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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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