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Drug Details

Drug Name(s)	RILUZOLE
FDA Application No.	(ANDA) 204048
Active Ingredient(s)	RILUZOLE
Company	ALKEM LABS LTD
Original Approval or Tentative Approval Date	March 30, 2016

- [Therapeutic Equivalents](#)
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- **Labels are not available**

Products on Application (ANDA) # 204048

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

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

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