### TAGAMET- cimetidine tablet Medtech Products Inc.

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#### Tagamet

### Drug Facts

### Active ingredient (in each tablet)

Cimetidine 200mg

### 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 foods and beverages

### Warnings

Allergy alert: Do not use if you are allergic to cimetidine 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

- 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
- kidney disease
- liver 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

- you need to take this product for more than 14 days
- stomach pain continues
- your heartburn continues or worsens

## 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 (1-800-222-1222) right away.

# Directions

- adults and children 12 years and over:
  - to **relieve** symptoms, swallow 1 tablet with a glass of water
  - to **prevent** symptoms, swallow 1 tablet with a glass of water **right before or any time up to 30 minutes before** eating food or drinking beverages that cause heartburn
  - do not take more than 2 tablets in 24 hours
- children under 12 years: ask a doctor

# Other information

store at 15-30°C (59-86°F)

## Inactive ingredients

cellulose, corn starch, hypromellose, magnesium stearate, polyethylene glycol, polysorbate 80, povidone, sodium lauryl sulfate, sodium starch glycolate, titanium dioxide

# Questions?

Call toll-free 1-800-482-4394 weekdays

# PRINCIPAL DISPLAY PANEL

Tagamet® HB 200 Cimetidine Tablets 200 mg Acid Reducer 70 tablets (70 doses)



#### PRINCIPAL DISPLAY PANEL

Tagamet® HB 200 Cimetidine Tablets 200 mg Acid Reducer

30 tablets (30 doses)



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Pı	oduct Infor	mation							
			HUMAN OTC DE		tom Code	(5	•		029-222
	oduct Type				Item Code (Source) NDC			NDC:03	029-222
Ro	oute of Admini	stration	ORAL						
۵۵	tive Ingredi	ent/Active	Mojety						
			dient Name			Basi	of Str	anath	Strengt
	METIDINE (UNII:	-		meBasis of StE - UNII:80061L1WGD)CIMETIDINE				ligti	200 mg
In	active Ingre	dients							
			Ingredie	nt Name					Strength
ST.	ARCH, CORN (UI	NII: 08232NY3S	-						
	PROMELLOSES								
MA	GNESIUM STEA	RATE (UNII: 70	097M6I30)						
PO	LYETHYLENE G	LYCOL, UNSPE	ECIFIED (UNII: 3	WJQ0SDW1A)					
РО	LYSORBATE 80	(UNII: 60ZP392	ZG8H)						
	VIDONE (UNII: FZ								
	DIUM LAURYL S								
	DIUM STARCH			(UNII: 5856J3	G2A2)				
	ANIUM DIOXIDI	E (FINITE FEXAN)							
			-						
MI	CROCRYSTALLII	NE CELLULOSI	E (UNII: OP1R32						
MI		NE CELLULOSI	E (UNII: OP1R32						
MI	CROCRYSTALLII	NE CELLULOSI	E (UNII: OP1R32						
MI(	CROCRYSTALLII	NE CELLULOSI ALUMINUM LA	E (UNII: OP1R32						
мі FD <b>Pr</b>	CROCRYSTALLII &C BLUE NO. 1	NE CELLULOSI ALUMINUM LA	e (UNII: OP1R321 Ake (UNII: J9EQA				no score		
мі FD Pr Co	CROCRYSTALLII &C BLUE NO. 1 roduct Chara	NE CELLULOSI ALUMINUM LA	E (UNII: OP1R320 AKE (UNII: J9EQA S	4352JM)			no score 13mm		
MI( FD Pr Co Sh	CROCRYSTALLII &C BLUE NO. 1 <b>Oduct Chara</b> lor	ALUMINUM LA	E (UNII: OP1R32I AKE (UNII: J9EQA S D S	A3S2JM)	e				
MI FD Pr Co Sh Fla	CROCRYSTALLII &C BLUE NO. 1 roduct Chara lor ape	ALUMINUM LA	E (UNII: OP1R32I AKE (UNII: J9EQA S D S	A3S2JM) Score Size	e		13mm		
MI FD Pr Co Sh Fla	CROCRYSTALLII &C BLUE NO. 1 oduct Chara lor ape ivor	ALUMINUM LA	E (UNII: OP1R32I AKE (UNII: J9EQA S D S	A3S2JM) Score Size	e		13mm		
MI FD Co Sh Fla Co	CROCRYSTALLII &C BLUE NO. 1 oduct Chara lor ape avor ntains	ALUMINUM LA	E (UNII: OP1R32I AKE (UNII: J9EQA S D S	A3S2JM) Score Size	e		13mm		
MI FD Co Sh Fla Co	CROCRYSTALLII &C BLUE NO. 1 oduct Chara lor ape ivor	ALUMINUM LA ACTERISTICS white DIAMON	E (UNII: OP1R32I AKE (UNII: J9EQA S D S	A3S2JM) Score Size mprint Cod		1arketing Date	13mm TAGAME	Γ;200	eting End Date
Pr Co Sh Fla Co	CROCRYSTALLII &C BLUE NO. 1 roduct Chara lor ape tvor ntains	ALUMINUM LA ACTERISTICS white DIAMON	E (UNII: OP1R32I AKE (UNII: J9EQA D S I	A3S2JM) Score Size mprint Cod	N		13mm TAGAME	Γ;200	_
Pr Co Sh Fla Co Pa #	CROCRYSTALLII &C BLUE NO. 1 roduct Chara lor ape ivor ntains ckaging Item Code NDC:63029-	ALUMINUM LA ACTERISTICS white DIAMON 1 in 1 BOX	E (UNII: OP1R32I AKE (UNII: J9EQA D S I	A3S2JM) Score Size mprint Code	<b>N</b> 06/	Date	13mm TAGAME	Γ;200	_
Pr Co Sh Fla Co Pa #	CROCRYSTALLII &C BLUE NO. 1 roduct Chara lor ape ivor ntains ckaging Item Code NDC:63029-	ALUMINUM LA ALUMINUM LA ACTERISTICS white DIAMON 1 in 1 BOX 6 in 1 BLISTER Product 1 in 1 BOX	E (UNII: OP1R32I AKE (UNII: J9EQA D S D I Ackage Desc R PACK; Type 0: 1	A3S2JM) Score Size mprint Code ription	06/ ation 06/	Date	13mm TAGAME	Γ;200	_
Pr Co Sh Fla Co Pa # 1	CROCRYSTALLII &C BLUE NO. 1 roduct Chara lor ape vor ntains ckaging item Code NDC:63029- 222-02	ALUMINUM LA ALUMINUM LA ACTERISTICS white DIAMON 1 in 1 BOX 6 in 1 BLISTER Product 1 in 1 BOX	E (UNII: OP1R320 AKE (UNII: J9EQA D S D I Ackage Desc	A3S2JM) Score Size mprint Code ription	06/ ation 06/	<b>Date</b> 01/2012	13mm TAGAME	Γ;200	_
MIR FD Co Sh Fla Co Pa # 1 1	CROCRYSTALLII &C BLUE NO. 1 roduct Chara lor ape ivor ntains ckaging item Code NDC:63029- 222-02	ALUMINUM LA ALUMINUM LA ACTERISTICS white DIAMON 1 in 1 BOX 6 in 1 BLISTER Product 1 in 1 BOX 30 in 1 BLISTER Product 1 in 1 BOX	E (UNII: OP1R32I AKE (UNII: J9EQA D S D I Ackage Desc R PACK; Type 0: 1	A3S2JM) Score Size mprint Code ription Not a Combin : Not a Combin	ation 06/ ination 06/ ination 06/	<b>Date</b> 01/2012	13mm TAGAME	Γ;200	_

