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## Drug Details

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

- [There are no Therapeutic Equivalents](#)
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- [Label Information](#)

### Products on Application (NDA) #203168

Click on a column header to re-sort the table:

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

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