UP AND UP FAMOTIDINE- famotidine tablet, film coated
Praxis, LLC

Target Corporation 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 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, titanium dioxide

Questions?

Call 1-888-547-7400

Principal Display Panel

see new warnings

Compare to active ingredient in Maximum Strength Pepcid ®AC

maximum strength

famotidine tablets, 20 mg

acid reducer

just one tablet prevents and relieves heartburn due to acid indigestion

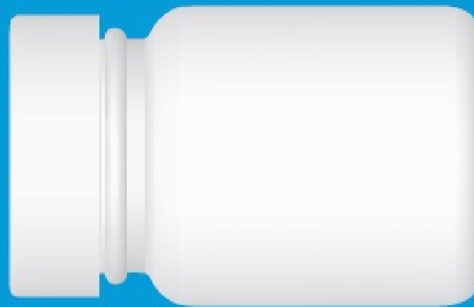
ACTUAL SIZE

50 + 50

100 TABLETS

2 x 50 TABLETS, 100 TOTAL

ACTUAL SIZE



see new warnings

NDC 11673-061-78

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Maximum Strength Pepcid® AC*

maximum strength
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up&up™

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100
TABLETS

2 x 50 TABLETS, 100 TOTAL

A8002 UW C1

245 05 0702 R00
C-001227-01-082

3 70030 27598 4

GLUTEN FREE

Distributed by Target Corporation
Minneapolis, MN 55403
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PAPER BOX | PLASTIC BOTTLE | Empty & Discard Cap

how2recycle.info

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Drug Facts (continued)

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maximum strength
famotidine tablets, 20 mg
acid reducer

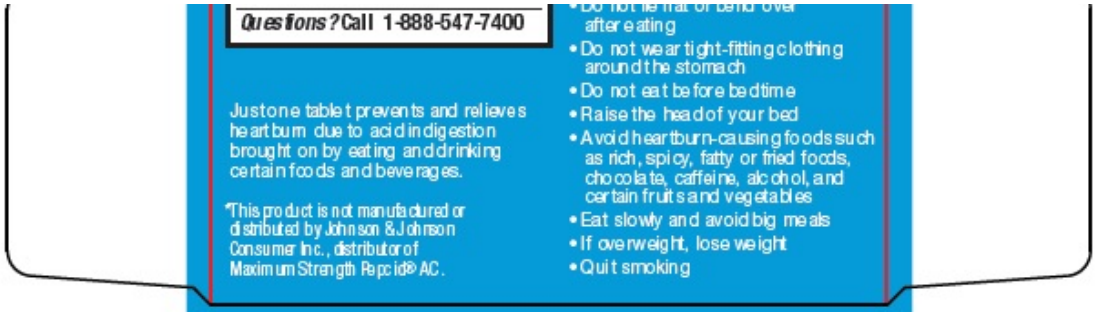
up&up™

DO NOT USE IF PRINTED FOIL UNDER CAP IS BROKEN OR MISSING

Drug Facts (continued)

Tips for Managing Heartburn

- Do not lie flat or bend over



UP AND UP FAMOTIDINE

famotidine tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59368-237
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: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
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
	NDC:59368-237			

1	NDC:59368-237-03	1 in 1 CARTON	05/14/2012	
1		50 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:59368-237-01	2 in 1 CARTON	04/17/2020	
2		50 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:59368-237-02	1 in 1 CARTON	11/24/2021	
3		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	05/03/2012	

Labeler - Praxis, LLC (016329513)

Establishment

Name	Address	ID/FEI	Business Operations
Praxis, LLC		016329513	manufacture(59368-237) , label(59368-237) , pack(59368-237)

Revised: 1/2023 Praxis, LLC