

**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.*

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**Heparin Sodium 25,000 USP Units Added to 5% Dextrose 500 mL Bag**

# **HEPARIN**

**Sodium**

# **25,000**

**USP Units**

**Added to 5% Dextrose 500 mL Bag**

**(50 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: 52533-106-32**



(01) 0 0352533 10632 3



**Rx Only**

Each mL Contains: Heparin Sodium 50 USP units, Dextrose 50 mg,  
NaCl 70 mcg, trace amount of Benzyl Alcohol. pH adj: HCl/NaOH.

**00003**

*Outsourced Compounded Drug. Not for Resale.*



**CANTRELL DRUG COMPANY**

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LOT: xxxxxx

BUD:

CMPD Date: 03/13



**HEPARIN SODIUM**

heparin sodium injection, solution

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

| Active Ingredient/Active Moiety |                   |                    |
|---------------------------------|-------------------|--------------------|
| Ingredient Name                 | Basis of Strength | Strength           |
| Heparin Sodium (Heparin)        | Heparin           | 50 [USP'U] in 1 mL |

| Inactive Ingredients |                   |
|----------------------|-------------------|
| Ingredient Name      | Strength          |
| DEXTROSE             | 50 mg in 1 mL     |
| BENZYL ALCOHOL       | 0.0001 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:52533-106-32 | 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