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## Drug Details

**Drug Name(s)** DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM; LEVOMEFOLATE CALCIUM  
**FDA Application No. (ANDA)** 203593  
**Active Ingredient(s)** DROSPIRENONE;ETHINYL ESTRADIOL;LEVOMEFOLATE CALCIUM;LEVOMEFOLATE CALCIUM  
**Company** WATSON LABS INC

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

### Products on Application (ANDA) #203593

Click on a column header to re-sort the table:

<a href="#">Drug Name</a>	<a href="#">Active Ingredients</a>	<a href="#">Strength</a>	<a href="#">Dosage Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLDTE Code</a>
DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLATE CALCIUM; LEVOMEFOLATE CALCIUM	DROSPIRENONE;ETHINYL ESTRADIOL;LEVOMEFOLATE CALCIUM;LEVOMEFOLATE CALCIUM	3MG; 0.02MG; 0.451MG; 0.451MG	TABLET;ORAL	None (Tentative Approval)	No None

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