

LABETALOL HCL - labetalol 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.

Labetalol HCl 5 mg/mL 4 mL Syringe

Label

Labetalol HCl **20 mg/4 mL**
(5 mg/mL)
Injection Solution

Store at Room Temperature. Protect from Light. Preserved. Single-Dose Syringe. For IV Use. Hospital/Office Use Only. **LOT: xxxxxx**
BUD: CMPD Date: 06/12

NDC: 52533-034-20 Outsourced Compounded Drug: Not for Resale

Rx Only

(01) 0 0352533 03420 6

Cantrell Drug Co. 7321 Cantrell Rd. Little Rock, AR
 877-666-5222 www.cantrelldrug.com

Labetalol 5 mg/mL

Each mL: Labetalol HCl 5mg, Dextrose 45mg, EDTA 0.1mg, Methylparaben 0.8mg, Propylparaben 0.1mg, pH adj: Citric Acid/NaOH.

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LABETALOL HCL

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Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-034
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
LABETALOL HYDROCHLORIDE (UNII: 1GEV3BAW9J) (LABETALOL - UNII:R5H8897N95)		LABETALOL HYDROCHLORIDE	5 mg in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
Water (UNII: 059QF0K00R)				
Anhydrous Dextrose (UNII: 5SL0G7R0OK)		45 mg in 1 mL		
Edetate Disodium (UNII: 7FLD91C86K)		0.1 mg in 1 mL		
Methylparaben (UNII: A2I8C7HI9T)		0.8 mg in 1 mL		
Propylparaben (UNII: Z8IX2SC1OH)		0.1 mg in 1 mL		
Citric Acid Monohydrate (UNII: 2968PHW8QP)				
Sodium Hydroxide (UNII: 55X04QC32I)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-034-20	4 mL in 1 SYRINGE, PLASTIC		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		02/23/2012		

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014

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