

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 1,000 USP Units Added to 0.9% Sodium Chloride 500 mL Bag

HEPARIN

Sodium

1,000

USP Units

Added to 0.9% Sodium Chloride 500 mL* Bag

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

**HIGH
ALERT**

NDC: 52533-138-30

(01) 0 0352533 13830 0


Rx Only

Each mL Contains: Heparin Sodium 2 USP units,
Sodium Chloride 9 mg. pH adj: HCl/NaOH.
Outsourced Compounded Drug. Not for Resale.

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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:52533-138
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	2 [USP'U] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	9 mg in 1 mL
Water (UNII: 059QF0K00R)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	HYDROCHLORIC ACID (UNII: QTT17582CB)	
May contain	SODIUM HYDRO XIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-138-30	500 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/29/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

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