4	NDC:63029- 222-04	4 in 1 BOX	06/01/2019			
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product				
5	NDC:63029- 222-05	4 in 1 BOX	06/01/2019			
5		9 in 1 BLISTER PACK; Type 0: Not a Combination Product				
6	NDC:63029- 222-70	1 in 1 CARTON	06/01/2012			
6		70 in 1 BOTTLE; Type 0: Not a Combination Product				
7	NDC:63029- 222-06	7 in 1 BOX	06/01/2012			
7		10 in 1 BLISTER PACK; Type 0: Not a Combination Product				
8	NDC:63029- 222-09	1 in 1 CARTON	06/01/2012			
8		90 in 1 BOTTLE; Type 0: Not a Combination Product				
9	NDC:63029- 222-11	1 in 1 CARTON	06/01/2012			
9		100 in 1 BOTTLE; Type 0: Not a Combination Product				
10	NDC:63029- 222-12	2 in 1 BOX	06/01/2012			
10		6 in 1 BLISTER PACK; Type 0: Not a Combination Product				
Marketing Information						

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020238	06/01/2012	
NDA	NDA020238	06/01/2012	

TAGAMET					
cimetidine tablet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (	Source)	NDC:63	029-223
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingree	dient Name		Basis of Stre	ength	Strength
CIMETIDINE (UNII: 80061L1WGD) (	CIMETIDINE - UNII:80061L1	WGD)	CIMETIDINE		200 mg
Inactive Ingredients					
	Ingredient Name				Strength
STARCH, CORN (UNII: 08232NY3S	J)				

HYPROMELLOSES	•	•							
		(UNII: 70097M6I30)							
POLYETHYLENE C	GLYCC	DL, UNSPECIFIED (UN	NII: 3WJQ	0SDW1A)					
POLYSORBATE 80 (UNII: 60ZP39ZG8H)									
POVIDONE (UNII: FZ 989GH94E)									
SODIUM LAURYL SULFATE (UNII: 368GB5141J)									
		OLATE TYPE A POTA	ATO (UN	III: 5856J3G2A2)					
TITANIUM DIOXID									
		ELLULOSE (UNII: OP1F							
FD&C BLUE NO. 3	1 ALU	MINUM LAKE (UNII: J9	9EQA3S2	2JM)					
	_								
Product Char	acte	ristics							
Color		blue	S	Score		2 pieces			
Shape		DIAMOND	S	Size 13mm					
Flavor		PEPPERMINT	Ir	Imprint Code TAGAM			1ET;200		
Contains									
Packaging									
# Item Code	Item Code Package Description		ion	Marketing S Date	itart	Marketing End Date			
<b>1</b> NDC:63029- 223-30				02/01/2023					
1 10 in 1 BLISTER PACK; Type 0: Not Product			a Combination						
Marketing	Inf	ormation							
Category			Application Number or Monograph Citation		Marketing Start Date		Date		
NDA	N	DA020238			02/01/2023				

Labeler - Medtech Products Inc. (122715688)

Revised: 5/2023

Medtech Products Inc.