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Drugs@FDA
FDA Approved Drug Products

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Drug Details

Drug Name(s)	**PROLENSA**
FDA Application No.	**(NDA) 203168**
Active Ingredient(s)	**BROMFENAC SODIUM**
Company	**BAUSCH AND LOMB**
Original Approval or Tentative Approval Date	**April 5, 2013**
Chemical Type	**5 New formulation or new manufacturer**
Review Classification	**S Standard review drug**

- **There are no Therapeutic Equivalents**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (NDA) #203168
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
PROLENSA	BROMFENAC SODIUM	EQ 0.07% ACID	SOLUTION/DROPS;OPHTHALMIC	Prescription	Yes	None

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