FOSTER AND THRIVE ACID REDUCER- famotidine tablet, film coated Praxis, LLC

McKesson Acid Reducer 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 preventsymptoms, 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

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

Questions or comments?

Call 833-358-6431 Monday to Friday 9:00am to 7:00pm EST

Principal Display Panel

COMPARE TO MAXIMUM STRENGTH PEPCID ® AC ACTIVE INGREDIENT

Foster & Thrive ™

MAXIMUM STRENGTH

Acid Reducer

Famotidine Tablets 20 mg

ACID REDUCER

Gluten Free

Just One Tablet Prevents & Relieves Heartburn Due to Acid Indigestion

ACTUAL SIZE
SEE NEW WARNINGS
50 TABLETS



■ do not use more than 2 tablets in 2.4 hours ■ children under 1.2 ye as: ask a doctor

Other information
■ read the direction sand warningsbefore use
■ keep the carton, it contains important information. ▶

Questions or comments? Cal 833-35 8-6431 Monday to Filday 9:00am to 7:00pm EST

"This product is not ma nufactured or dishribute dby Johnson & John son: Consumer Inc., dishributor of Maximum Strength Pepcid™ AC.

FOSTER AND THRIVE ACID REDUCER

famotidine tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG Item Code (Source)		NDC:59368-286
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: 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			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59368-286- 01	1 in 1 CARTON	08/31/2023	
1		50 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	08/31/2023	

Labeler - Praxis, LLC (016329513)

Establishment			
Name	Address	ID/FEI	Business Operations
Praxis, LLC		016329513	manufacture(59368-286) , label(59368-286) , pack(59368-286)

Revised: 1/2023 Praxis, LLC