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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

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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	RLDTE Code
NORGESTIMATE AND ETHINYL ESTRADIOL	ETHINYL ESTRADIOL; NORGESTIMATE	0.035MG; 0.25MG	TABLET;ORAL-28	Prescription No	AB

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U.S. Food and Drug Administration
10903 New Hampshire Avenue
Silver Spring, MD 20993
Ph. 1-888-INFO-FDA (1-888-463-6332)

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