MIDAZOLAM HCL- midazolam hcl 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.

Midazolam HCl 1 mg/mL in 0.9% Sodium Chloride 50 mL Bag

Midazolam HCI 50 mg/50 mL* in 0.9% Sodium Chloride

LOT: xxxxx

BUD: 09/09/9999

Compounded Date: 09/99

Volume: 50 mL*

*Prefilled 25 mL Bag with 50 mL Total Volume.

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

Rx Only



Hospital/Office Use Only



Each mL: Midazolam 1 mg (as HCl), Sodium Chloride 8.8 mg. pH adj: Hydrochloric Acid/Sodium Hydroxide.

Outsourced Compounded Drug. Not for Resale

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MIDAZOLAM HCL

midazolam hcl injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-001
Route of Administration	INTRAVENOUS	DEA Schedule	CIV

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MIDAZOLAM HYDRO CHLO RIDE (UNII: W7TTW573JJ) (MIDAZOLAM - UNII:R60L0SM5BC)	MIDAZOLAM	1 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8.8 mg in 1 mL		
WATER (UNII: 059QF0KO0R)			

Other Ingredients	ther Ingredients		
Ingredient Kind	Ingredient Name	Quantity	
May contain	HYDRO CHLO RIC ACID (UNII: QTT17582CB)		
May contain	SO DIUM HYDRO XIDE (UNII: 55X04QC32I)		

Packaging				
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:52533-001-77	50 mL in 1 BAG; Type 0: Not a Combination Product	06/28/2013	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/28/2013	

Labeler - Cantrell Drug Company (035545763)

Revised: 4/2016 Cantrell Drug Company