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Drug Details

Drug Name(s)	NITROFURANTOIN
FDA Application No.	(ANDA) 205180
Active Ingredient(s)	NITROFURANTOIN
Company	ACTAVIS MID ATLANTIC
Original Approval or Tentative Approval Date	May 3, 2016

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

Products on Application (ANDA) #205180

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
NITROFURANTOIN	NITROFURANTOIN	25MG/5ML	SUSPENSION;ORAL	Prescription	No	AB

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