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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 Strength: EQ 0.09% ACID **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**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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 Office of Generic Drugs
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Generic Drug Product Information & Patent Information - **Daily**

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Patent and Generic Drug Product Data Last Updated March 11, 2016

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