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Drug Details

Drug Name(s)	NEVANAC
FDA Application No.	(NDA) 021862
Active Ingredient(s)	NEPAFENAC
Company	ALCON PHARMS LTD
Original Approval or Tentative Approval Date	August 19, 2005
Chemical Type	1 New molecular entity (NME)
Review Classification	P Priority review drug

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #021862

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NEVANAC	NEPAFENAC	0.1%	SUSPENSION/DROPS;OPHTHALMIC	Prescription	Yes	None

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

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Lupin EX1166
Page 1

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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)
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