

MORPHINE SULFATE - morphine sulfate 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.

Morphine Sulfate 2 mg/mL in 0.9% Sodium Chloride Syringe

MORPHINE SULFATE

morphine sulfate injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-161
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, INTRATHECAL, EPIDURAL	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Morphine Sulfate (UNII: X3P646A2J0) (Morphine - UNII:76I7G6D29C)	Morphine Sulfate	2 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Sodium Chloride (UNII: 451W47IQ8X)	9 mg in 1 mL
Water (UNII: 059QF0KO0R)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	SULFURIC ACID (UNII: O40 UQP6 WCF)	
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-16 1-45	1 mL in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/22/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014

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