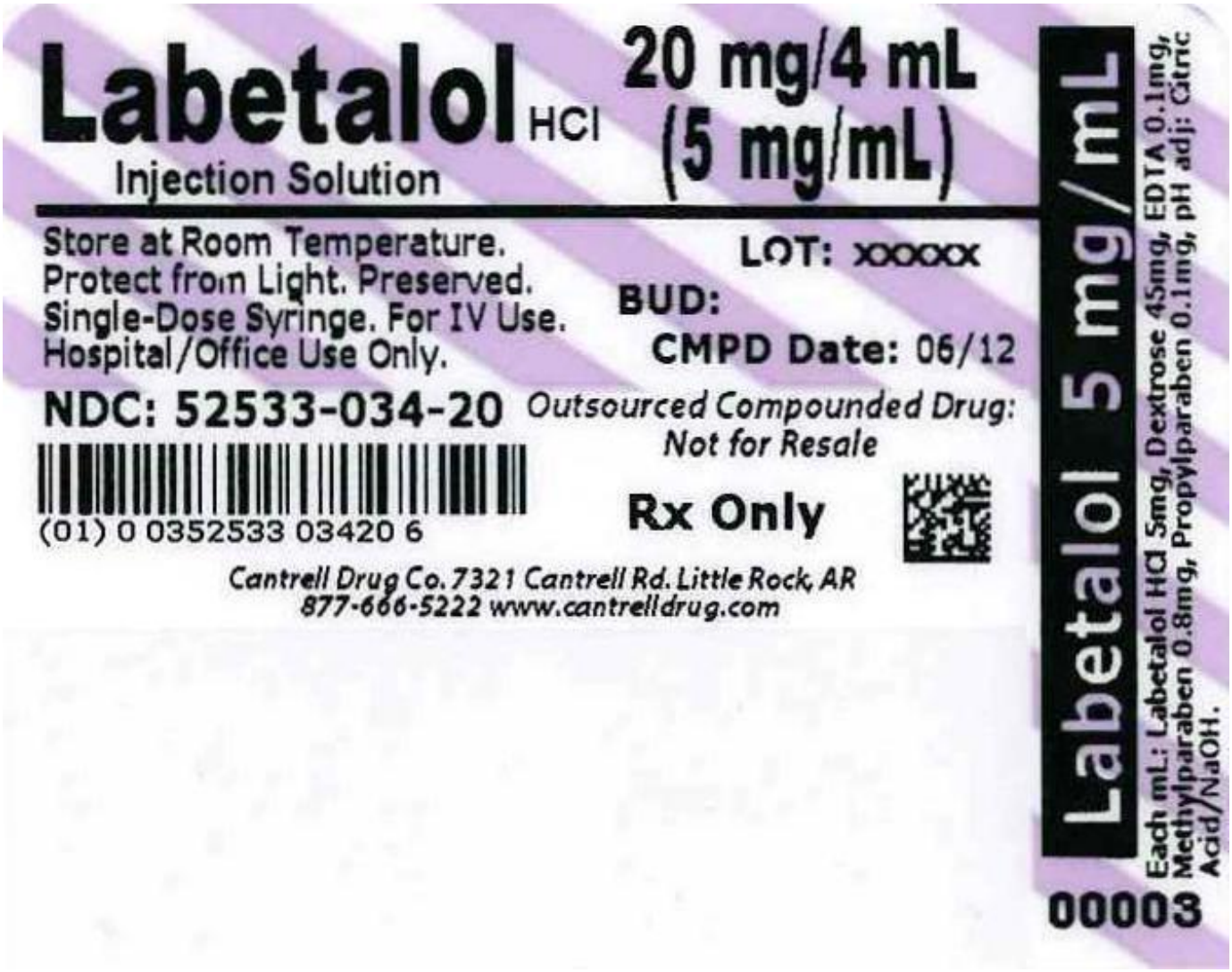
LABETALOL HCL - labetalol hcl injection, solution
Cantrell Drug Company

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

labetalol hcl injection, solution

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: 059QF0KO0R)	
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 Cantrell Drug Company