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## Drug Details

<b>Drug Name(s)</b>	<b>RISEDRONATE SODIUM</b>
<b>FDA Application No.</b>	<b>(ANDA) 206768</b>
<b>Active Ingredient(s)</b>	<b>RISEDRONATE SODIUM</b>
<b>Company</b>	<b>AUROBINDO PHARMA LTD</b>
<b>Original Approval or Tentative Approval Date</b>	<b>October 21, 2016</b>

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### Products on Application (ANDA) #206768

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

<b>Drug Name</b>	<b>Active Ingredients</b>	<b>Strength</b>	<b>Dosage Form/Route</b>	<b>Marketing Status</b>	<b>RLD TE Code</b>
RISEDRONATE SODIUM	RISEDRONATE SODIUM	150MG	TABLET;ORAL	Prescription	No AB

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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