Drug Details

Drug Name(s)	**GRANISETRON HYDROCHLORIDE**
FDA Application No.	**(ANDA) 204238**
Active Ingredient(s)	**GRANISETRON HYDROCHLORIDE**
Company	**AUROBINDO PHARMA LTD**
Original Approval or Tentative Approval Date	**July 6, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #204238
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
GRANISETRON HYDROCHLORIDE	GRANISETRON HYDROCHLORIDE	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	INJECTABLE;INJECTION	Prescription	No	AP
GRANISETRON HYDROCHLORIDE	GRANISETRON HYDROCHLORIDE	EQ 4MG BASE/4ML (EQ 1MG BASE/ML)	INJECTABLE;INJECTION	Prescription	No	AP

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