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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

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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	RLDTE 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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