

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 3 mL Syringe

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 HYDROCHLORIDE (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	
EDETATE DISODIUM (UNII: 7FLD91C86K)		0.02 mg in 1 mL	

WATER (UNII: 059QF0KO0R)

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:52533-001-60	3 mL in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/01/2013	

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014

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