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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

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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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