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Drug Details

Drug Name(s)	NORETHINDRONE AND ETHINYL ESTRADIOL
FDA Application No.	(ANDA) 200488
Active Ingredient(s)	ETHINYL ESTRADIOL; NORETHINDRONE
Company	JAI PHARMA LTD
Original Approval or Tentative Approval Date	October 21, 2015

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

Products on Application (ANDA) #200488

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
NORETHINDRONE AND ETHINYL ESTRADIOL	ETHINYL ESTRADIOL; NORETHINDRONE	0.035MG; 0.5MG	TABLET;ORAL-28	Prescription No	AB

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