

FENTANYL CITRATE - fentanyl citrate 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.

Fentanyl Citrate 20 mcg/mL in 0.9% Sodium Chloride 30 mL PCA Vial, Non-MedNet

**fentaNYL Citrate 20 mcg/mL in
 0.9% Sodium Chloride 30 mL PCA Vial**



(20 mcg/mL) Total Dose: 600mcg/30mL Qty: 30 mL

LOT: xxxxxxxx BUD: 09/21/2014 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. EQ TO BASE.**

Each mL Contains: Fentanyl Citrate 20 mcg, Sodium Chloride 9 mg, pH Adjuster: Hydrochloric Acid/Sodium Hydroxide.

NDC: 52533-025-10



(01) 0 0352533 02510 5

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Rx Only

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 877-666-5222 www.cantrelldrug.com*

Non-MedNet

FENTANYL CITRATE

fentanyl citrate injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-025
Route of Administration	INTRAVENOUS	DEA Schedule	CII

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FENTANYL CITRATE (UNII: MUN5LYG46H) (FENTANYL - UNII:UF599785JZ)	FENTANYL	20 ug 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
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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:52533-025-10	30 mL in 1 SYRINGE, 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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