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Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations

Search results from the "OB_Disc" table for query on "021664."

Active Ingredient:	BROMFENAC SODIUM
Dosage Form;Route:	SOLUTION/DROPS;OPHTHALMIC
Proprietary Name:	XIBROM
Applicant:	BAUSCH AND LOMB INC
Strength:	EQ 0.09% ACID
Application Number:	N021664
Product Number:	001
Approval Date:	Mar, 24, 2005
RX/OTC/DISCN:	DISCN
Patent and Exclusivity Info for this product:	View

Active Ingredient:	BROMFENAC SODIUM
Dosage Form;Route:	SOLUTION/DROPS;OPHTHALMIC
Proprietary Name:	BROMDAY
Applicant:	BAUSCH AND LOMB INC
Strength:	EQ 0.09% ACID **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**
Application Number:	N021664
Product Number:	002
Approval Date:	Oct, 16, 2010
RX/OTC/DISCN:	DISCN
Patent and Exclusivity Info for this product:	View

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