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Drug Details

Drug Name(s)	DROSPIRENONE;ETHINYL ESTRADIOL;LEVOMEFOLATE CALCIUM;LEVOMEFOLATE CALCIUM
FDA Application No.	(ANDA) 203594
Active Ingredient(s)	DROSPIRENONE;ETHINYL ESTRADIOL;LEVOMEFOLATE CALCIUM;LEVOMEFOLATE CALCIUM
Company	WATSON LABS
Original Approval or Tentative Approval Date	October 11, 2016

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

Products on Application (ANDA) #203594
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
DROSPIRENONE;ETHINYL ESTRADIOL;LEVOMEFOLATE CALCIUM;LEVOMEFOLATE CALCIUM	DROSPIRENONE;ETHINYL ESTRADIOL;LEVOMEFOLATE CALCIUM;LEVOMEFOLATE CALCIUM	3MG; 0.03MG; 0.451MG; 0.451MG	TABLET;ORAL	None (Tentative Approval)	No None

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