

**CAREONE ACID RELIEF- famotidine tablet, film coated**  
**Retail Business Services, LLC**

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**Retail Business Services, LLC. Acid Relief 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 or comments?**

**1-800-719-9260**

### **Package/Label Principal Display Panel**

CAREone™

Compare to the active ingredient in Maximum Strength Pepcid® AC

MAXIMUM STRENGTH

ACID RELIEF

Famotidine Tablets, 20mg

Acid Reducer

Just one tablet prevents & relieves heartburn due to acid indigestion

Releases a cooling sensation in mouth & throat

Cool Mint Flavor

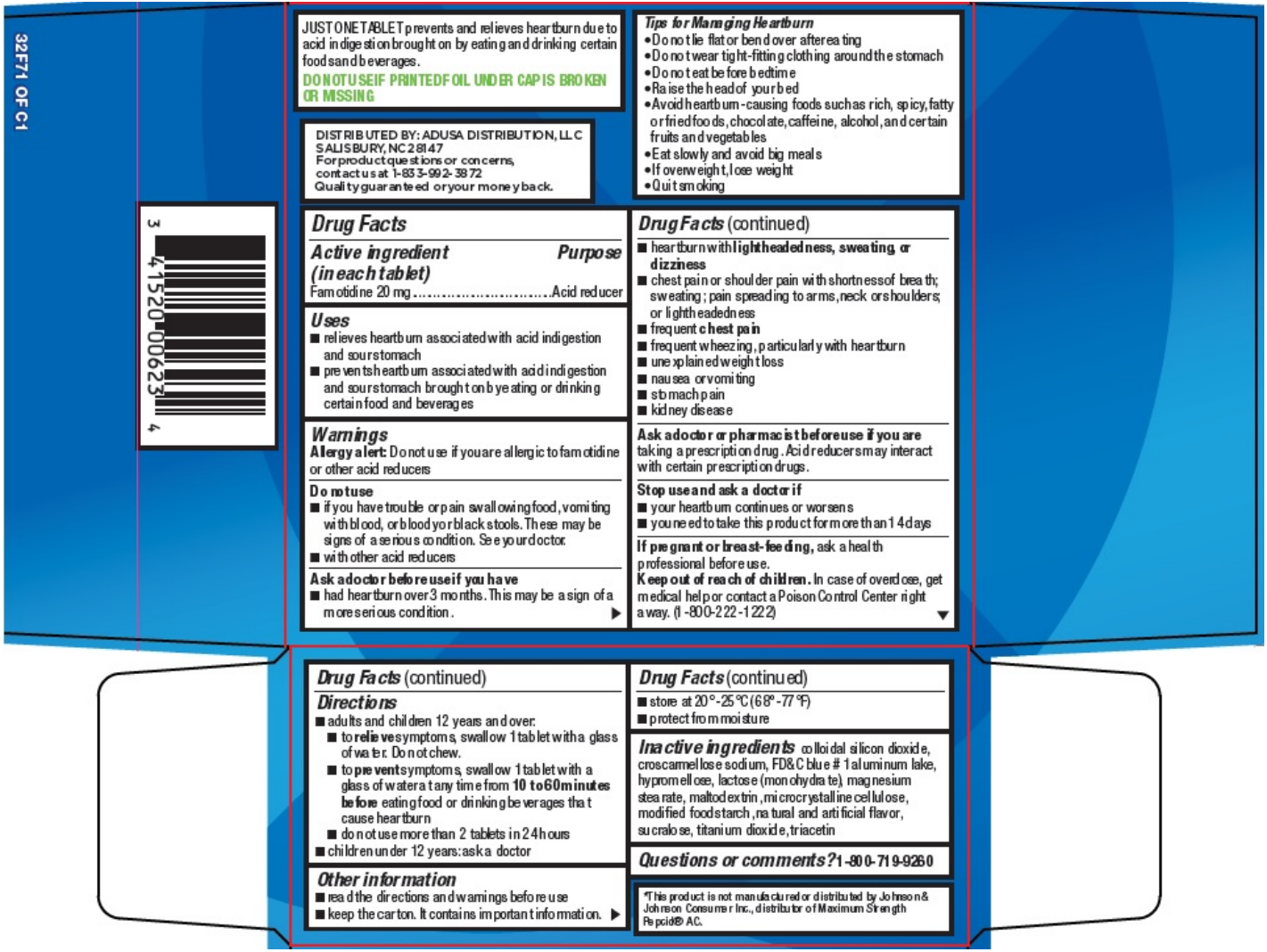
Gluten Free

Actual Size

OUR PHARMACISTS RECOMMEND

50 TABLETS





Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72476-299
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			

<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	

### Product Characteristics

<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>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72476-299-71	1 in 1 CARTON	06/03/2022	
1		50 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	06/03/2022	

**Labeler** - Retail Business Services, LLC (967989935)

Revised: 6/2022

Retail Business Services, LLC