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Drug Details

Drug Name(s)	RUFINAMIDE
FDA Application No.	(ANDA) 205075
Active Ingredient(s)	RUFINAMIDE
Company	GLENMARK PHARMS LTD
Original Approval or Tentative Approval Date	May 16, 2016

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

Products on Application (ANDA) #205075

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
RUFINAMIDE	RUFINAMIDE	200MG	TABLET;ORAL	Prescription	No	AB
RUFINAMIDE	RUFINAMIDE	400MG	TABLET;ORAL	Prescription	No	AB

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

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