FAMOTIDINE- famotidine tablet
Little Pharma, Inc.

Tips for Managing Heartburn

- Do not lie flat or bend over after eating
- Do not wear tight fitting clothing around th estomach
- Do not eat before bedtime
- Raise the head of your bed
- Avoid heartburn-causing foods such as rich, spicy, fatty or fried foods, chocolate, caffeine, alcohol, and certain fruits and vegetables

Just One Tablet prevents and relieves heartburn die to acid indigestion brough on by leating and drinking certain foods and beverages

Important: Read all directions and warnings before use. Tamper Evident: Do not use if imprented inner safety seal is torn or missing

Drug Facts

Active Ingredient (in each tablet)

Famotidine USP 10mg

Acid reducer

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

Alergy Alert

Do not us if 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 Contol Center right away. (1-800-222-1222)

Directions

- adults and children 12 years and over:
- to **relieve** symtoms, 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 **15 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 20º to 25ºC (68º to 77ºF)
- protect from moisture

carnuba wax, corn starch, hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, sodium starch glycolate, talc, titanium dioxide

Questions or comments?

Call **1-844-243-1241** or email hi@curistrelief.com

curist

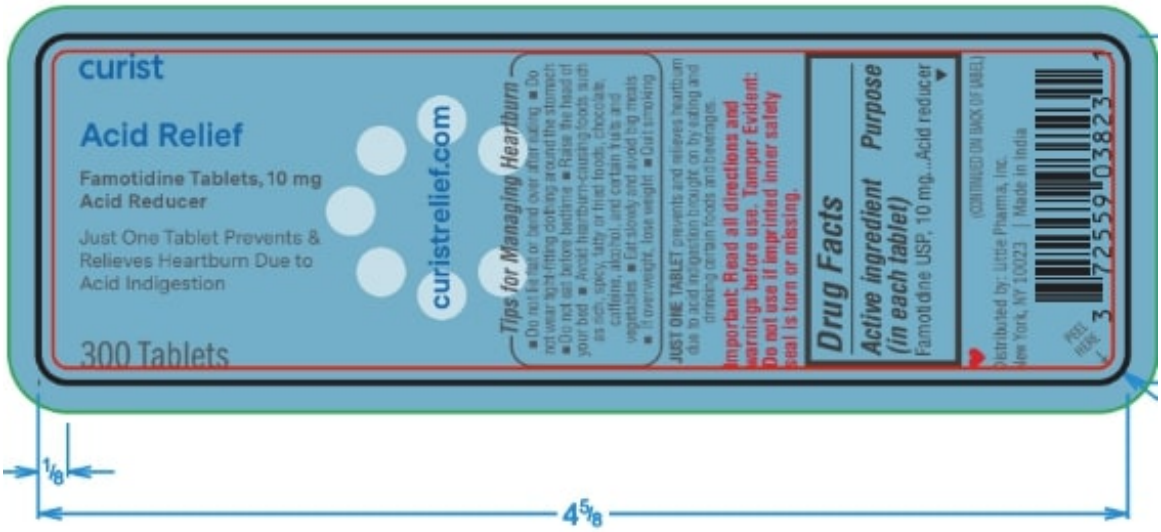
Acid Relief

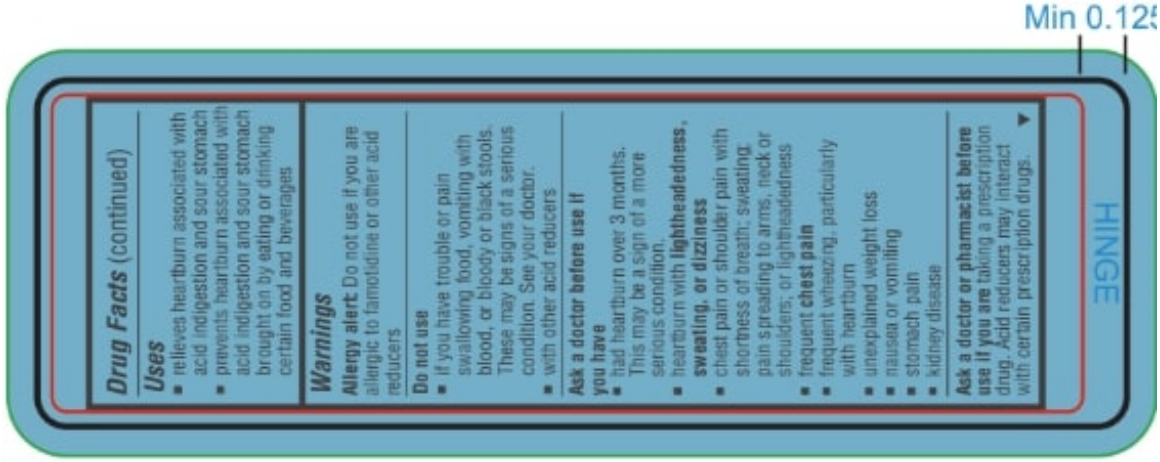
