

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 20 USP Units Added to Lactated Ringer's 1,000 mL Bag

Oxytocin

Added to Lactated Ringer's 1,000 mL Bag

20 USP
Units

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 20 USP Units added to Lactated Ringer's. pH adj: Glacial Acetic Acid/Sodium Hydroxide.

NDC: 52533-051-24



(01) 0 0352533 05124 1

Hospital/Office Use Only

Outsourced Compounded Drug. Not for Resale.

Lot: xxxxx

BUD:

CMPD Date: 03/13

LR

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CANTRELL DRUG COMPANY

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OXYTOCIN

oxytocin injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-051
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYTOCIN (UNII: 1JQS135EYN) (OXYTOCIN - UNII:1JQS135EYN)	OXYTOCIN	2 [USP'U] in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	600 mg in 100 mL
SODIUM LACTATE (UNII: TU7HW0W0QT)	310 mg in 100 mL
POTASSIUM CHLORIDE (UNII: 660YQ98110)	30 mg in 100 mL
CALCIUM CHLORIDE (UNII: M410D6VV5M)	20 mg in 100 mL

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:52533-051-24	1000 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/20/2011	

Labeler - Cantrell Drug Company (035545763)

Revised: 1/2015

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