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## Drug Details

<b>Drug Name(s)</b>	<b>GUAIFENESIN</b>
<b>FDA Application No.</b>	<b>(ANDA) 091009</b>
<b>Active Ingredient(s)</b>	<b>GUAIFENESIN</b>
<b>Company</b>	<b>ACTAVIS LABS FL INC</b>
<b>Original Approval or Tentative Approval Date</b>	<b>September 3, 2015</b>

- [Other OTC Drugs with the same Active Ingredient, Strength and Dosage Form/Route](#)
- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

### Products on Application (ANDA) #091009

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 TE Code</a>
GUAIFENESIN	GUAIFENESIN	1.2GM	TABLET, EXTENDED RELEASE;ORAL	Over-the-counter	No None

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