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## Drug Details

<b>Drug Name(s)</b>	<b>GABAPENTIN</b>
<b>FDA Application No.</b>	<b>(ANDA) 205101</b>
<b>Active Ingredient(s)</b>	<b>GABAPENTIN</b>
<b>Company</b>	<b>SCIEGEN PHARMS INC</b>
<b>Original Approval or Tentative Approval Date</b>	<b>February 4, 2016</b>

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

### Products on Application (ANDA) #205101

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="#">RLD</a>	<a href="#">TE Code</a>
GABAPENTIN	GABAPENTIN	600MG	TABLET;ORAL	Prescription	No	AB
GABAPENTIN	GABAPENTIN	800MG	TABLET;ORAL	Prescription	No	AB

[Back to Top](#) | [Back to Previous Page](#) | [Back to Drugs@FDA Home](#)

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