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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](#)
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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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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)
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