Drug Details

Drug Name(s)	**NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE**
FDA Application No.	**(ANDA) 205049**
Active Ingredient(s)	**ETHINYL ESTRADIOL; NORETHINDRONE ACETATE**
Company	**JAI PHARMA LTD**
Original Approval or Tentative Approval Date	**May 31, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Label Information**

Products on Application (ANDA) #205049
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE	ETHINYL ESTRADIOL; NORETHINDRONE ACETATE	0.01MG,0.01MG;1MG,N/A	TABLET;ORAL	Prescription	No	AB

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