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## Drug Details

<b>Drug Name(s)</b>	<b>PROFENAL</b>
<b>FDA Application No.</b>	<b>(NDA) 019387</b>
<b>Active Ingredient(s)</b>	<b>SUPROFEN</b>
<b>Company</b>	<b>ALCON</b>
<b>Original Approval or Tentative Approval Date</b>	<b>December 23, 1988</b>
<b>Chemical Type</b>	<b>3 New dosage form</b>
<b>Review Classification</b>	<b>S Standard review drug</b>

- There are no Therapeutic Equivalents
- [Approval History, Letters, Reviews, and Related Documents](#)
- Labels are not available

## Products on Application (NDA) #019387

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>
PROFENAL	SUPROFEN	1%	SOLUTION/DROPS;OPHTHALMIC	Discontinued	No None

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