

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.

Heparin Sodium 4,000 USP Units Added to 0.45% Sodium Chloride 1,000 mL Bag

HEPARIN

Sodium

4,000

**USP
Units**

Added to 0.45% Sodium Chloride 1,000 mL* Bag

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

**HIGH
ALERT**

NDC: 52533-211-26



(01) 0 0352533 21126 3

0.45%
Rx Only

Each mL Contains: Heparin Sodium 4 USP units, Sodium Chloride 4.53 mg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

Outsourced Compounded Drug. Not for Resale.

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CANTRELL DRUG COMPANY

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LOT: xxxxxx

BUD:

CMPD Date: 03/13



HEPARIN SODIUM

heparin sodium injection, solution

Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-211	
Route of Administration	INTRAVENOUS			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)		Heparin	4 [USP'U] in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
Sodium Chloride (UNII: 451W47IQ8X)		4.5 mg in 1 mL		
BENZYL ALCOHOL (UNII: LKG8494WBH)		0.00004 mL in 1 mL		
Water (UNII: 059QF0K00R)				
Other Ingredients				
Ingredient Kind	Ingredient Name		Quantity	
May contain	HYDROCHLORIC ACID (UNII: QTT17582CB)			
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-211-26	1000 mL in 1 BAG		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		11/13/2012		

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

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