ACID REDUCER- famotidine tablet, film coated Praxis, LLC

HEB 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 ating 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?

1-800-719-9260

Package/Label Principal Display Panel

Compare to Maximum Strength Zantac 360° [™]active ingredient

H-E-B ®

Maximum Strength

Acid Reducer

Famotidine Tablets, 20 mg

Acid Reducer

JUST ONE TABLET Prevents & Relieves Heartburn Due to Acid Indigestion

actual size



ACID REDUCER famotidine tablet, film coated famotidine tablet, film coated Product Information Product Type HUMAN OTC DRUG Route of Administration ORAL Active Ingredient/Active Woiety Ingredient Name Basis of Strength Strength

	Inactive Ingredients Ingredient Name						
CARNAUBA WAX (UNII: R12CBM0EIZ)							
CROSCARMELLOSI	· · · · · · · · · · · · · · · · · · ·	280L1HH48)					
LACTOSE MONOH							
MAGNESIUM STEARATE (UNII: 70097M6I30)							
MICROCRYSTALLIN	NE CELLULOSE (UN	III: OP1R32D61U)					
POLYETHYLENE G	LYCOL, UNSPECIF	IED (UNII: 3WJQ0SDW1A)					
POLYVINYL ALCOH	IOL, UNSPECIFIED	(UNII: 532B59J990)					
TALC (UNII: 7SEV7J4	4R1U)						
TITANIUM DIOXIDE	(UNII: 15FIX9V2JP)						
Product Chara	acteristics						
Color	white	Score	Score		no score		
Shape	ROUND	Size	Size		8mm		
Flavor		Imprint Co	Imprint Code		L194		
Contains							
Packaging							
	Packa	ge Description	Ма	rketing Start Date	Marketing End Date		
	Packa 1 in 1 CARTON	ge Description	Ma 12/19/	Date	-		
 # Item Code 1 NDC:59368-294- 01 	1 in 1 CARTON	ge Description ype 0: Not a Combinatior	12/19/	Date	-		
 # Item Code 1 NDC:59368-294- 01 	1 in 1 CARTON 50 in 1 BOTTLE; T		12/19/	Date	-		
 # Item Code 1 NDC:59368-294- 01 	1 in 1 CARTON 50 in 1 BOTTLE; T		12/19/	Date	-		
 # Item Code 1 NDC:59368-294- 01 1 	1 in 1 CARTON 50 in 1 BOTTLE; Ty Product	ype 0: Not a Combination	12/19/	Date	-		
1 NDC:59368-294-	1 in 1 CARTON 50 in 1 BOTTLE; Ty Product	ype 0: Not a Combination	12/19/	Date	-		

Labeler - Praxis, LLC (016329513)

Establishment

Name	Address	ID/FEI	Business Operations		
Praxis, LLC		016329513	label(59368-294) , pack(59368-294) , manufacture(59368-294)		

Revised: 1/2023