

MIDAZOLAM HCL- midazolam hydrochloride 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_{HCl} 50 mg/50 mL

in 0.9% Sodium Chloride (1 mg/mL)

LOT: xxxxx

BUD: 09/02/2014

Compounded Date: 09/99

Volume: 50 mL



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

NDC: 52533-001-31

Hospital/Office
Use Only

Rx Only



(01) 0 0352533 00131 4



Each mL: Midazolam HCl 1 mg, NaCl 9 mg, EDTA 0.02 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 LABEL	Item Code (Source)	NDC:52533-001
Route of Administration	INTRAVENOUS	DEA Schedule	CIV

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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MIDAZOLAM HYDROCHLORIDE (MIDAZOLAM)		MIDAZOLAM	1 mg in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
SODIUM CHLORIDE		9 mg in 1 mL		
EDETATE DISODIUM		0.02 mg in 1 mL		
WATER				
Other Ingredients				
Ingredient Kind	Ingredient Name		Quantity	
May contain	HYDROCHLORIC ACID			
May contain	SODIUM HYDROXIDE			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-001-31	50 mL in 1 BAG		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		04/25/2013		

Labeler - Cantrell Drug Company (035545763)

Revised: 1/2015

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