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Drug Details

Drug Name(s)	VORICONAZOLE
FDA Application No.	(ANDA) 207049
Active Ingredient(s)	VORICONAZOLE
Company	VERSAPHARM INC
Original Approval or Tentative Approval Date	September 7, 2016

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

Products on Application (ANDA) #207049

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
VORICONAZOLE	VORICONAZOLE	50MG	TABLET;ORAL	Prescription	No	AB
VORICONAZOLE	VORICONAZOLE	200MG	TABLET;ORAL	Prescription	No	AB

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