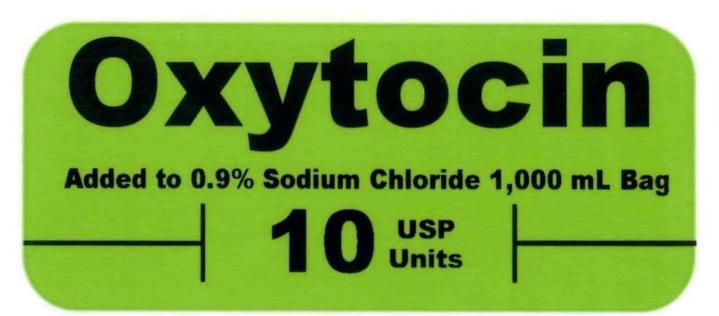
OXYTOCIN - oxytocin injection, solution Cantrell Drug Company

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 10 USP Units Added to 0.9% Sodium Chloride 1,000 mL Bag



Store at Room Temperature. Preservative Free. Single-Dose Bag. Injection Solution for IV Use Only.

Volume: 1,000 mL*

Rx Only

*Volume Excludes Manufacturer Overfill

Each 1,000 mL Bag Contains: Oxytocin 10 USP Units added to 0.9% Sodium Chloride. pH adj: Glacial Acetic Acid/Sodium Hydroxide.



NDC: 52533-048-24

Lot: xxxxx

BUD:

CMPD Date: 03/13

00002

Hospital/Office Use Only Outsourced Compounded Drug. Not for Resale.





OXYTOCIN

oxytocin injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-048
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OXYTOCIN (UNII: 1JQS135EYN) (OXYTOCIN - UNII:1JQS135EYN)	OXYTOCIN	10 [USP'U] in 1000 mL	

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM CHLORIDE (UNII: 451W47IQ8X)	9 g in 1000 mL		
WATER (UNII: 059QF0KO0R)			

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

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-048-24	1000 mL in 1 BAG		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/30/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014 Cantrell Drug Company