

FAMOTIDINE - famotidine tablet, film coated
Wal-Mart Stores, Inc.

Drug Facts

Active ingredient (in each tablet)

Famotidine USP 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 (1-800-222-1222) right away.

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° to 25°C (68° to 77°F)
- protect from moisture

Inactive ingredients

carnauba wax, corn starch, hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, red iron oxide, sodium starch glycolate, talc, titanium dioxide, yellow iron oxide.

Questions or comments?

1-888-287-1915

Tips for Managing Heartburn

- Do not lie flat or bend over after eating
- Do not wear tight-fitting clothing around the stomach
- 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
- Eat slowly and avoid big meals
- If overweight, lose weight
- Quit smoking

***JUST ONE TABLET* prevents and relieves heartburn due to acid indigestion brought on by eating and drinking certain foods and beverages.**

Do not use if carton is open or if printed foil seal under bottle cap is open or torn.

**DISTRIBUTED BY: Wal-Mart Inc.,
Bentonville, AR 72716
MADE IN INDIA**

Code: TS/DRUGS/22/2009

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL -20 mg (25 Tablets, Container Label)

**SEE NEW WARNINGS
NDC 49035-505-35
equate™
MAXIMUM STRENGTH
Famotidine
Tablets USP, 20 mg**

Acid Reducer

- Just one tablet prevents and relieves heartburn due to acid indigestion

TABLETS

SEE NEW WARNINGS

equate™ NDC 49035-505-35

MAXIMUM STRENGTH

Famotidine
Tablets USP, 20 mg

Acid Reducer

- Just one tablet prevents and relieves heartburn due to acid indigestion

25 TABLETS

Do not use if printed foil seal on bottle is broken or missing.

Active ingredient (in each tablet)
Famotidine USP 20 mg.....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 (1-800-222-1222) right away. **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° to 25°C (68° to 77° F) ■ protect from moisture **Inactive ingredients** See carton for complete list of inactive ingredients

Questions or comments? 1-888-281-1915

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716
MADE IN INDIA

Code: TS/DRUGS/22/2009 Lot: P 1421077
UM-3884

Lot:
Expiry:

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL -20 mg (25 Tablets, Container Carton Label)

SEE NEW WARNINGS
NDC 49035-505-35

equate™

**Compare
to Maximum
Strength
Pepcid® AC
active
ingredient***

**MAXIMUM STRENGTH
Famotidine
Tablets USP, 20 mg**

Acid Reducer

- Just one tablet prevents and relieves heartburn due to acid indigestion

Smaller tablet than the brand with the same effect

Actual Size

**20
mg**

**25
TABLETS**



FAMOTIDINE

famotidine tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-505
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)	
STARCH, CORN (UNII: O8232NY3SJ)	
HYDROXYPROPYL CELLULOSE (160000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

Product Characteristics

Color	YELLOW	Score	no score
Shape	ROUND (Square shaped Biconvex)	Size	5mm
Flavor		Imprint Code	CC;59
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-505-35	1 in 1 CARTON	04/26/2016	
1		25 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49035-505-14	1 in 1 CARTON	08/06/2020	
2		50 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:49035-505-21	1 in 1 CARTON	08/06/2020	
3		100 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:49035-505-34	1 in 1 CARTON	08/06/2020	
4		200 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:49035-505-95	2 in 1 CARTON	08/06/2020	
5		50 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:49035-505-80	2 in 1 CARTON	08/06/2020	
6		100 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:49035-505-39	1 in 1 CARTON	01/21/2021	
7		365 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206531	04/26/2016	

Labeler - Wal-Mart Stores, Inc. (051957769)

Registrant - Aurohealth LLC (078728447)

Establishment

Name	Address	ID/FEI	Business Operations
APL HEALTHCARE LIMITED		650844777	ANALYSIS(49035-505) , MANUFACTURE(49035-505)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(49035-505) , MANUFACTURE(49035-505)

Revised: 12/2021

Wal-Mart Stores, Inc.