

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 30 mL PCA Vial

HYDROmorphone HCl 15 mg/30 mL

in 0.9% Sodium Chloride (0.5 mg/mL)

**Store at Room Temperature. Protect from Light.
Preservative Free. Single-Dose PCA Vial.
Injection Solution For Slow IV Use.**



NDC: 52533-005-05

BUD:

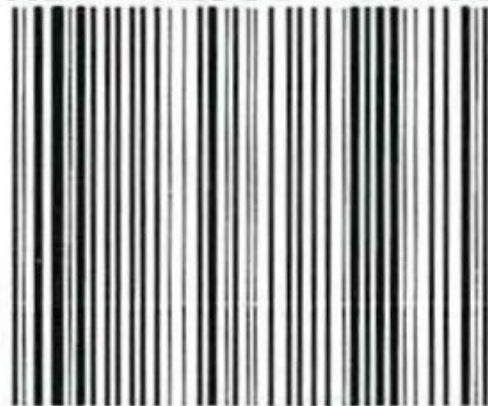
LOT: xxxxxx

CMPD Date: 03/13

Hospital/Office Use Only

Rx Only

30 mL



(01) 0 0352533 00505 3



Each mL: Hydromorphone HCl 0.5 mg, Sodium Chloride 9 mg. pH adj: HCl/NaOH.

Outsourced Compounded Drug. Not for Resale. 00001

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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	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: 059QF0K00R)				
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:52533-005-05	30 mL in 1 VIAL, GLASS		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			01/18/2012	

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014

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