Drugs@FDA
FDA Approved Drug Products

Drug Details

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

- **There are no Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (NDA) #207131
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
CEFAZOLIN	CEFAZOLIN	2 G per 100 ML	SOLUTION;INJECTION	Prescription	TBD [11]	None

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