

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 25,000 USP Units Added to 0.45% Sodium Chloride 500 mL Bag

HEPARIN

Sodium

25,000

USP Units

Added to 0.45% Sodium Chloride 500 mL* Bag

(50 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: 52533-216-40



(01) 0 0352533 21640 4

0.45%
Rx Only

Each mL Contains: Heparin Sodium 50 USP units, Sodium Chloride 4.57 mg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

Outsourced Compounded Drug. Not for Resale.



CANTRELL DRUG COMPANY

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LOT: xxxxxx

BUD:

CMPD Date: 03/13

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HEPARIN SODIUM

heparin sodium injection, solution

Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-216	
Route of Administration	INTRAVENOUS			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)		Heparin	50 [USP'U] in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
Sodium Chloride (UNII: 451W47IQ8X)		4.5 mg in 1 mL		
BENZYL ALCOHOL (UNII: LKG8494WBH)		0.0001 mL in 1 mL		
Water (UNII: 059QF0KO0R)				
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:52533-216-40	500 mL in 1 BAG		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			12/21/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

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