Drug Details

Drug Name(s)	**NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE**
FDA Application No.	**(ANDA) 091332**
Active Ingredient(s)	**ETHINYL ESTRADIOL; NORETHINDRONE**
Company	**LUPIN LTD**
Original Approval or Tentative Approval Date	**March 23, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #091332
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 AND FERROUS FUMARATE	ETHINYL ESTRADIOL; NORETHINDRONE	0.035MG; 0.4MG	TABLET, CHEWABLE;ORAL	Prescription	No AB

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