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Drug Details

Drug Name(s)	ALMOTRIPTAN MALATE
FDA Application No.	(ANDA) 078027
Active Ingredient(s)	ALMOTRIPTAN MALATE
Company	TEVA PHARMS USA
Original Approval or Tentative Approval Date	July 7, 2015

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

Products on Application (ANDA) #078027

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
ALMOTRIPTAN MALATE	ALMOTRIPTAN MALATE	EQ 6.25MG BASE	TABLET;ORAL	Prescription	No AB
ALMOTRIPTAN MALATE	ALMOTRIPTAN MALATE	EQ 12.5MG BASE	TABLET;ORAL	Prescription	No AB

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