OMEPRAZOLE- omeprazole tablet, delayed release Walgreens

OMEPRAZOLE MAGNESIUM DELAYED-RELEASE TABLETS

Active ingredient (in each tablet)

Omeprazole delayed-release tablet 20 mg

(equivalent to 20.6 mg omeprazole magnesium)

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
- Omeprazole may cause severe skin reactions. Symptoms may include:
 - skin reddening
 - blisters
 - rash

If an allergic reaction occurs, stop use and seek medical help right away.

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

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 everyday 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 or crush 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° to 25°C (68° to 77°F) and protect from moisture

Inactive ingredients

acetyl tributyl citrate, colloidal silicon dioxide, corn starch, croscarmellose sodium, hydroxypropyl cellulose, hypromellose 2910, magnesium stearate, methacrylic acid copolymer type C, microcrystalline cellulose, polyethylene glycol 400, polyethylene glycol 3350, polyethylene glycol 8000, polyvinyl alcohol, red iron oxide, sucrose, talc, titanium dioxide, triethyl citrate

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

Principal display panel

†Compare to the active ingredient in Prilosec OTC®

24-hour

Omeprazole

Delayed-release tablets 20 mg

Acid reducer

Treats FREQUENT heartburn!

MAY TAKE 1 TO 4 DAYS FOR FULL EFFECT

Tablets

One 14-day course of treatment

†This product is not manufactured by AstraZeneca AB, owner of the registered trademark Prilosec OTC®.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed By

Walgreen Co. Deerfield, IL 60015 100% Satisfaction Guaranteed ©2025 Walgreen Co.

Package label



Omeprazole Delayed Release Tablets 20 mg

OMEPRAZOLE omeprazole tablet, delayed release							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9510				
Route of Administration	ORAL						

9	lient/Active Moiety				
	Ingredient Name			Basis of Strength	Strength
OMEPRAZOLE MA UNII:KG60484QX9)	AGNESIUM (UNII: 426QFE7XLK) (O	MEPRAZ OLE -	OME	PRAZOLE	20.6 mg
Inactive Ingr	edients				
	Ingredien	nt Name			Strength
ACETYLTRIBUTY	LCITRATE (UNII: 0ZBX0N59RZ)				
SILICON DIOXIDE	E (UNII: ETJ7Z6XBU4)				
STARCH, CORN (UNII: O8232NY3SJ)				
CROSCARMELLO	SE SODIUM (UNII: M28OL1HH48)				
HYDROXYPROPY	L CELLULOSE, UNSPECIFIED (UN	NII: 9XZ8H6N6OH)			
HYPROMELLOSE	2910 (15 MPA.S) (UNII: 365FW2)	Z0W)			
MAGNESIUM STE	ARATE (UNII: 70097M6I30)				
METHACRYLIC A	CID AND ETHYL ACRYLATE COP	OLYMER (UNII: NX7	6LV5T8J)		
POLYETHYLENE	GLYCOL 400 (UNII: B697894SGQ)				
POLYETHYLENE	GLYCOL 3350 (UNII: G2M7P15E5F	?)			
POLYETHYLENE	GLYCOL 8000 (UNII: Q662QK8M3I	B)			
POLYVINYL ALCO	HOL, UNSPECIFIED (UNII: 532B5	i9J990)			
FERRIC OXIDE RI	ED (UNII: 1K09F3G675)				
SUCROSE (UNII: C	C151H8M554)				
TALC (UNII: 7SEV7	/J4R1U)				
TITANIUM DIOXII					
	JE (UNII: ISFIA9VZJP)				
	TE (UNII: 8Z96QXD6UM)				
TRIETHYL CITRA	-	510)			
TRIETHYL CITRA	TE (UNII: 8Z96QXD6UM)	510)			
TRIETHYL CITRA MICROCRYSTALL	TE (UNII: 8Z96QXD6UM) INE CELLULOSE (UNII: OP1R32D6	510)			
TRIETHYL CITRA	TE (UNII: 8Z96QXD6UM) INE CELLULOSE (UNII: OP1R32D6	51U)			
TRIETHYL CITRA MICROCRYSTALL	TE (UNII: 8Z96QXD6UM) INE CELLULOSE (UNII: OP1R32D6	51U) Scor	e	no	score
TRIETHYL CITRA MICROCRYSTALL Product Cha i Color	TE (UNII: 8Z96QXD6UM) INE CELLULOSE (UNII: OP1R32D6 racteristics		e		score
TRIETHYL CITRA MICROCRYSTALL Product Cha i Color Shape	TE (UNII: 8Z96QXD6UM) INE CELLULOSE (UNII: OP1R32D6 racteristics red (reddish brown)	Scor Size	e int Code		
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Labeler - Walgreens (008965063)

Revised: 5/2025

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