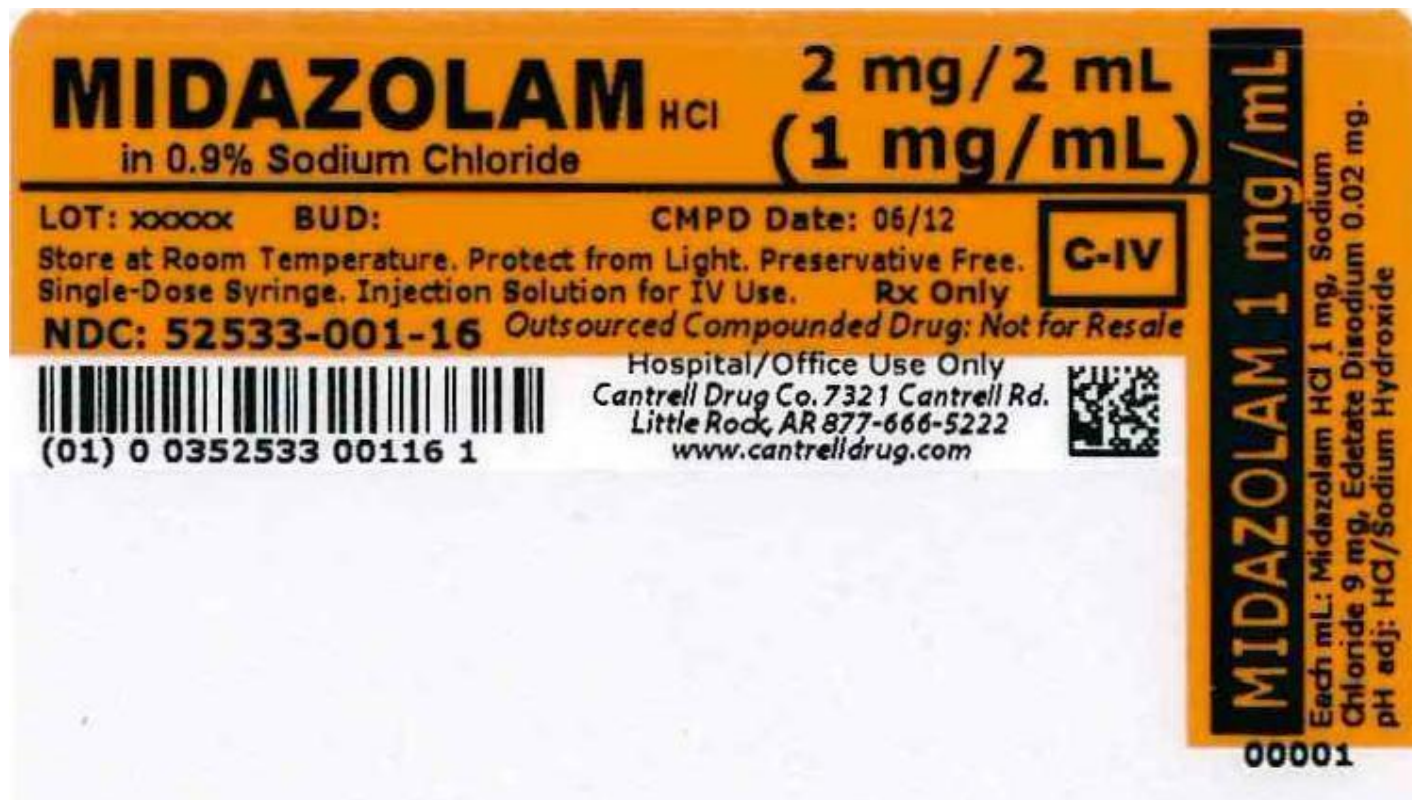


MIDAZOLAM HCL - midazolam hcl injection, solution
Cantrell Drug Company

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 2 mL Syringe

Label



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 CHLORIDE (UNII: W7TTW573JJ) (MIDAZOLAM - UNII:R60L0SM5BC)	MIDAZOLAM	1 mg in 1 mL
Inactive Ingredients			
	Ingredient Name	Strength	
	Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL	

WATER (UNII: 059QF0KO0R)				
EDETATE DISODIUM (UNII: 7FLD91C86K)		0.02 mg in 1 mL		
Other Ingredients				
Ingredient Kind	Ingredient Name		Quantity	
May contain	HYDROCHLORIC ACID (UNII: QTT17582CB)			
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-001-16	2 mL in 1 SYRINGE, PLASTIC		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		12/05/2011		

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014

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