

FAMOTIDINE- famotidine 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.

Famotidine 2 mg/mL in 0.9% Sodium Chloride 10 mL Syringe

Famotidine

in 0.9% Sodium Chloride 10 mL

20 mg/10 mL (2 mg/mL)

Lot: xxxxx BUD: Quantity: 10 mL


Store Refrigerated. Single-Dose Syringe.

Injection Solution for IV Use.

Rx Only


NDC: 52533-172-12 CMPD Date: 09/99 Hospital/Office Use

Outsourced Compounded Drug. Not for Resale.



(01) 0 0352533 17212 0

Cantrell Drug Company
 7321 Cantrell Road Little Rock, AR 72207
 877-666-5222 www.cantrelldrug.com



FAMOTIDINE 2 mg/mL

Each mL: Famotidine 2 mg, Sodium Chloride 9 mg, Mannitol 4 mg, L-Aspartic Acid 0.8 mg. pH adj: L-Aspartic Acid / NaOH

00002

FAMOTIDINE

famotidine injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52533-172
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FAMOTIDINE (UNII: 5QZO15J2Z8) (FAMOTIDINE - UNII:5QZO15J2Z8)	FAMOTIDINE	2 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	9 mg in 1 mL
MANNITOL (UNII: 3OWL53L36A)	4 mg in 1 mL
ASPARTIC ACID (UNII: 30KYC7MIAD)	0.005 mg in 1 mL
WATER (UNII: 059QF0KO0R)	

Other Ingredients		
Ingredient Kind	Ingredient Name	Quantity
May contain	HYDROCHLORIC ACID (UNII: QTT17582CB)	
May contain	SODIUM HYDRO XIDE (UNII: 55X04QC32I)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52533-172-12	10 mL in 1 SYRINGE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/11/2014	

Labeler - Cantrell Drug Company (035545763)

Revised: 6/2014

Cantrell Drug Company