Drug Details

Drug Name(s)	**DROSPIRENONE AND ETHINYL ESTRADIOL**
FDA Application No.	**(ANDA) 205876**
Active Ingredient(s)	**DROSPIRENONE; ETHINYL ESTRADIOL**
Company	**APOTEX INC**
Original Approval or Tentative Approval Date	**September 21, 2016**

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

Products on Application (ANDA) #205876
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
DROSPIRENONE AND ETHINYL ESTRADIOL	DROSPIRENONE; ETHINYL ESTRADIOL	3MG; 0.03MG	TABLET;ORAL-28	Prescription	No AB

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