Famotidine Tablets, 10mg

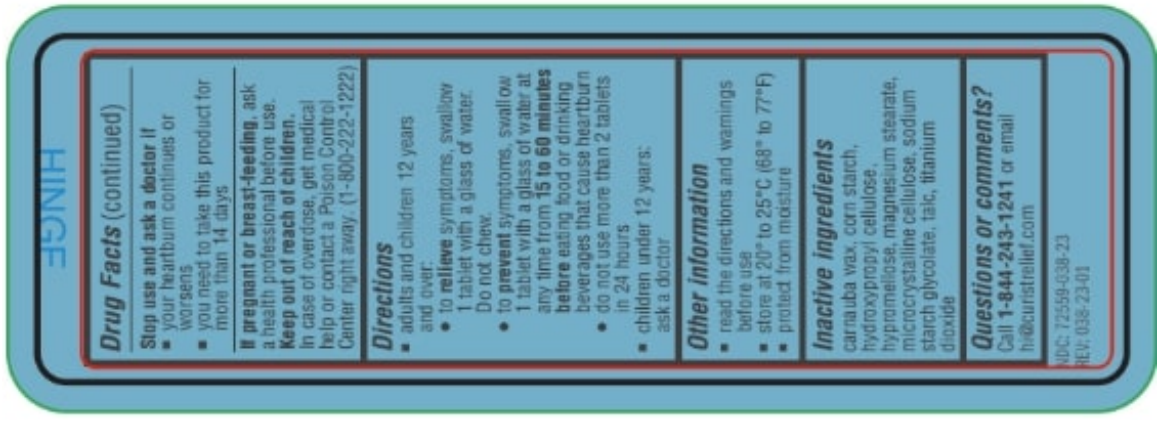
Acid Reducer

Just One Tablet Prevents & Relieves Heartburn Dut to Acid Indigestion

300 Tablets







FAMOTIDINE

famotidine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72559-038
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FAMOTIDINE (UNII: 5QZO15J2Z8) (FAMOTIDINE - UNII:5QZO15J2Z8)	FAMOTIDINE	10 mg in 10 mg

Inactive Ingredients

Ingredient Name	Strength
TALC (UNII: 7SEV7J4R1U)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
MICROCRYSTALLINE CELLULOSE 102 (UNII: PNR0YF693Y)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STARCH, CORN (UNII: O8232NY3SJ)	
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)	
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)	
HYPROMELLOSE 2910 (15 MPA.S) (UNII: 36SFW2JZ0W)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	

Product Characteristics

Color	white	**Score**	score with uneven pieces
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	T;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72559-038-23	300 mg in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA215766	05/15/2025	

Labeler - Little Pharma, Inc. (074328189)

Establishment

Name	Address	ID/FEI	Business Operations
Annora Pharma Private Limited		650980746	manufacture(72559-038)

Establishment

Name	Address	ID/FEI	Business Operations
Nutra-Med Packaging LLC		022004902	pack(72559-038)

Revised: 4/2025

Little Pharma, Inc.