ATTACHMENT 1 REFERENCE LISTED DRUG (RLD)

Orange Book: Proprietary Name Search

Page 1 of 1

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

Appl No	<u>TE</u> Code	RLD	Active Ingredient	Dosage Form; Route	Strength	Proprietary Name	Applicant
020232	AB		Indiana I formati	Capsule; Oral	325MG;50MG;40MG;30MG		WATSON PHARMS

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Search results from the "Rx" table for query on "020232."

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE Active Ingredient:

WATSON PHARMS

PHOSPHATE

Dosage Form; Route: Capsule; Oral

Proprietary Name FIORICET W/ CODEINE

Applicant: Strength: 325MG;50MG;40MG;30MG

Application Number: 020232 Product Number: 001

Approval Date: JUL 30, 1992

Reference Listed Drug: Yes RX/OTC/DISCN: RX TE Code: AB

Patent and Exclusivity Info for this

product:

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Patent and Exclusivity Search Results from query on 020232 001.

Patent Data

There are no unexpired patents for this product in the Orange Book Database.

[Note: Title I of the 1984 Amendments does not apply to drug products submitted or approved under the former Section 507 of the Federal Food, Drug and Cosmetic Act (antibiotic products). Drug products of this category will not have patents listed.]

Exclusivity Data

There is no unexpired exclusivity for this product.

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Patent and Exclusivity Terms

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