PHENYLEPHRINE HCL - phenylephrine hcl 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.

Phenylephrine HCl 40 mg Added to 0.9% Sodium Chloride 250 mL Bag

Phenylephrine 40 mg HCl

Added to
0.9% Sodium Chloride
250 mL* Bag

(160 mcg/mL*)

*Volume & Concentration Exclude Manufacturer Overfill

Store at Room Temperature. Protect from Light.
Contains No Antimicrobial Preservative.
Single-Dose Bag. Injection Solution for IV Use.

NDC: 52533-110-18

(01) 0 0352533 11018 4

Rx Only

Each mL Contains: Phenylephrine HCl 160 mcg, Sodium Chloride 9.056 mg, Sodium Citrate (Dihydrate) 64 mcg, Sodium Metabisulfite 32 mcg, Citric Acid 14.6 mcg. pH adj: Citric Acid/Sodium Hydroxide.

Hospital/Office Use Only

Outsourced Compounded Drug. Not for Resale

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LOT: xxxxx
BUD:
CMPD Date: 03/13

PHENYLEPHRINE HCL
phenylephrine hcl injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:52533-110
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	160 ug in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Sodium Chloride (UNII: 451W47IQ8X)	9.056 mg in 1 mL
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	64 ug in 1 mL
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	32 ug in 1 mL
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	14.6 ug in 1 mL
WATER (UNII: 059QF0KO0R)	

Other Ingredients		
Ingredient Kind	**Ingredient Name**	**Quantity**
May contain	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52533-110-18	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		06/30/2011	

Labeler - Cantrell Drug Company (035545763)

Revised: 8/2015 Cantrell Drug Company