FDA

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FDA Approved Drug Products

Drug Details

Drug Name(s)	**NORGESTIMATE AND ETHINYL ESTRADIOL**
FDA Application No.	**(ANDA) 205630**
Active Ingredient(s)	**ETHINYL ESTRADIOL; NORGESTIMATE**
Company	**LUPIN LTD**
Original Approval or Tentative Approval Date	**October 27, 2016**

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

Products on Application (ANDA) #205630
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NORGESTIMATE AND ETHINYL ESTRADIOL	ETHINYL ESTRADIOL; NORGESTIMATE	0.035MG; 0.25MG	TABLET;ORAL-28	Prescription	No	AB

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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