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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 | RLDTE 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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U.S. Food and Drug Administration
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