Drugs@FDA
FDA Approved Drug Products

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**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

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

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