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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

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- [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; ESTRADIOL DIENOGEST	N/A,2MG,3MG,N/A,N/A; VALERATE	N/A,N/A; 3MG,2MG,2MG,1MG,N/A	TABLET;ORAL	Prescription No	AB

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