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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

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- [Approval History, Letters, Reviews, and Related Documents](#)
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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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