

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



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.

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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 LABEL	Item Code (Source)	NDC:52533-216
Route of Administration	INTRAVENOUS	DEA Schedule	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Heparin Sodium (Heparin)	Heparin	50 [USP'U] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Sodium Chloride	4.5 mg in 1 mL
BENZYL ALCOHOL	0.0001 mL in 1 mL
Water	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	HYDROCHLORIC ACID	
May contain	SODIUM HYDROXIDE	

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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