Drug Details

Drug Name(s)	**DARIFENACIN HYDROBROMIDE**
FDA Application No.	**(ANDA) 207664**
Active Ingredient(s)	**DARIFENACIN HYDROBROMIDE**
Company	**CIPLA LTD**
Original Approval or Tentative Approval Date	**September 1, 2016**

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

Products on Application (ANDA) #207664
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
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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