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## Drug Details

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

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- [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:

<a href="#">Drug Name</a>	<a href="#">Active Ingredients</a>	<a href="#">Strength</a>	<a href="#">Dosage Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLDTE Code</a>
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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