U.S. Food and Drug Administration
Department of Health and Human Services
CENTER FOR DRUG EVALUATION AND RESEARCH

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FDA Approved Drug Products

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Drug Details

Drug Name(s)	**LAMICTAL (Brand Name Drug)**
FDA Application No.	**(NDA) 020241**
Active Ingredient(s)	**LAMOTRIGINE**
Company	**GLAXOSMITHKLINE**
Original Approval or Tentative Approval Date	**December 27, 1994**
Chemical Type	**1 New molecular entity (NME)**
Review Classification	**P Priority review drug**

- Therapeutic Equivalents
- Approval History, Letters, Reviews, and Related Documents
- Dear Health Professional Letter
- Label Information
- Dear Health Professional Letter
- Other Important Information from FDA

Products on Application (NDA) #020241

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
LAMICTAL	LAMOTRIGINE	100MG	TABLET; ORAL	Prescription	No	AB
LAMICTAL	LAMOTRIGINE	150MG	TABLET; ORAL	Prescription	No	AB

LAMICTAL	LAMOTRIGINE	200MG	TABLET; ORAL	Prescription	No	AB
LAMICTAL	LAMOTRIGINE	250MG	TABLET; ORAL	Discontinued	No	None
LAMICTAL	LAMOTRIGINE	25MG	TABLET; ORAL	Prescription	Yes	AB
LAMICTAL	LAMOTRIGINE	50MG	TABLET; ORAL	Discontinued	No	None

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