

FAMOTIDINE- famotidine tablet, film coated
Precision Dose, Inc.

Famotidine Tablets

20 mg

For Hospital Use Only

LI1731 Rev. 10/24

Drug Facts

Active ingredient (in each tablet)

Famotidine 20 mg

Purpose

Acid reducer

Uses

- relieves heartburn associated with acid indigestion and sour stomach
- prevents heartburn associated with acid indigestion and sour stomach brought on by eating or drinking certain food and beverages

Warnings

Allergy alert

Do not use if you are allergic to famotidine or other acid reducers

Do not use

- if you have trouble or pain swallowing food, vomiting with blood, or bloody or black stools. These may be signs of a serious condition.
See your doctor.
- with other acid reducers

Ask a doctor before use if you have

- had heartburn over 3 months. This may be a sign of a more serious condition.
- heartburn with **lightheadedness, sweating, or dizziness**
- chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or lightheadedness
- frequent **chest pain**
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain
- kidney disease

Ask a doctor or pharmacist before use if you are taking a prescription drug. Acid reducers may interact with certain prescription drugs.

Stop use and ask a doctor if

- your heartburn continues or worsens
- you need to take this product for more than 14 days

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- **Use as directed per healthcare professional.**
- adults and children 12 years and over:
 - to **relieve** symptoms, swallow 1 tablet with a glass of water. Do not chew.
 - to **prevent** symptoms, swallow 1 tablet with a glass of water at any time from **10 to 60 minutes before** eating food or drinking beverages that cause heartburn
 - do not use more than 2 tablets in 24 hours
- children under 12 years: ask a doctor

Other information

- read the directions and warnings before use
- store at 20°-25°C (68°-77°F)
- protect from moisture

Inactive ingredients

carnauba wax, colloidal silicon dioxide, croscarmellose sodium, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, talc, titanium dioxide

Questions or comments?

1-800-397-9228

GLUTEN FREE

How Supplied

NDC 68094-054-65

Unit Dose Packages of 200 Tablets (20 × 10) per Carton

Packaged by:

Precision Dose, Inc.

South Beloit, IL 61080

For inquiries call Precision Dose, Inc. at 1-800-397-9228 or email

druginfo@precisiondose.com

LI1731 Rev. 10/24

PRINCIPAL DISPLAY PANEL - 20 mg Tablet Blister Pack Carton

PrecisionDose™

NDC 68094-054-65

Unit Dose

Famotidine

Tablets 20 mg

200 Tablets

(20 x 10)

(in each tablet)

Famotidine 20 mg

Acid reducer

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GLUTEN FREE

USUAL DOSE: SEE ENCLOSED DRUG FACTS

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Hospital Use Only.

LC1735

R0

Packaged by:

Precision Dose, Inc.

South Beloit, IL 61080



PrecisionDose™

NDC 68094-054-65

Unit Dose

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Tablets 20 mg**

**200 Tablets
(20 x 10)**

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Famotidine 20 mg.....Acid reducer

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FAMOTIDINE

famotidine tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68094-054(NDC:0113-0194)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FAMOTIDINE (UNII: 5QZO15J2Z8) (Famotidine - UNII:5QZO15J2Z8)	FAMOTIDINE	20 mg

Inactive Ingredients

Ingredient Name	Strength
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ05DW1A)	

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

Product Characteristics

Color	WHITE	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	L194
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68094-054-65	20 in 1 CARTON	03/28/2024	
1	NDC:68094-054-59	10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077351	03/28/2024	

Labeler - Precision Dose, Inc. (035886746)

Revised: 5/2025

Precision Dose, Inc.