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Drug Details

Drug Name(s)	LEVETIRACETAM
FDA Application No.	(ANDA) 205102
Active Ingredient(s)	LEVETIRACETAM
Company	SECAN PHARMS
Original Approval or Tentative Approval Date	October 24, 2016

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

Products on Application (ANDA) #205102

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	250MG	TABLET;ORAL	None (Tentative Approval)	No None
LEVETIRACETAM	LEVETIRACETAM	750MG	TABLET;ORAL	None (Tentative Approval)	No None
LEVETIRACETAM	LEVETIRACETAM	1GM	TABLET;ORAL	Prescription	No AB
LEVETIRACETAM	LEVETIRACETAM	500MG	TABLET;ORAL	Prescription	No AB

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