Drug Details

Drug Name(s)	**ESTRADIOL VALERATE AND DIENOGEST**
FDA Application No.	**(ANDA) 202349**
Active Ingredient(s)	**DIENOGEST; ESTRADIOL VALERATE**
Company	**WATSON LABS INC**
Original Approval or Tentative Approval Date	**May 6, 2016**

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

Products on Application (ANDA) #202349
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
ESTRADIOL VALERATE AND DIENOGEST	DIENOGEST; ESTRADIOL VALERATE	N/A,2MG,3MG,N/A,N/A; 3MG,2MG,2MG,1MG,N/A	TABLET;ORAL	Prescription No	AB

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