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 3 g Added to 0.9% Sodium Chloride 100 mL Bag

ceFAZolin Sodium 3 g

added to 0.9% Sodium Chloride 100 mL* Bag

BUD:

LOT: xxxxx

Compounded Date: 09/99

Volume: 100 mL*

*Volume & Concentration Excludes Manufacturer Overfill Store Refrigerated. Protect from Light. Single-Dose Bag. Hospital/Office Use Only. Injection Solution For IV Use Only.

NDC: 52533-113-42

Rx Only



*Each mL Contains: Cefazolin Sodium 30 mg, Sodium Chloride 9 mg.

Outsourced Compounded Drug. Not for Resale

Cantrell Drug Co. 7321 Cantrell Road Little Rock, AR 72207

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CEFAZOLIN SODIUM

cefazolin sodium injection, solution

Product Type HUMAN PRESCRIPTION DRUG

Item Code (Source)

NDC:52533-113

Route of Administration

INTRAVENOUS

Active Ingredient/Active Moiety

	Ingredient Name	Basis of Strength	Strength
ı	Cefazolin Sodium (UNII: P380M0454Z) (Cefazolin - UNII:IHS69L0Y4T)	Cefazolin	30 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL			
Water (UNII: 059QF0KO0R)				

1	Packaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-113-42	100 mL in 1 BAG		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved drug other		12/05/2013			

Labeler - Cantrell Drug Company (035545763)

Revised: 6/2014 Cantrell Drug Company