

**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 15 USP Units Added to 0.9% Sodium Chloride 250 mL Bag

# Oxytocin

**Added to 0.9% Sodium Chloride 250 mL Bag**

**15** **USP**  
**Units**

**Store at Room Temperature. Preservative Free.**

**Single-Dose Bag. Injection Solution for IV Use Only.**

**Volume: 250 mL\***

**Rx Only**

**\*Volume Excludes Manufacturer Overfill**

**Each 250 mL Bag Contains: Oxytocin 15 USP Units added to 0.9% Sodium Chloride. pH adj: Glacial Acetic Acid/Sodium Hydroxide.**



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**NDC: 52533-056-18**



(01) 0 0352533 05618 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**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG LABEL	<b>Item Code (Source)</b>	NDC:52533-056
<b>Route of Administration</b>	INTRAVENOUS	<b>DEA Schedule</b>	

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
OXYTOCIN (OXYTOCIN)	OXYTOCIN	15 [USP'U] in 250 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
Sodium Chloride	2.25 g in 250 mL
Water	

**Other Ingredients**

<b>Ingredient Kind</b>	<b>Ingredient Name</b>	<b>Quantity</b>
May contain	ACETIC ACID	
May contain	SODIUM HYDRO XIDE	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:52533-056-18	250 mL in 1 BAG		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
unapproved drug other		04/25/2013	

**Labeler** - Cantrell Drug Company (035545763)

Revised: 12/2014

Cantrell Drug Company