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Drug Details

Drug Name(s)	ESOMEPRAZOLE SODIUM
FDA Application No.	(ANDA) 204657
Active Ingredient(s)	ESOMEPRAZOLE SODIUM
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	August 11, 2016

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

Products on Application (ANDA) #204657

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
ESOMEPRAZOLE SODIUM	ESOMEPRAZOLE SODIUM	EQ 20MG BASE/VIAL	INJECTABLE;INTRAVENOUS	Prescription No	AP
ESOMEPRAZOLE SODIUM	ESOMEPRAZOLE SODIUM	EQ 40MG BASE/VIAL	INJECTABLE;INTRAVENOUS	Prescription No	AP

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