#### FAMOTIDINE- famotidine tablet Bi-Mart

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# **Drug Facts**

#### Active ingredient (in each tablet)

Famotidine 10 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

- with other acid reducers
- 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.

## 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 wheezing, particularly with heartburn
- frequent chest pain
- 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 (1-800-222-1222) 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 preventsymptoms, swallow 1 tablet with a glass of water at any time from 15 to 60 minutes before ating food or drinking beverages that cause heartburn
  - $\circ~$  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
- store at 20° 25°C (68° 77°F)
- protect from moisture

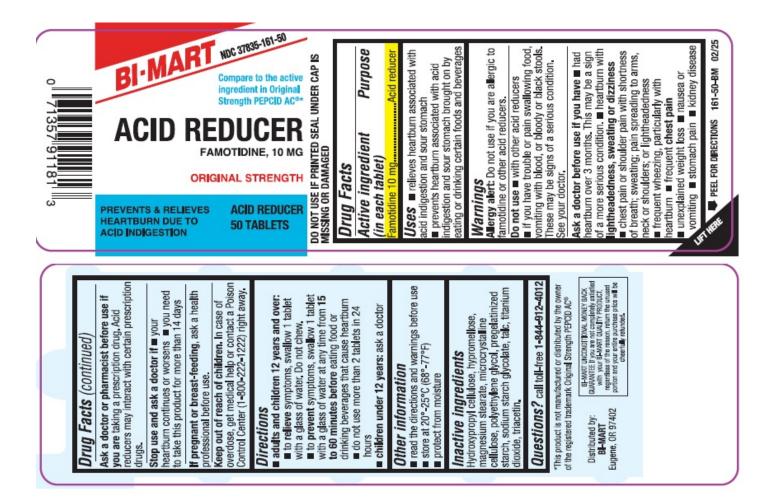
# Inactive ingredients

Hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, pregelatinized starch, sodium starch glycolate, polyethylene glycol, talc, titanium dioxide, triacetin.

# **Questions?**

call toll-free **1-844-912-4012** 

## PACKAGE LABEL-PRINCIPAL DISPLAY PANEL



FAMOTIDINE									
famotidine tablet									
Product Information									
Product Type	HUMAN OTC DRUG	ltem Code (Source)		NDC:37835-161					
Route of Administration	ORAL								
Active Ingredient/Active Moiety									
Ingredient Name Basis of Stre					Strength				
FAMOTIDINE (UNII: 5QZ015J2Z8) (FAMOTIDINE - UNII:5QZ015J2Z8)			FAMOTIDINE		10 mg				
Inactive Ingredients									
		Strength							
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)									
STARCH, CORN (UNII: 08232NY3SJ)									
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)									
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ 8H6N6OH)									
MAGNESIUM STEARATE (UNII: 70097M6I30)									
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)									
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)									

PO	POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)							
TALC (UNII: 7SEV7J4R1U)								
Pı	roduct Chara	cteristics						
		white (white to off-white)	S	Score	no score			
Shape		ROUND	5	Size	5mm			
Flavor			I	mprint Code	V;21			
Contains								
Pa	ackaging							
	ackaging Item Code	Package Desc	ription	Marketing Start Date	Marketing End Date			
#	Item Code	Package Desc 50 in 1 BOTTLE; Type 0: Not Product	- Combination	-	-			
#	Item Code NDC:37835-161-	50 in 1 BOTTLE; Type 0: Not	- Combination	Date	-			
#	Item Code NDC:37835-161- 50	50 in 1 BOTTLE; Type 0: Not Product	- Combination	Date	-			
# 1	Item Code NDC:37835-161- 50	50 in 1 BOTTLE; Type 0: Not	- Combination	Date	-			
#	Item Code NDC:37835-161- 50	50 in 1 BOTTLE; Type 0: Not Product	a Combination 0/	Date	-			

Labeler - Bi-Mart (027630078)

Registrant - Bi-Mart (027630078)

Establishment									
Name	Address	ID/FEI	<b>Business Operations</b>						
VKT Pharma Private Limited		871408062	manufacture(37835-161)						

Revised: 4/2025

Bi-Mart