# ACID CONTROLLER- famotidine tablet, film coated Walgreen Company

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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.
- 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 60minutes 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

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**

Walgreens

WALGREENS PHARMACIST RECOMMNEDED

Compare to the active ingredient in Maximum Strength Pepcid® AC

Acid Controller

FAMOTIDINE TABLETS, 20 mg / ACID REDUCER

Maximum Strength

Just one tablet prevents & relieves heartburn due to acid indigestion
 85 TABLETS
 ACTUAL SIZE

SEE NEW WARNINGS



Drug Facts (continued)

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#### **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	<b>Basis of Strength</b>	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: M280L1HH48)			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POLYVINYL ALCOHOL, UNSPECIFIED (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				

P	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			
	NDC 03C3 0701				

2	NDC:0303-0701-	1 in 1 CARTON	06/02/2011	09/30/2020
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: 5/2022 Walgreen Company