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Drug Details

Drug Name(s)	DICLOFENAC SODIUM
FDA Application No.	(ANDA) 206493
Active Ingredient(s)	DICLOFENAC SODIUM
Company	ACTAVIS MID ATLANTIC
Original Approval or Tentative Approval Date	December 2, 2015

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #206493

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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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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