FAMOTIDINE- famotidine tablet, film coated DISCOUNT DRUG MART

Famotidine

Drug Facts

Active ingredient (in each tablet)

Famotidine, USP 10 mg Famotidine, USP 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

Famotidine USP, 10 mg

- 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.

Famotidine USP, 20 mg

- with other acid reducers
- if you have kidney disease, except under the advice and supervision of a doctor.
- 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

- frequent **chest pain**
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain
- 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

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

Famotidine USP, 10 mg

- 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 **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

Famotidine USP, 20 mg

- 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

- store at 20° to 25° C (68° to 77° F)
- protect from moisture
- read the directions and warnings before use
- keep the carton. It contains important information.

Inactive ingredients

colloidal silicon dioxide, hydroxypropyl cellulose, hypromellose, iron oxide red (only for Famotidine USP, 10 mg), magnesium stearate, microcrystalline cellulose, polyethylene glycol 400, pregelatinized starch, talc, titanium dioxide

Questions?

call 1-800-406-7984

DISTRIBUTED BY DRUG MART-FOOD FAIR MEDINA, OHIO 44256

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

DISCOUNT drug mart FOOD FAIR MAY CONTAIN ANTI-THEFT DEVICE

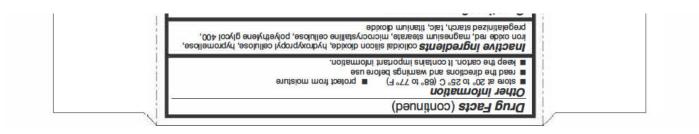
Original Strength ACID CONTROLLER Famotidine Tablets, USP 10 mg

Acid Reducer Just One Tablet Prevents & Relieves Heartburn Due to Acid Indigestion

*COMPARE TO THE ACTIVE INGREDIENT OF ORIGINAL STRENGTH PEPCID AC®

30 TABLETS





PRINCIPAL DISPLAY PANEL - 20 mg Tablet Bottle Carton

DISCOUNT drug mart FOOD FAIR

MAY CONTAIN ANTI-THEFT DEVICE

Maximum Strength ACID CONTROLLER Famotidine Tablets, USP 20 mg

Acid Reducer Just One Tablet Prevents & Relieves Heartburn Due to Acid Indigestion

*COMPARE TO THE ACTIVE INGREDIENT OF MAXIMUM STRENGTH PEPCID AC®

25 TABLETS



Drug Facts Drug Facts Active ingredient Purpose

amotidine tablet, film c	oated						
Product Information							
Product Type	HUMAN	OTC DRUG	Item Code (Sou	rce)	NDC:539	43-036	
Route of Administration	ORAL						
Active Ingredient/Active Ingre	ctive Moiety						
	Ingredient	Name		Basis of Str	rength	Strength	
FAMOTIDINE (UNII: 5QZC	•		28)	FAMOTIDINE		10 mg	
Inactive Ingredients							
		ngredient Name				Strength	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)							
HYDROXYPROPYL CELL	,	AW) (UNII: RFW2ET6	71P)				
HYPROMELLOSE, UNSPI							
FERRIC O XIDE RED (UNII: 1K09F3G675)							
MAGNESIUM STEARATE (UNII: 70097M6I30)							
MICRO CRYSTALLINE CH		21R32D61U)					
POLYETHYLENE GLYCO	L 400 (UNII: B69789	94SGQ)					
STARCH, CORN (UNII: 08	232NY3SJ)						
TALC (UNII: 7SEV7J4R1U)							
TITANIUM DIO XIDE (UNI							
, , , , , , , , , , , , , , , , , , ,	,						
Product Characteris	tics						
Color	pink	Score	Score			no score	
Shape	ROUND	Size	Size			8 mm	
Flavor		Imprint C	Imprint Code		035		
Contains							
Packaging							
# Item Code	Marke	ting Start Date	e Marke	ting End Dat			
1 NDC:53943-036-25 30 i	-	e Description Ype 0: Not a Combin		•		U III	
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Marketing Inform	mation						

06/25/2010

FAMOTIDINE							
famotidine tablet, film coate	ed						
Product Information							
Product Type	HUMA	N OTC DRUG	Item Code (Sour	rce)	NDC:5394	43-037	
Route of Administration	ORAL						
Active Ingredient/Activ	e Moiety						
Ingredient Name Basis of Strength						Strength	
FAMOTIDINE (UNII: 5QZO15J	•			FAMOTIDINE	0	20 mg	
		. ,				0	
Inactive Ingredients							
mactive ingreatents		Ingredient Name				Strength	
SILICON DIOXIDE (UNII: ETJ)		Ingreulent Name				Strength	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D6 1U)							
MAGNESIUM STEARATE (UNII: 70097M6I30)							
HYDROXYPROPYL CELLULOSE (1200000 MW) (UNII: RFW2ET671P)							
STARCH, CORN (UNII: 08232NY3SJ)							
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)							
TALC (UNII: 7SEV7J4R1U)							
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)							
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)							
Product Characteristics	_						
				-	no score		
Color	white			8mm			
Shape	ROUND						
Flavor		Imprint Code 036					
Contains							
Packaging							
# Item Code	em Code Package Description			Marketing Start Date		Marketing End Date	
1 NDC:53943-037-31 25 in 1 BOTTLE; Type 0: Not a Combination Product 07/23/2010							
Marketing Informa	tion						
		han an Maragana I. C'	- 41	ng Chart Day	Master		
					Market	ing End Date	
ANDA ANDA	A090283		07/23/201	U			

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment					
Name	Address	ID/FEI	Business Operations		
Ohm Laboratories Inc.		184769029	manufacture(53943-036, 53943-037)		

Revised: 9/2018

DISCOUNT DRUG MART