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Drug Details

Drug Name(s)	DICLOFENAC SODIUM
FDA Application No.	(ANDA) 208301
Active Ingredient(s)	DICLOFENAC SODIUM
Company	GLENMARK PHARMS LTD
Original Approval or Tentative Approval Date	September 13, 2016

- [Therapeutic Equivalents](#)
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- **Labels are not available**

Products on Application (ANDA) #208301

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
DICLOFENAC SODIUM	DICLOFENAC SODIUM	3%	GEL;TOPICAL	Prescription	No	AB

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