Drug Details

Drug Name(s)	**BROMDAY**
FDA Application No.	**(NDA) 021664**
Active Ingredient(s)	**BROMFENAC SODIUM**
Company	**BAUSCH AND LOMB INC**
Original Approval or Tentative Approval Date	**March 24, 2005**
Chemical Type	**3 New dosage form**
Review Classification	**S Standard review drug**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**

Products on Application (NDA) #021664
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
BROMDAY	BROMFENAC SODIUM	EQ 0.09% ACID	SOLUTION/DROPS;OPHTHALMIC	Prescription	Yes	AT2
XIBROM	BROMFENAC SODIUM	EQ 0.09% ACID	SOLUTION/DROPS;OPHTHALMIC	Discontinued	No	None

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