



U.S. Food and Drug Administration

Protecting and Promoting *Your* Health

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Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations
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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
 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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FDA/Center for Drug Evaluation and Research
 Office of Generic Drugs
 Division of Labeling and Program Support
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Generic Drug Product Information & Patent Information - **Daily**

Orange Book Data Updated Through September 2015

Patent and Generic Drug Product Data Last Updated November 09, 2015

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