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Drug Details

Drug Name(s)	TOLTERODINE TARTRATE
FDA Application No.	(ANDA) 077006
Active Ingredient(s)	TOLTERODINE TARTRATE
Company	IVAX SUB TEVA PHARMS
Original Approval or Tentative Approval Date	February 23, 2015

- [Therapeutic Equivalents](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #077006

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
TOLTERODINE TARTRATE	TOLTERODINE TARTRATE	1MG	TABLET;ORAL	Prescription	No	AB
TOLTERODINE TARTRATE	TOLTERODINE TARTRATE	2MG	TABLET;ORAL	Prescription	No	AB

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