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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**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

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

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