

HYDROMORPHONE HCL - hydromorphone 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.

Hydromorphone HCl 0.5 mg/mL in 0.9% Sodium Chloride 1 mL Syringe

HYDROMORPHONE HCl **0.5 mg / 1 mL**
(0.5 mg / mL)

in 0.9% Sodium Chloride 1 mL

LOT: 12345 BUD: 12/12/9999 CMPD Date: 04/14

Store at Room Temp. Protect from Light. Preservative Free. Isotonic. Single-Dose Syringe. Injection Solution for IV, IM, SQ Use.

C-II

NDC: 52533-005-45 Outsourced Compounded Drug: Not for Resale

Rx Only. Hospital/Office Use Only
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(01) 0 0352533 00545 9

HYDROMORPHONE 0.5 mg/mL
Each mL: Hydromorphone HCl 0.5 mg,
Sodium Chloride 9 mg; pH adj: Hydrochloric
Acid/Sodium Hydroxide.

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HYDROMORPHONE HCL

hydromorphone hcl injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-005
Route of Administration	INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROMORPHONE HYDROCHLORIDE (UNII: L960UP2KRW) (HYDROMORPHONE - UNII:Q812464R06)	HYDROMORPHONE HYDROCHLORIDE	0.5 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	HYDROCHLORIC ACID (UNII: QTT17582CB)	
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-005-45	1 mL in 1 SYRINGE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/11/2013	

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014

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