Drugs@FDA
FDA Approved Drug Products

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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