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## Drug Details

<b>Drug Name(s)</b>	<b>POLYMYXIN B SULFATE</b>
<b>FDA Application No.</b>	<b>(ANDA) 206589</b>
<b>Active Ingredient(s)</b>	<b>POLYMYXIN B SULFATE</b>
<b>Company</b>	<b>AUROBINDO PHARMA LTD</b>
<b>Original Approval or Tentative Approval Date</b>	<b>April 5, 2016</b>

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

### Products on Application (ANDA) #206589

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="#">RLDTE Code</a>
POLYMYXIN B SULFATE	POLYMYXIN B SULFATE	EQ 500,000 UNITS BASE/VIAL	INJECTABLE;INJECTION	Prescription No	AP

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