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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