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Drug Details

Drug Name(s)	EPTIFIBATIDE
FDA Application No.	(ANDA) 206127
Active Ingredient(s)	EPTIFIBATIDE
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	December 8, 2015

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

Products on Application (ANDA) #206127

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
EPTIFIBATIDE	EPTIFIBATIDE	2MG/ML	INJECTABLE;INJECTION	Prescription	No	AP
EPTIFIBATIDE	EPTIFIBATIDE	75MG/100ML	INJECTABLE;INJECTION	Prescription	No	AP

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