

CEFAZOLIN SODIUM - cefazolin sodium injection, solution
Cantrell Drug Company

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Cefazolin Sodium 2 g Added to 0.9% Sodium Chloride 100 mL Bag

ceFAZolin Sodium

2 g

Added to
0.9% Sodium Chloride 100 mL* Bag

Volume: 100 mL*

Rx Only

***Volume Excludes Manufacturer Overfill**

Store Refrigerated. Protect From Light.

Single-Dose Bag. For IV Use Only.

NDC: 52533-014-42



00001

(01) 0 0525330 14420 5

Outsourced & Compounded by:



CANTRELL DRUG COMPANY

7321 Cantrell Road Little Rock, AR 72207
(877) 666-5222 www.cantrelldrug.com

Lot: xxxxxx

BUD:

CEFAZOLIN SODIUM

cefazolin sodium injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-014
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Cefazolin Sodium (UNII: P380M0454Z) (Cefazolin - UNII:IHS69L0Y4T)	Cefazolin	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Sodium Chloride (UNII: 451W47IQ8X)	0.9 g in 100 mL
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-014-42	100 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/01/2011	

Labeler - Cantrell Drug Company (035545763)

Revised: 4/2014

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