

**UP AND UP FAMOTIDINE- famotidine tablet, film coated**  
**Target Corporation**

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**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 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**

colloidal silicon dioxide, croscarmellose sodium, FD&C blue #1 aluminum lake, hypromellose, lactose (monohydrate), magnesium stearate, maltodextrin, microcrystalline cellulose, modified food starch, natural and artificial flavor, sucralose, titanium dioxide, triacetin

**Questions?**

**Call 1-888-547-7400**

**Package/Label Principal Display Panel**

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

release a cooling sensation in mouth and throat

ACTUAL SIZE

COOL MINT FLAVOR

up & up™

25 TABLETS

25 TABLETS



## UP AND UP FAMOTIDINE

famotidine tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11673-414	
<b>Route of Administration</b>	ORAL			
<b>Active Ingredient/Active Moiety</b>				
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
	FAMOTIDINE (UNII: 5QZO15J2Z8) (FAMOTIDINE - UNII:5QZO15J2Z8)	FAMOTIDINE	20 mg	
<b>Inactive Ingredients</b>				
	<b>Ingredient Name</b>		<b>Strength</b>	
	CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)			
	FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
	HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
	LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
	MAGNESIUM STEARATE (UNII: 70097M6I30)			
	MALTODEXTRIN (UNII: 7CVR7L4A2D)			
	MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
	SUCRALOSE (UNII: 96K6UQ3ZD4)			
	TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
	TRIACETIN (UNII: XHX3C3X673)			
	SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
<b>Product Characteristics</b>				
<b>Color</b>	BLUE	<b>Score</b>	no score	
<b>Shape</b>	ROUND	<b>Size</b>	8mm	
<b>Flavor</b>		<b>Imprint Code</b>	32F	
<b>Contains</b>				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:11673-414-63	1 in 1 CARTON	01/25/2022	
1		25 in 1 BOTTLE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
ANDA	ANDA077351	01/25/2022		

**Labeler** - Target Corporation (006961700)

