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 20,000 USP Units Added to 5% Dextrose 500 mL Bag

HEPARIN Sodium 20,000 USP Units

Added to 5% Dextrose 500 mL Bag

(40 USP units/mL) *Volume & Concentration Exclude Manufacturer Overfill

Store at Room Temperature. Single-Dose Bag. Hospital/Office Use Only. Injection Solution For IV Use.



NDC: 52533-215-32

D5

Each mL Contains: Heparin Sodium 40 USP units, Dextrose 50 mg, NaCl 56 mcg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

Outsourced Compounded Drug. Not for Resale.

00002



LOT: XXXXXX BUD:

CMPD Date: 03/13



HEPARIN SODIUM

heparin sodium injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-215
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	40 [USP'U] in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
DEXTROSE (UNII: IY9 XDZ35W2)	50 mg in 1 mL		
BENZYL ALCOHOL (UNII: LKG8494WBH)	0.00008 mL in 1 mL		
Water (UNII: 059QF0KO0R)			

Other Ingredients			
Ingredient Kind	Ingredient Name	Quantity	
May contain	HYDRO CHLO RIC ACID (UNII: QTT17582CB)		
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-215-32	500 mL in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/08/2014	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014 Cantrell Drug Company