CURIST ACID RELIEF- famotidine 20 mg tablet
Little Pharma, Inc.

Curist Acid Relief (famotidine 20 mg tablets)

curistrelief.com

Important: Read all directions and warnings before use.

Tamper Evident: do not use if imprinted inner safety seal is torn or missing

Drug Facts

Active ingredient (in each tablet)

Famotidine USP, 20 mg

Purpose

Acid reducer

(CONTINUED ON BACK OF LABEL)

REV: 032-23-01

NDC 72559-032-23

Distributed by: Little Pharma, Inc.

New York, NY 10023 | Made in India

*Maximum strength famotidine tablet available over-the-counter (OTC) without a prescription

PEEL HERE

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 **lighheadedness, sweating, or dizziness**
- chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or lightheadness
- 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)

- 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 warning before use
- store at 20°-25°C (68°-77°F)
- protect from moisture

hydroxypropyl cellulose, hypromellose, macrogol, magnesium stearate, microcrystalline cellulose, pre-gelatinized starch, sodium starch glycolate, talc, titanium dioxide, triacetin

Questions or comments?

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

REV: 032-23-01

STOP PEELING

curist

Acid Relief

Maximum Strength OTC*

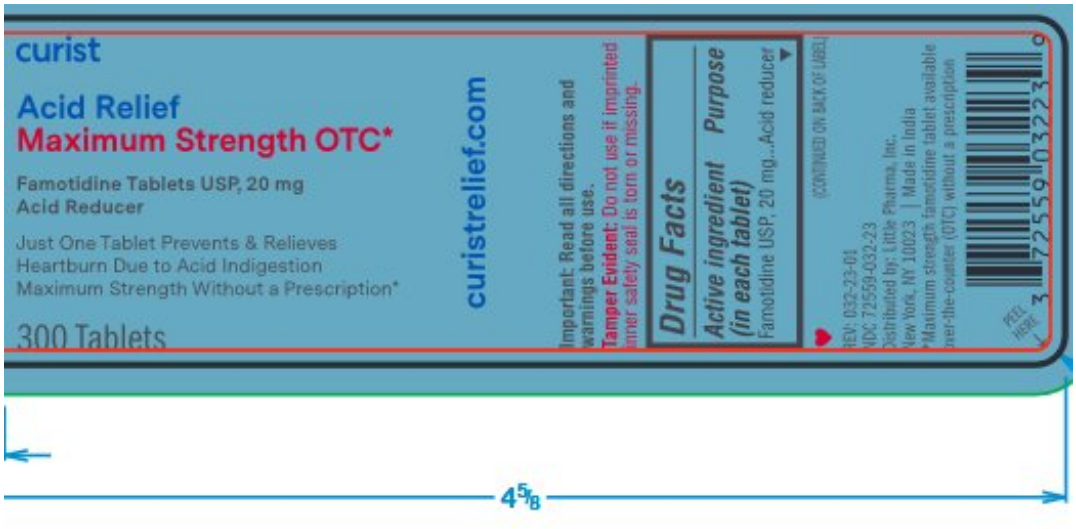
Famotidine Tablets USP, 20 mg

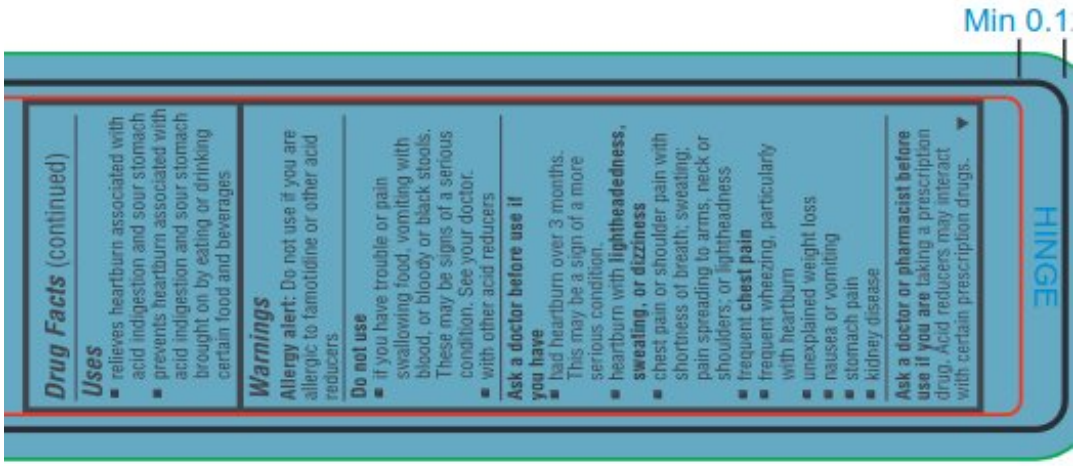
Acid Reducer

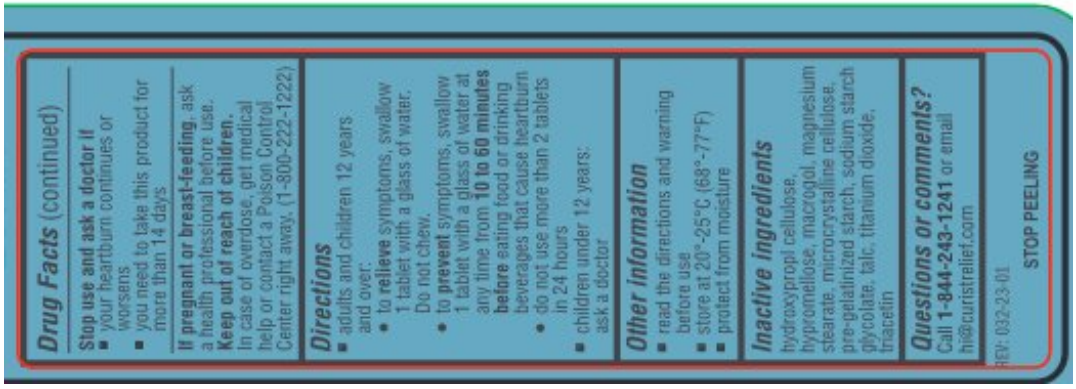
Just One Tablet Prevents & Relieves Heartburn Due to Acid Indigestion

Maximum Strength Without a Prescription*

300 Tablets







CURIST ACID RELIEF

famotidine 20 mg tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72559-032
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
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TRIACETIN (UNII: XHX3C3X673)	
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
HYDROXYPROPYL CELLULOSE (110000 WAMW) (UNII: 5Y0974F5PW)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (white to off-white)	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	V;15
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72559-032-23	300 in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA215822	01/31/2024	

Labeler - Little Pharma, Inc. (074328189)

Revised: 1/2024 Little Pharma, Inc.