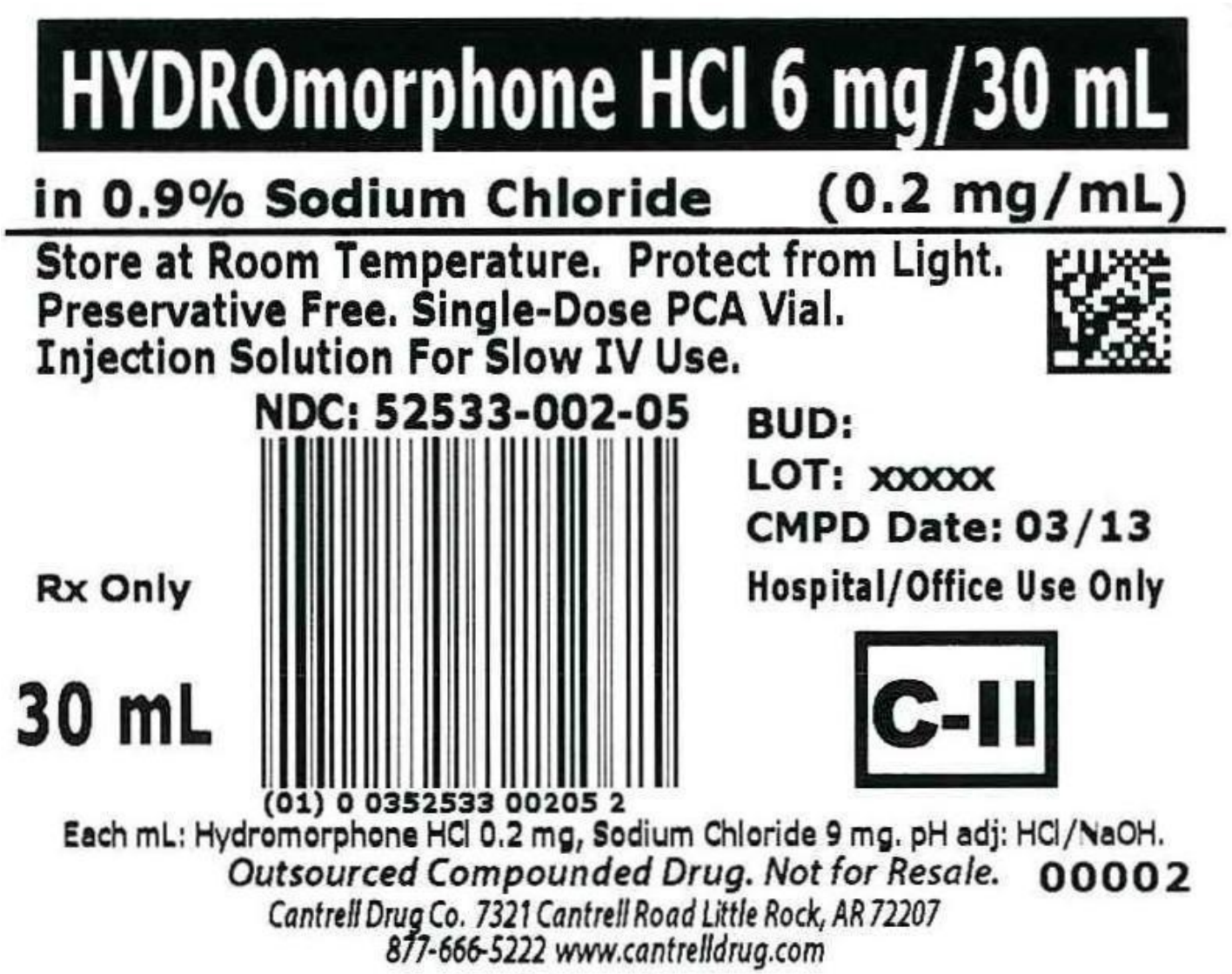
HYDROMORPHONE HCL - hydromorphone hydrochloride injection, solution
Cantrell Drug Company

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 0.2 mg/mL in 0.9% Sodium Chloride 30 mL PCA Vial



HYDROMORPHONE HCL

hydromorphone hcl injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG LABEL	**Item Code (Source)**	NDC:52533-002
Route of Administration	INTRAVENOUS	**DEA Schedule**	CII

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROMORPHONE HYDROCHLORIDE (HYDROMORPHONE)	HYDROMORPHONE HYDROCHLORIDE	0.2 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Sodium Chloride	9 mg in 1 mL
WATER	

Other Ingredients		
Ingredient Kind	**Ingredient Name**	**Quantity**
May contain	**HYDROCHLORIC ACID**	
May contain	**SODIUM HYDROXIDE**	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52533-002-05	30 mL in 1 VIAL, GLASS		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
unapproved drug other		08/23/2010	

Labeler - Cantrell Drug Company (035545763)

Revised: 5/2014 Cantrell Drug Company