

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 10 mg/mL in 0.9% Sodium Chloride 30 mL PCA Vial, Non-MedNet**HYDROMORPHONE HCl 10 mg/mL in
0.9% Sodium Chloride 30 mL PCA Vial****(10 mg/mL) Total Dose: 300mg/30mL Qty: 30 mL****LOT: 134672 BUD: 09/04/2014 CMPD Date: 06/14***Outsourced Compounded Drug. Hospital/Office Use Only.***STORE ROOM TEMP. PROTECT FROM LIGHT. PRESERVATIVE FREE.
SINGLE-DOSE. INJECTION SOLUTION FOR SLOW IV USE.**

Each mL Contains: Hydromorphone HCl 10 mg, Sodium Chloride 9 mg, pH Adjuster: Hydrochloric Acid/Sodium Hydroxide

NDC: 52533-008-10**(01) 0 0352533 00810 8****0003****Rx Only**Cantrell Drug Co. 7321 Cantrell Rd. Little Rock, AR 72207
877-666-5222 www.cantrelldrug.com**Non-MedNet****HYDROMORPHONE HCL**

hydromorphone hcl injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-008
Route of Administration	INTRAVENOUS	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROMORPHONE HYDROCHLORIDE (UNII: L960UP2KRW) (HYDROMORPHONE - UNII:Q812464R06)	HYDROMORPHONE HYDROCHLORIDE	10 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:52533-008-10	30 mL in 1 VIAL, GLASS		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			08/13/2014	

Labeler - Cantrell Drug Company (035545763)

Revised: 8/2014

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