ACID RELIEF- famotidine tablet, film coated Rite Aid Corporation

Rite Aid Corporation Acid Relief 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

- 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
- keep the carton. It contains important information.
- store at 20°-25°C (68°-77°F)
- · protect from moisture

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, FD&C blue #1 aluminum lake, hypromellose, lactose (monohydrate), magnesium stearate, maltodextrin, microcrystalline cellulose, modified food starch, natural and artificial flavor, sucralose, titanium dioxide, triacetin

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

FREE FROM | GLUTEN FREE

SOY FREE

Compare to the active ingredient of Maximum Strength Pepcid® AC

MAXIMUM STRENGTH

ACID RFI IFF

Famotidine Tablets, 20 mg

Acid Reducer

Just one tablet prevents & relieves heartburn due to acid indigestion

COOL MINT FLAVOR

Releases a cooling sensation in mouth & throat

ACTUAL SIZE

50 TABLETS



ACID RELIEF

famotidine tablet, film coated

| Product Information | | | |
|----------------------------|----------------|--------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:11822-1014 |
| 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 |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| CROSCARMELLOSE SODIUM (UNII: M280L1HH48) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) | |
| LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | |
| MALTODEXTRIN (UNII: 7CVR7L4A2D) | |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| SUCRALOSE (UNII: 96K6UQ3ZD4) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| TRIACETIN (UNII: XHX3C3X673) | |

| Product Characteristics | | | | |
|-------------------------|-------|--------------|----------|--|
| Color | BLUE | Score | no score | |
| Shape | ROUND | Size | 8mm | |
| Flavor | | Imprint Code | 32F | |
| Contains | | | | |

| P | ackaging | | | |
|---|----------------------|---|-------------------------|-----------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:11822- 1014-0 | 1 in 1 CARTON | 03/31/2022 | |
| 1 | | 50 in 1 BOTTLE; Type 0: Not a Combination Product | | |

| Marketing Start Date | Marketing End Date |
|-------------------------|-----------------------|
| 03/31/2022 | |
| | Date |

Labeler - Rite Aid Corporation (014578892)

Revised: 4/2022 Rite Aid Corporation