

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 500 mL Bag

Oxytocin

Added to 0.9% Sodium Chloride 500 mL Bag

10 USP
Units

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

Volume: 500 mL*

Rx Only

***Volume Excludes Manufacturer Overfill**

Each 500 mL Bag Contains: Oxytocin 10 USP Units added to 0.9% Sodium Chloride. pH adj: Glacial Acetic Acid/Sodium Hydroxide.



00003

NDC: 52533-050-30



(01) 0 0352533 05030 5

Hospital/Office Use Only

Outsourced Compounded Drug. Not for Resale.



CANTRELL DRUG COMPANY

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Lot: xxxxx

BUD:

CMPD Date: 03/13



OXYTOCIN

oxytocin injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG LABEL	Item Code (Source)	NDC:52533-050
Route of Administration	INTRAVENOUS	DEA Schedule	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYTOCIN (OXYTOCIN)	OXYTOCIN	2 [USP'U] in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Sodium Chloride	9 g in 100 mL
Water	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	ACETIC ACID	
May contain	SODIUM HYDROXIDE	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-050-30	500 mL in 1 BAG		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/09/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 12/2014

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