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

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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