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Drug Details

Drug Name(s)	NATEGLINIDE
FDA Application No.	(ANDA) 205248
Active Ingredient(s)	NATEGLINIDE
Company	ZYDUS PHARMS USA INC
Original Approval or Tentative Approval Date	July 6, 2016

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

Products on Application (ANDA) #205248

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NATEGLINIDE	NATEGLINIDE	60MG	TABLET;ORAL	Prescription	No	AB
NATEGLINIDE	NATEGLINIDE	120MG	TABLET;ORAL	Prescription	No	AB

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