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Drug Details

Drug Name(s)	LEVETIRACETAM
FDA Application No.	(ANDA) 203308
Active Ingredient(s)	LEVETIRACETAM
Company	MYLAN LABS LTD
Original Approval or Tentative Approval Date	September 16, 2016

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

Products on Application (ANDA) #203308

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
LEVETIRACETAM	LEVETIRACETAM	500MG/5ML (100MG/ML)	INJECTABLE;IV (INFUSION)	Prescription	No AP

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