Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

Search results from the "OB_Disc" table for query on "019329."

Active Ingredient:	SODIUM CHLORIDE
Dosage Form;Route:	INJECTABLE;INJECTION
Proprietary Name:	SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER
Applicant:	ABRAXIS PHARM
Strength:	234MG/ML
Application Number:	N019329
Product Number:	001
Approval Date:	Apr, 22, 1987
RX/OTC/DISCN:	DISCN
Patent and Exclusivity Info for this product:	View

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FDA/Center for Drug Evaluation and Research
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