Drug Details

Drug Name(s)	**LAMOTRIGINE**
FDA Application No.	**(ANDA) 206382**
Active Ingredient(s)	**LAMOTRIGINE**
Company	**SCIEGEN PHARMS INC**
Original Approval or Tentative Approval Date	**June 17, 2016**

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (ANDA) #206382
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
LAMOTRIGINE	LAMOTRIGINE	25MG	TABLET, ORALLY DISINTEGRATING;ORAL	None (Tentative Approval)	No	None
LAMOTRIGINE	LAMOTRIGINE	50MG	TABLET, ORALLY DISINTEGRATING;ORAL	None (Tentative Approval)	No	None
LAMOTRIGINE	LAMOTRIGINE	100MG	TABLET, ORALLY DISINTEGRATING;ORAL	None (Tentative Approval)	No	None
LAMOTRIGINE	LAMOTRIGINE	200MG	TABLET, ORALLY DISINTEGRATING;ORAL	None (Tentative Approval)	No	None

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