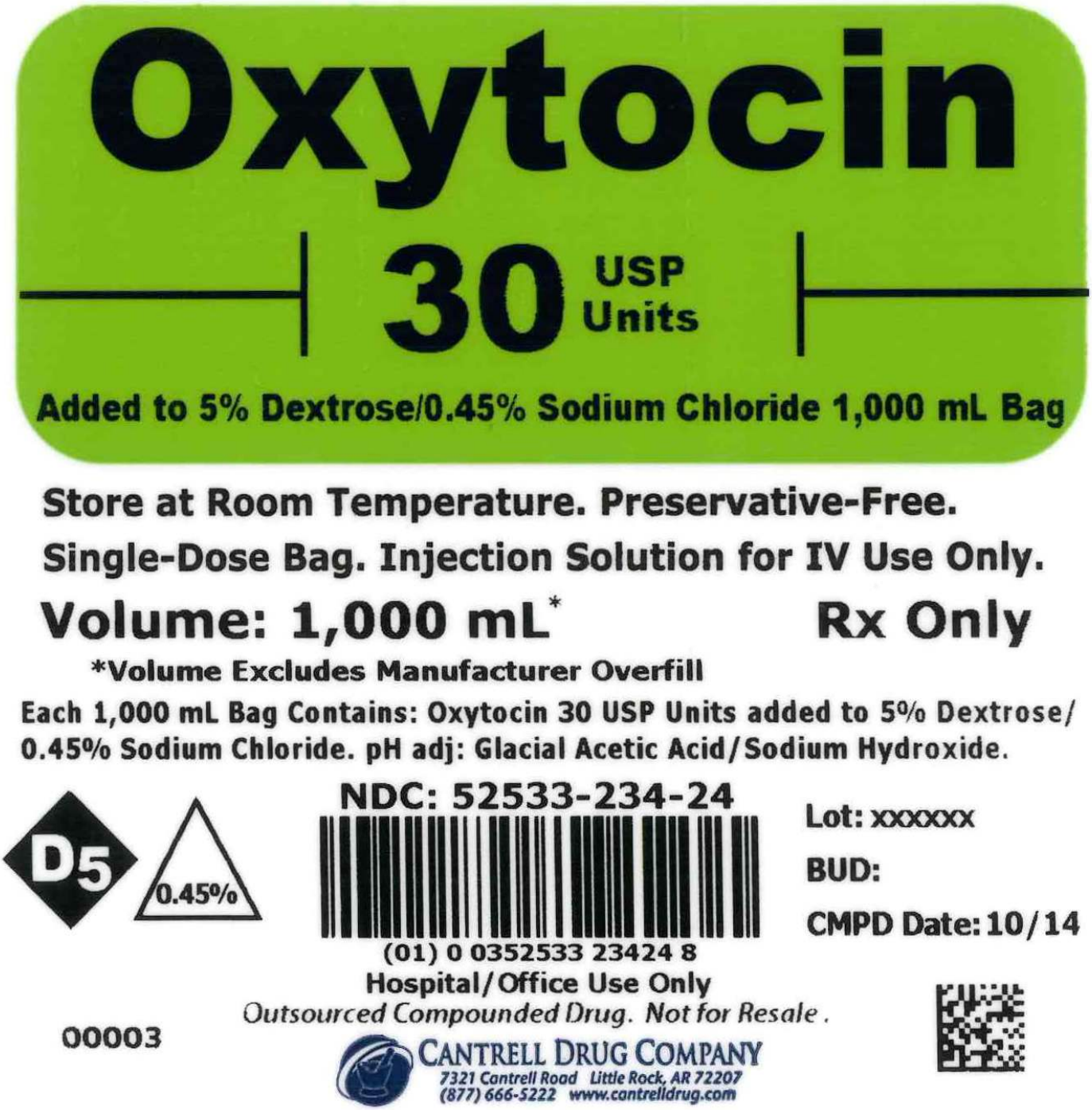
OXYTOCIN- oxytocin injection, solution
Cantrell Drug Company

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.

Oxytocin 30 USP Units Added to 5% Dextrose/0.45% Sodium Chloride 1,000 mL Bag



OXYTOCIN
oxytocin injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:52533-234
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OXYTOCIN (UNII: 1JQS135EYN) (OXYTOCIN - UNII:1JQS135EYN)	OXYTOCIN	30 [USP'U] in 1000 mL

Inactive Ingredients	
Ingredient Name	**Strength**
DEXTROSE MONOHYDRATE (UNII: LX22YL083G)	50 g in 1000 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X)	4.5 g in 1000 mL
WATER (UNII: 059QF0KO0R)	

Other Ingredients		
Ingredient Kind	**Ingredient Name**	**Quantity**
May contain	**ACETIC ACID** (UNII: Q40Q9N063P)	
May contain	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52533-234-24	1000 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		03/18/2016	

Labeler - Cantrell Drug Company (035545763)

Revised: 3/2016 Cantrell Drug Company