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	RLD	TE 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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