

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 0.5 USP Units/mL in 0.45% Sodium Chloride 2 mL Syringe

Label

HEPARIN SODIUM			
heparin sodium injection, solution			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-148
Route of Administration	INTRAVENOUS		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	0.5 [USP'U] in 1 mL	
Inactive Ingredients			
Ingredient Name	Strength		
Sodium Chloride (UNII: 451W47IQ8X)	4.5 mg in 1 mL		

WATER (UNII: 059QF0K00R)

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:52533-148-16	2 mL in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/22/2012	

Labeler - Cantrell Drug Company (035545763)

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