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## Drug Details

<b>Drug Name(s)</b>	<b>CEFAZOLIN</b>
<b>FDA Application No.</b>	<b>(NDA) 207131</b>
<b>Active Ingredient(s)</b>	<b>CEFAZOLIN</b>
<b>Company</b>	<b>CELERITY PHARMACEUTICALS LLC</b>
<b>Original Approval or Tentative Approval Date</b>	<b>August 7, 2015</b>
<b>Chemical Type</b>	<b>5 New formulation or new manufacturer</b>
<b>Review Classification</b>	<b>S Standard review drug</b>

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

### Products on Application (NDA) #207131

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>
CEFAZOLIN	CEFAZOLIN	2 G per 100 ML	SOLUTION;INJECTION	Prescription	TBD  <sup>11</sup>	None

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