Drug Details

Drug Name(s)	**LEVONORGESTREL AND ETHINYL ESTRADIOL**
FDA Application No.	**(ANDA) 203871**
Active Ingredient(s)	**ETHINYL ESTRADIOL; LEVONORGESTREL**
Company	**HAUPT PHARMA**
Original Approval or Tentative Approval Date	**November 13, 2015**

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

Products on Application (ANDA) #203871
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
LEVONORGESTREL AND ETHINYL ESTRADIOL	ETHINYL ESTRADIOL; LEVONORGESTREL	0.03MG; 0.15MG	TABLET;ORAL	Prescription No	AB

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