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## Drug Details

<b>Drug Name(s)</b>	<b>SUMATRIPTAN</b>
<b>FDA Application No.</b>	<b>(ANDA) 204841</b>
<b>Active Ingredient(s)</b>	<b>SUMATRIPTAN</b>
<b>Company</b>	<b>LANNETT</b>
<b>Original Approval or Tentative Approval Date</b>	<b>February 19, 2016</b>

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

### Products on Application (ANDA) #204841

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>
SUMATRIPTAN	SUMATRIPTAN	5MG/SPRAY	SPRAY;NASAL	Prescription	No	AB
SUMATRIPTAN	SUMATRIPTAN	20MG/SPRAY	SPRAY;NASAL	Prescription	No	AB

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