

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 5 mg/mL in 0.9% Sodium Chloride 30 mL Monoject Syringe Barrel

MORPHINE Sulfate **150 mg/30 mL**
in 0.9% Sodium Chloride

MORPHINE 5 mg/mL

LOT: xxxxx

BUD:

Compounded Date: 01/13

(5 mg/mL)

Qty: 30 mL

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



Each mL: Morphine Sulfate 5 mg, Sodium Chloride 9 mg, pH adj: Sulfuric Acid/NaOH.

NDC: 52533-132-48 Hospital/Office Use Only



(01) 0 0352533 13248 3

Rx Only



Outsourced Compounded Drug: Not for Resale.

Cantrell Drug Co. 7321 Cantrell Road Little Rock, AR
877-666-5222 www.cantrelldrug.com

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MORPHINE SULFATE

morphine sulfate injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-132
Route of Administration	INTRAVENOUS	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MORPHINE SULFATE (UNII: X3P646A2J0) (MORPHINE - UNII:76I7G6D29C)	MORPHINE SULFATE	5 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: O40UQP6WCF)	
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-132-48	30 mL in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/25/2013	

Labeler - Cantrell Drug Company (035545763)

Revised: 4/2014

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