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## Drug Details

<b>Drug Name(s)</b>	<b>DARIFENACIN HYDROBROMIDE</b>
<b>FDA Application No.</b>	<b>(ANDA) 206743</b>
<b>Active Ingredient(s)</b>	<b>DARIFENACIN HYDROBROMIDE</b>
<b>Company</b>	<b>AUROBINDO PHARMA LTD</b>
<b>Original Approval or Tentative Approval Date</b>	<b>September 19, 2016</b>

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

### Products on Application (ANDA) #206743

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>RLDTE Code</b>
DARIFENACIN HYDROBROMIDE	DARIFENACIN HYDROBROMIDE	EQ 7.5MG BASE	TABLET, EXTENDED RELEASE;ORAL	Prescription No	AB
DARIFENACIN HYDROBROMIDE	DARIFENACIN HYDROBROMIDE	EQ 15MG BASE	TABLET, EXTENDED RELEASE;ORAL	Prescription No	AB

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