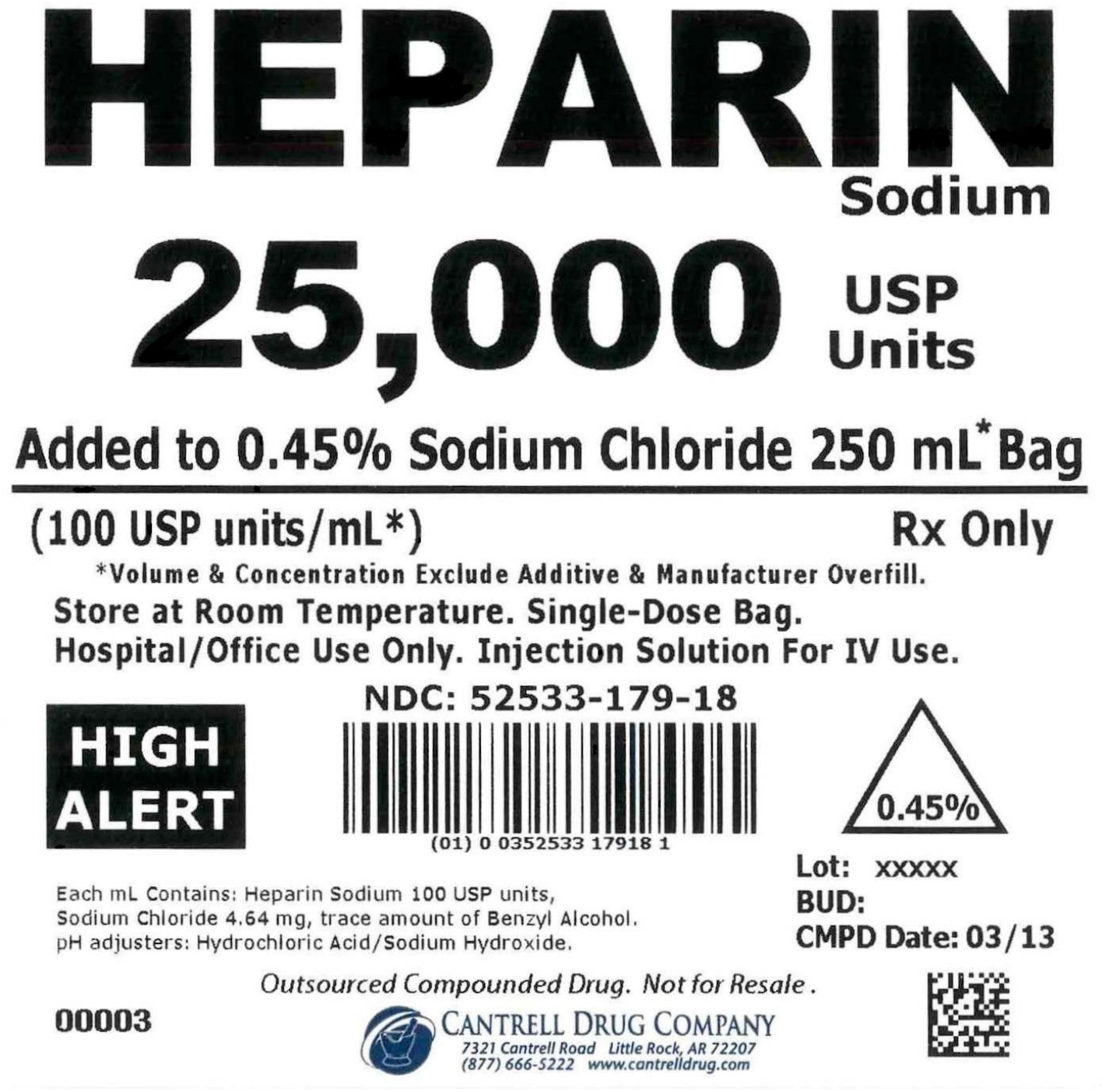
HEPARIN SODIUM - heparin sodium injection, solution
Cantrell Drug Company

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 250 mL Bag



• WARNINGS AND PRECAUTIONS

Outsourced Compounded Drug. Not for Resale. Hospital/Office Use Only.

• ADVERSE EVENTS

To facilitate Adverse Event Reporting: www.fda.gov/medwatch or 1-800-FDA-1088.

• HOW SUPPLIED

Contains 25,000 USP Units of Heparin Sodium in 0.45% Sodium Chloride in a 250 mL Single-Dose Bag. Volume & Concentration exlude additive & manufacturer overfill.

This product is Sterile, Nonpyrogenic, and Latex Free.

• INGREDIENTS

Each 1 mL contains contains Heparin Sodium 100 USP Units, Sodium Chloride 4.5 mg, Benzyl Alcohol 0.0002 mL. May contain Hydrochloric Acid and/or Sodium Hydroxide for pH adjustment.

• STORAGE AND HANDLING

Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature].

• DOSAGE AND ADMINISTRATION.

FOR INTRAVENOUS USE.

Rx Only

Rev. 05/15

CANTRELL DRUG COMPANY

LITTLE ROCK, AR 72207

HEPARIN SODIUM

heparin sodium injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:52533-179
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	100 [USP'U] in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Sodium Chloride (UNII: 451W47IQ8X)	4.5 mg in 1 mL
BENZYL ALCOHOL (UNII: LKG8494WBH)	0.0002 mL in 1 mL
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:52533-179-18	250 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/15/2015	

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2015 Cantrell Drug Company