

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 2,500 USP Units Added to 4.5% Sodium Chloride 500 mL Bag

HEPARIN

Sodium

2,500

USP Units

Added to 0.45% Sodium Chloride 500 mL* Bag

(5 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: 52533-096-40



(01) 0 0352533 09640 2

0.45%
Rx Only

Each mL Contains: Heparin Sodium 5 USP units, Sodium Chloride 4.54 mg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

Outsourced Compounded Drug. Not for Resale.

00004



CANTRELL DRUG COMPANY

7321 Cantrell Road Little Rock, AR 72207
(877) 666-5222 www.cantrelldrug.com

LOT: xxxxxx

BUD:

CPD Date: 03/13



HEPARIN SODIUM

heparin sodium injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG LABEL	Item Code (Source)	NDC:52533-096
Route of Administration	INTRAVENOUS	DEA Schedule	

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Heparin Sodium (Heparin)	Heparin	5 [USP'U] in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM CHLORIDE	4.5 mg in 1 mL
BENZYL ALCOHOL	0.00005 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:52533-096-40	500 mL in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/18/2011	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

Cantrell Drug Company