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Drug Details

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

- 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:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
PROFENAL	SUPROFEN	1%	SOLUTION/DROPS;OPHTHALMIC	Discontinued	No None

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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