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## Drug Details

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

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- **Labels are not available**

### Products on Application (ANDA) #203871

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LEVONORGESTREL AND ETHINYL ESTRADIOL	ETHINYL ESTRADIOL; LEVONORGESTREL	0.03MG; 0.15MG	TABLET;ORAL	Prescription No	AB

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