## HEPARIN SODIUM - heparin 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.

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Heparin Sodium 5,000 USP Units Added to 0.9% Sodium Chloride 500 mL Bag



Added to 0.9% Sodium Chloride 500 mL\* Bag

(10 USP units/mL)\*Volume & Concentration Exclude Manufacturer Overfill Store at Room Temperature. Single-Dose Bag. Hospital/Office Use Only. Injection Solution For IV Use.







Each mL Contains: Heparin Sodium 10 USP units, Sodium Chloride 9.086 mg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

Outsourced Compounded Drug. Not for Resale.

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LOT: XXXXX BUD: CMPD Date: 03/13



## HEPARIN SODIUM

heparin sodium injection, solution

## Product Information Product Type HUMAN PRESCRIPTION DRUG Item Code (Source) NDC:52533-098 Route of Administration INTRAVENOUS

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	10 [USP'U] in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL	
BENZYL ALCOHOL (UNII: LKG8494WBH)	0.001 mL in 1 mL	
Water (UNII: 059QF0KO0R)		

Other Ingredients		
Ingredient Kind	Ingredient Name	Quantity
May contain	HYDRO CHLO RIC ACID (UNII: QTT17582CB)	
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-098-30	500 mL in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/30/2011	

## Labeler - Cantrell Drug Company (035545763)

Establishment			
Name	Address	ID/FEI	Business Operations
Cantrell Drug Company		035545763	manufacture(52533-098), human drug compounding outsourcing facility(52533-098) (No intent to compound 506E (drug shortage) drugs), (Not compounding from bulk ingredient)

Revised: 12/2014 Cantrell Drug Company