TOPCARE OMEPRAZOLE- omeprazole tablet, delayed release Topco Associates LLC

Topco Associates LLC. Omeprazole Drug Facts

Active ingredient (in each tablet)

Omeprazole 20 mg

Purpose

Acid reducer

Use

- treats frequent heartburn (occurs <u>2 or more</u> days a week)
- not intended for immediate relief of heartburn; this drug may take 1 to 4 days for full effect

Warnings

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

Do not use if you have:

- trouble or pain swallowing food, vomiting with blood, or bloody or black stools
- 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

These may be signs of a serious condition. See you doctor.

Ask a doctor before use if you have:

- had heartburn over 3 months. This may be a sign of a more serious condition.
- frequent wheezing, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain

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
- you need to take more than 1 course of treatment every 4 months
- you get diarrhea
- you develop a rash or joint pain

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

- for adults 18 years of age and older
- this product is to be used once a day (every 24 hours), every day for 14 days
- it may take 1 to 4 days for full effect; some people get complete relief of symptoms within 24 hours

14-Day Course of Treatment

- swallow 1 tablet with a glass of water before eating in the morning
- take every day for 14 days
- do not take more than 1 tablet a day
- do not use for more than 14 days unless directed by your doctor
- · swallow whole. Do not chew, crush, or suck tablets.

Repeated 14-Day Courses (if needed)

- you may repeat a 14-day course every 4 months
- do not take for more than 14 days or more often than every 4 months unless directed by a doctor
- children under 18 years of age: ask a doctor. Heartburn in children may sometimes be caused by a serious condition.

Other information

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

Inactive ingredients

carnauba wax, FD&C blue #1/brilliant blue FCF aluminum lake, hypromellose, hypromellose acetate succinate, lactose monohydrate, menthol, monoethanolamine, sodium lauryl sulfate, sodium starch glycolate, sodium stearate, sodium stearyl fumarate, sucralose, talc, titanium dioxide, triacetin, triethyl citrate

Questions or comments?

1-888-423-0139

Package/Label Principal Display Panel

TopCare® health

VALUE PACK! THREE-14 TABLET BOTTLES

Omeprazole DELAYED RELEASE TABLETS 20 mg

COMPARE TO PRILOSEC OTC®

3 Bottles Inside

ACID REDUCER

24 HR

Treats Frequent Heartburn!

COOL MINT COATED TABLET

SWALLOW - DO NOT CHEW

42 TABLETS

THREE 14-DAY COURSES OF TREATMENT • MAY TAKE 1 TO 4 DAYS FOR FULL EFFECT actual size

Tips for Managing Heartburn

- Do not lie flat or bend over after eating
- Do not wear tight-fitting clothing 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 If overweight, lose weight
- Quit smoking



VALUE PACK!

NDC 76162-470-03

DIE CUT WINDOW DIE CUT WINDOW



PRILOSEC OTC®*

3 Rottles Inside

COOL MINT

COATED

TABLET

actual



3 Bottles Inside

TopCare .

Omeprazole

DELAYED RELEASE TABLETS 20 mg ACID REDUCER

COOL MINT COATED TABLET

SWALLOW -DO NOT CHEW

Do not use if printed seal under cap is broken or

*This product is not manufactured or distributed by Procter & Gamble, distributor of Philoses OTCB.

Omeprazole **DELAYED RELEASE TABLETS 20 mg**



42 TABLETS

THREE 14-DAY COURSES OF TREATMENT • MAY TAKE 1 TO 4 DAYS FOR FULL EFFECT



47KD7 88 C1



Drug Facts

Active ingredient (in each tablet)

Purpose

■ you get diarrhea ■ you develop a rash or joint pain

If pregnant or breast-feeding, ask a health professional before use.

Warmings Allergy alert: Do not use if you are allergic to omeprazole

■ treats frequent heartburn (occurs 2 or more days a week)
■ not intended for immediate relief of heartburn; this drug may take 1 to

Do not use if you have:

trouble or pain swallowing food, vomiting with blood, or bloody or black stools

hearthum with lightheadedness, oweating or dizziness

chest pain or shoulder pain with shortness of breath; sweating pain spreading to arms, neck or shoulders; or lighthe adedness

frequent chest pain

These may be signs of a serious condition. See your doctor.

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- consisten.

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 nausea or vomiting stomach pain

Drug Facts (continued)

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- 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).

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- this product is to be used once a day (every 24 hours), every day for
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Drug Facts (continued)

Inactive ingredients

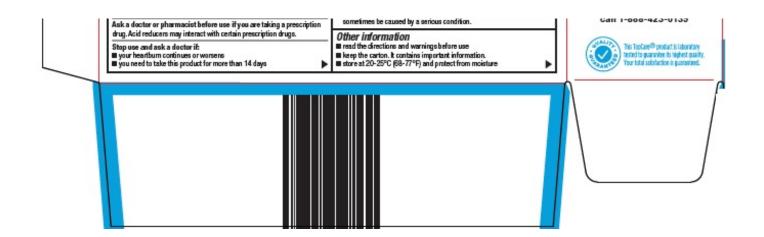
camauba wax, FD&C blue #1/brilliant blue FCF aluminum lake, hypromellose, hypromellose acetate succinate, lactose monohydrate, menthol, monoethanolamine sodium lauryl sulfate, sodium starch glycolate, sodium stearate, sodium stearyl marate, sucralose, talc, titanium dioxide, triacetin, triethyl citrate

Questions or comments? 1-989-423-0139

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TOPCARE OMEPRAZOLE

omeprazole tablet, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76162-470

Route of Administration ORAL

TRIETHYL CITRATE (UNII: 8Z96QXD6UM)

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII: KG60484QX9) OMEPRAZOLE 20 mg

Inactive Ingredients		
Ingredient Name	Strength	
CARNAUBA WAX (UNII: R12CBM0EIZ)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)		
MONOETHANOLAMINE (UNII: 5KV86114PT)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
SODIUM STEARATE (UNII: QU7E2XA9TG)		
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
TRIACETIN (UNII: XHX3C3X673)		

Product Characteristics			
Color	BLUE	Score	no score
Shape	OVAL	Size	12mm
Flavor	MINT (COOL)	Imprint Code	20

Contains

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76162-470- 03	3 in 1 CARTON	09/22/2022	
1		14 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:76162-470- 01	1 in 1 CARTON	09/22/2022	
2		14 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA022032	09/22/2022	

Labeler - Topco Associates LLC (006935977)

Revised: 10/2022 Topco Associates LLC