

PEPCID AC MAXIMUM STRENGTH- famotidine tablet, film coated
Johnson & Johnson Consumer Inc.

PEPCID® AC

Maximum Strength

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, hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, pregelatinized starch, talc, titanium dioxide

Questions or comments?

1-800-755-4008 (toll-free) or **215-273-8755** (collect)

PRINCIPAL DISPLAY PANEL

NDC 16837-855-14

MAXIMUM STRENGTH

Pepcid®

AC

Famotidine Tablets 20 mg

Acid Reducer

Just One Tablet!

Prevents & Relieves Heartburn

Due to Acid Indigestion

actual size

25 Tablets



Drug Facts (continued)

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.

Questions or Comments?
 1-800-755-4008 (toll-free) or 215-273-8755 (collect)

Drug Facts (continued)

Active ingredient
 Famotidine 20 mg.....Acid reducer

Purpose
 • Heartburn with high acid levels, sweating, or chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or light-headedness
 • Frequent chest pain
 • Frequent heartburn, particularly with heartburn
 • Unexplained weight loss
 • Nausea or vomiting
 • Stomach pain • Kidney disease

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 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.

Uses
 • Relieves heartburn associated with acid indigestion and sour stomach
 • Prevents heartburn associated with eating or drinking certain food and beverages

Drug Facts
 • Heartburn with high acid levels, sweating, or dizziness
 • Chest pain or shoulder pain with shortness of breath; sweating; pain spreading to arms, neck or shoulders; or light-headedness
 • Frequent chest pain
 • Frequent heartburn, particularly with heartburn
 • Unexplained weight loss
 • Nausea or vomiting
 • Stomach pain • Kidney disease

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 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.

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.

Questions or Comments?
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MAXIMUM STRENGTH
Pepcid[®] AC

Famotidine (active ingredient) made in Hungary
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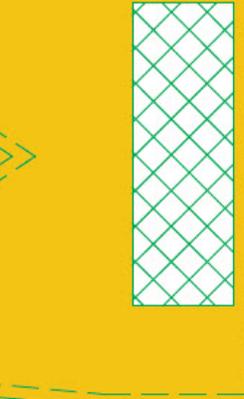
MAXIMUM STRENGTH
Pepcid[®] AC
 25 Tablets

NDC 16837-855-14

MAXIMUM STRENGTH
Pepcid[®] AC

Tips for Managing Heartburn

- Do not lie flat or bend over after eating around the stomach. • Do not eat before bedtime. • Raise the head of your bed. • Avoid heartburn-causing foods such as rich, spicy, fatty, or fried foods, chocolate, caffeine, alcohol, and certain fruits and vegetables.
- Eat slowly and avoid big meals.
- Quit smoking.
- If overweight, lose weight.



30049137





PEPCID AC MAXIMUM STRENGTH

famotidine tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:16837-855
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)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	no score
Shape	SEMI-CIRCLE (D shaped)	Size	9mm
Flavor		Imprint Code	PAC;20
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:16837-855-05	1 in 1 CARTON	09/01/2003	12/31/2023
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:16837-855-25	5 in 1 CARTON	09/01/2003	03/31/2014
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:16837-855-49	1 in 1 CARTON	09/01/2003	12/31/2023
3		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:16837-855-50	1 in 1 CARTON	09/01/2003	
4		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
5	NDC:16837-855-52	50 in 1 TRAY	09/01/2003	
5		1 in 1 POUCH; Type 0: Not a Combination Product		
6	NDC:16837-855-65	1 in 1 CARTON	09/01/2003	10/25/2013
6		65 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
7	NDC:16837-855-67	1 in 1 CARTON	09/01/2003	12/31/2023
7		65 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
8	NDC:16837-855-70	1 in 1 CARTON	09/01/2003	
8		70 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
9	NDC:16837-855-75	1 in 1 CARTON	09/01/2003	
9		75 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
10	NDC:16837-855-90	125 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	03/15/2021	
11	NDC:16837-855-80	1 in 1 CARTON	09/01/2003	
11		80 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
12	NDC:16837-855-85	1 in 1 CARTON	09/01/2003	
12		85 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
13	NDC:16837-855-09	1 in 1 PACKAGE	09/01/2003	
13		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
14	NDC:16837-855-12	1 in 1 CARTON	09/01/2003	
14		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
15	NDC:16837-855-13	1 in 1 POUCH; Type 0: Not a Combination Product	09/01/2003	
16	NDC:16837-855-14	5 in 1 CARTON	09/01/2003	
16		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
	NDC:16837			

17	NDC:16837-855-15	1 in 1 CARTON	09/01/2003	
17		35 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
18	NDC:16837-855-17	1 in 1 CARTON	09/01/2003	12/31/2023
18		65 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
19	NDC:16837-855-18	1 in 1 CARTON	09/01/2003	
19		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
20	NDC:16837-855-40	5 in 1 CARTON	09/01/2003	12/31/2023
20		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
21	NDC:16837-855-51	12 in 1 CARTON	09/01/2003	12/31/2023
21		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
22	NDC:16837-855-16	1 in 1 CARTON	09/01/2003	
22		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
23	NDC:16837-855-20	2 in 1 PACKAGE	08/22/2014	
23		1 in 1 CARTON		
23		50 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
24	NDC:16837-855-19	1 in 1 CARTON	04/27/2015	
24		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
25	NDC:16837-855-24	2500 in 1 CARTON	06/05/2023	
25		1 in 1 POUCH; Type 0: Not a Combination Product		
26	NDC:16837-855-02	2 in 1 CARTON	06/05/2023	
26		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020325	09/01/2003	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 12/2023

Johnson & Johnson Consumer Inc.