## 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 1,000 mL Bag



Added to 0.9% Sodium Chloride 1,000 mL\* Bag

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





0.9% Rx Only

Each mL Contains: Heparin Sodium 5 USP units, Sodium Chloride 9.043 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:52533-097
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	5 [USP'U] in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL		
BENZYL ALCOHOL (UNII: LKG8494WBH)	0.00005 mL in 1 mL		
Water (UNII: 059QF0KO0R)			

Other Ingredients			
Ingredient Kind	Ingredient Name	Quantity	
May contain	HYDRO CHLO RIC ACID (UNII: QTT17582CB)		
May contain	SO DIUM HYDRO XIDE (UNII: 55X04QC32I)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:52533-097-24	1000 mL in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/09/2012	

## Labeler - Cantrell Drug Company (035545763)

Revised: 1/2015 Cantrell Drug Company