BERKLEY AND JENSEN FAMOTIDINE- famotidine tablet, film coated BJWC

BJWC Famotidine Tablets, 20 mg 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 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?

1-800-934-1204

Principal Display Panel

Compare to the Active Ingredient in Maximum Strength Pepcid[®] AC

berkley jensen®

MAXIMUM STRENGTH

FAMOTIDINE TABLETS, 20mg

ACID REDUCER

JUST ONE TABLET PREVENTS & RELIEVES HEARTBURN DUE TO ACID INDIGESTION

ACTUAL SIZE 200 TABLETS | 2x100 COUNT BOTTLES 100% MONEY-BACK GUARANTEE



BERKLEY AND JENSE	EN FAMOTIDINE					
famotidine tablet, film coated						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:68	DC:68391-300	
Route of Administration	ORAL					
Active Ingredient/Active Moiety						
Ingredient Name Basis of Stren			ength	Strength		
FAMOTIDINE (UNII: 5QZO15J2Z8)	(FAMOTIDINE - UNII:5QZ01	5J2Z8)	FAMOTIDINE		20 mg	
Inactive Ingredients						
	Ingredient Name				Strength	
CARNAUBA WAX (UNII: R12CBM08	EIZ)					
SILICON DIOXIDE (UNII: ETJ7Z6XE	3U4)					

LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) MAGNESIUM STEARATE (UNII: 70097M6I30) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)
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

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68391-300- 78	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/09/2013		
2	NDC:68391-300- 82	2 in 1 PACKAGE	12/09/2013		
2		100 in 1 BOTTLE; Type 0: Not a Combination Product			
Marketing Information					
	Marketing	Application Number or Monograph	Marketing Start	Marketing End	

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077351	12/09/2013	

Labeler - BJWC (159082692)

Revised: 12/2023

BJWC