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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](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- [Label Information](#)

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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