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## Drug Details

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

- **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:**

<a href="#">Drug Name</a>	<a href="#">Active Ingredients</a>	<a href="#">Strength</a>	<a href="#">Dosage Form/Route</a>	<a href="#">Marketing Status</a>	<a href="#">RLDTE Code</a>
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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