ACID CONTROLLER- famotidine tablet, film coated
Walgreen Company

Walgreen Co. Acid Controller 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.
- if you have kidney disease, except under the advice and supervision of a 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

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.

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 and light

Inactive ingredients

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

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to Maximum Strength Pepcid® AC active ingredient

Acid Controller

FAMOTIDINE TABLETS, 20 mg / ACID REDUCER

MAXIMUM STRENGTH

Just one tablet prevents & relieves heartburn due to acid indigestion

25 TABLETS

ACTUAL SIZE


Walgreens
Acid Controller
FAMOTIDINE TABLETS, 20 mg /
ACID REDUCER
ORG0318-F
Walgreens
Compare to Maximum Strength Pepcid® AC active ingredient††
NDC 0363-0701-02
Acid Controller
FAMOTIDINE TABLETS, 20 mg /
ACID REDUCER
MAXIMUM STRENGTH
• Just one tablet prevents & relieves heartburn due to acid indigestion
25 TABLETS
ACTUAL SIZE

ACID CONTROLLER

famotidine tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-0701
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)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL (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:0363-0701-71	1 in 1 CARTON	09/26/2006	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0363-0701-72	1 in 1 CARTON	06/02/2011	
2		60 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0363-0701-01	1 in 1 CARTON	10/29/2007	
3		85 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:0363-0701-02	25 in 1 CARTON	09/26/2006	
4		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:0363-0701-39	30 in 1 CARTON	04/10/2015	04/10/2015
5		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:0363-0701-82	1 in 1 CARTON	05/08/2019	
6		200 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	ANDA077351	09/26/2006	

Labeler - Walgreen Company (008965063)

Revised: 5/2019 Walgreen Company