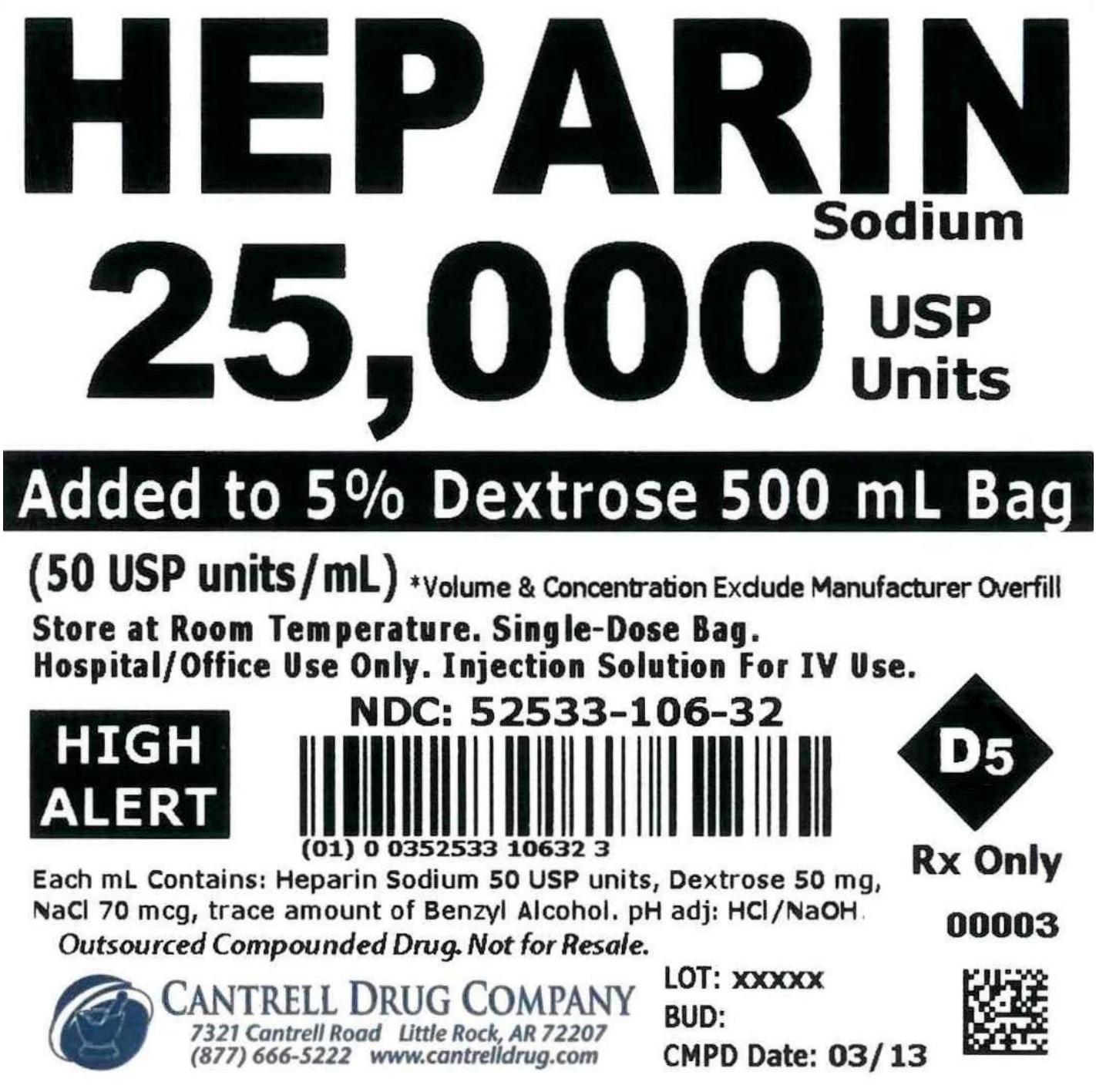
HEPARIN SODIUM - heparin sodium injection, solution
Cantrell Drug Company

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 5% Dextrose 500 mL Bag



HEPARIN SODIUM
heparin sodium injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:52533-106
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Heparin Sodium (UNII: ZZ45AB24CA) (Heparin - UNII:T2410KM04A)	Heparin	50 [USP'U] in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
DEXTROSE (UNII: IY9XDZ35W2)	50 mg in 1 mL
BENZYL ALCOHOL (UNII: LKG8494WBH)	0.0001 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:52533-106-32	500 mL in 1 BAG		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
unapproved drug other		08/09/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014 Cantrell Drug Company