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Drug Details

Drug Name(s)	PANTOPRAZOLE SODIUM
FDA Application No.	(ANDA) 205675
Active Ingredient(s)	PANTOPRAZOLE SODIUM
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	March 30, 2016

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- **Labels are not available**

Products on Application (ANDA) #205675

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
PANTOPRAZOLE SODIUM	PANTOPRAZOLE SODIUM	EQ 40MG BASE/VIAL	INJECTABLE;IV (INFUSION)	Prescription No	AP

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