREST LABS
020822		No	CITALOPRAM HYDROBROMIDE	Tablet; Oral	EQ 20MG BASE	CELEXA	FOREST LABS
020822		Yes	CITALOPRAM HYDROBROMIDE	Tablet; Oral	EQ 40MG BASE	CELEXA	FOREST LABS

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