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## Drug Details

<b>Drug Name(s)</b>	<b>LANSOPRAZOLE</b>
<b>FDA Application No.</b>	<b>(NDA) 208025</b>
<b>Active Ingredient(s)</b>	<b>LANSOPRAZOLE</b>
<b>Company</b>	<b>DEXCEL PHARMA</b>
<b>Original Approval or Tentative Approval Date</b>	<b>June 7, 2016</b>
<b>Chemical Type</b>	<b>5 New formulation or new manufacturer</b>

- **There are no Therapeutic Equivalents**
- **Labels are not available**
- [Approval History, Letters, Reviews, and Related Documents](#)

### Products on Application (NDA) #208025

**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>
LANSOPRAZOLE	LANSOPRAZOLE	15MG	TABLET, DELAYED RELEASE, ORALLY DISINTEGRATING;ORAL	Prescription	TBD	 None
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