

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 30,000 USP Units Added to 0.9% Sodium Chloride 1,000 mL Bag



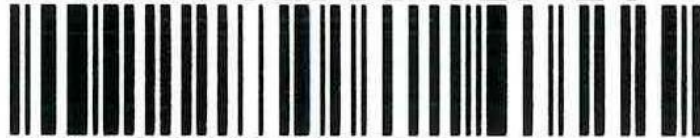
HEPARIN
Sodium
30,000 USP
Units

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

(30 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: 52533-029-24



(01) 0 0352533 02924 0

0.9%
Rx Only

Each mL Contains: Heparin Sodium 30 USP units, Sodium Chloride 9.042 mg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

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Outsourced Compounded Drug. Not for Resale.



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:52533-029
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	30 [USP'U] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL
BENZYL ALCOHOL (UNII: LKG8494WBH)	
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:52533-029-24	1000 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/22/2013	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

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