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Drug Details

Drug Name(s)	DROSPIRENONE AND ETHINYL ESTRADIOL
FDA Application No.	(ANDA) 202594
Active Ingredient(s)	DROSPIRENONE; ETHINYL ESTRADIOL
Company	JAI PHARMA LTD
Original Approval or Tentative Approval Date	October 22, 2015

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

Products on Application (ANDA) #202594

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
DROSPIRENONE AND ETHINYL ESTRADIOL	DROSPIRENONE; ETHINYL ESTRADIOL	3MG; 0.02MG	TABLET;ORAL	Prescription No	AB

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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)

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