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Drug Details

Drug Name(s)	PALONOSETRON HYDROCHLORIDE
FDA Application No.	(ANDA) 201533
Active Ingredient(s)	PALONOSETRON HYDROCHLORIDE
Company	DR REDDYS LABS LTD
Original Approval or Tentative Approval Date	April 21, 2016

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

Products on Application (ANDA) #201533

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
PALONOSETRON HYDROCHLORIDE	PALONOSETRON HYDROCHLORIDE	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	INJECTABLE;INTRAVENOUS	Prescription No	AP
PALONOSETRON HYDROCHLORIDE	PALONOSETRON HYDROCHLORIDE	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	INJECTABLE;INTRAVENOUS	Prescription No	AP

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