OMEPRAZOLE MAGNESIUM- omeprazole magnesium capsule, delayed release Walgreens Company

Active ingredient (in each capsule)

*Omeprazole delayed-release capsules 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
- If you develop 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 capsule with a glass of water before eating in the morning
- take every day for 14 days
- do not take more than 1 capsule a day
- do not use for more than 14 days unless directed by your doctor
- swallow whole. Do not chew or crush capsules.

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

black iron oxide, dibasic calcium phosphate, gelatin, glyceryl monostearate, hypromellose 3 cps, magnesium oxide, magnesium stearate, methacrylic acid copolymer dispersion, methacrylic acid copolymer Type B, microcrystalline cellulose, polysorbate 80, potassium hydroxide, propylene glycol, red iron oxide, shellac, silicon dioxide, sodium lauryl sulfate, sugar spheres, talc, titanium dioxide, triethyl citrate

Questions

call **1-888-375-3784**

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

Omeprazole Delayed-Release Capsules, 20 mg*, 14-count - Carton Label

Well at Walgreens WALLGREENS PHARMACIST RECOMMENDED[‡]

NDC 0363-0042-52

ACID REDUCER Omeprazole Delayed-Release Capsules 20 mg* / Acid Reducer

- Treats frequent heartburn
- May take 1 to 4 days for full effect

14 CAPSULES (SAFETY SEALED) ONE 14-DAY COURSE OF TREATMENT

Compare to Prilosec OTC[®] active ingredient^{‡‡}



Omeprazole Delayed-Release Capsules, 20 mg*, 14-count - Bottle Label

Well at Walgreens NDC 0363-0042-52

Omeprazole Delayed-Release Capsules 20 mg* / Acid Reducer

- Treats frequent
- May take 1 to 4 days for full effect

14 CAPSULES (SAFETY SEALED) ONE 14-DAY COURSE OF TREATMENT



OMEPRAZOLE MAGN omeprazole magnesium cap		se		
Product Information				
Product Type	HUMAN OTC DRUG	ltem Code (Source)	NDC:0363-0042(NDC:55111-397)	
Route of Administration	ORAL			
Active Ingredient/Active	Moiety			
Ingredient Name			Basis of Strength Strength	
OMEPRAZOLE MAGNESIUM (UNI	I: 426QFE7XLK) (omepr	azole - UNII:KG60484QX9)	omeprazole	20 mg
Inactive Ingredients				
-	Ingredient N	lame		Strength
FERROSOFERRIC OXIDE (UNII: X	•			
ANHYDROUS DIBASIC CALCIUM	PHOSPHATE (UNII: L1	.1K75P92J)		
Gelatin (UNII: 2G86QN327L)				
GLYCERYL MONOSTEARATE (UN	III: 2300U9XXE4)			
HYPROMELLOSE 2208 (3 MPA.S	5) (UNII: 9H4L916OBU)			
Magnesium Oxide (UNII: 3A3U0GI71G)				
MAGNESIUM STEARATE (UNII: 70	0097M6I30)			
METHACRYLIC ACID - METHYL I	METHACRYLATE COP	OLYMER (1:2) (UNII: 5KY6	i8S2577)	
Methacrylic Acid - Ethyl Acryla	te Copolymer (1:1) T	ype A (UNII: NX76LV5T8J)		
CELLULOSE, MICROCRYSTALLI	NE (UNII: OP1R32D61U)			
Polysorbate 80 (UNII: 60ZP39ZC	58H)			
POTASSIUM HYDROXIDE (UNII: V	VZH3C48M4T)			
PROPYLENE GLYCOL (UNII: 6DC9	Q167V3)			
FERRIC OXIDE RED (UNII: 1K09F3	G675)			
SHELLAC (UNII: 46N107B710)				
SILICON DIOXIDE (UNII: ETJ7Z6XI	BU4)			
RAW SUGAR (UNII: 8M707QY5GH)				
TALC (UNII: 7SEV7J4R1U)				

TRIETHYL CITRATE (UNII: 8Z96QXD6UM) Sodium Lauryl Sulfate (UNII: 368GB5141J)

Product Characteristics							
Color	WHITE, PINK	Score	no score				
Shape	CAPSULE	Size	22mm				
Flavor		Imprint Code	OMP20				
Contains							

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0363-0042- 52	1 in 1 CARTON	09/01/2015					
1		14 in 1 BOTTLE; Type 0: Not a Combination Product						
2	NDC:0363-0042- 27	2 in 1 CARTON	09/01/2015					
2		14 in 1 BOTTLE; Type 0: Not a Combination Product						
3	NDC:0363-0042- 33	3 in 1 CARTON	09/01/2015					
3		14 in 1 BOTTLE; Type 0: Not a Combination Product						
4	NDC:0363-0042- 01	3 in 1 CARTON	09/01/2015					
4		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				
AN	IDA	ANDA078878	09/01/2015					

Labeler - Walgreens Company (008965063)

Revised: 10/2023

Walgreens Company