ESOMEPRAZOLE MAGNESIUM - esomeprazole magnesium capsule, delayed release

Aurohealth LLC

Esomeprazole Magnesium Delayed-Release Capsules USP 20 mg*

Drug Facts

Active ingredient (in each capsule)

*Esomeprazole 20 mg (Each delayed-release capsule corresponds to 22.3 mg esomeprazole magnesium trihydrate USP)

Purpose

Acid reducer

Uses

- 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 esomeprazole.
- Esomeprazole 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 right away. (1-800-222-1222)

Directions

- adults 18 years of age and older
- this product is to be used once a day (every 24 hours), every day for 14 days
- may take 1 to 4 days for full effect

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
- swallow whole. Do not crush or chew capsules.
- do not use for more than 14 days unless directed by your doctor

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 before use. 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)
- Meets USP dissolution test 2

Inactive ingredients

colloidal silicon dioxide, FD&C blue no.1, gelatin, hydroxypropyl cellulose, hypromellose, magnesium carbonate, magnesium oxide, methacrylic acid and ethyl acrylate copolymer dispersion, mono and di glycerides, polysorbate 80, propylene glycol, shellac, sodium lauryl sulfate, strong ammonia solution, sugar spheres (which contains starch (maize) and sucrose), talc, titanium dioxide, triethyl citrate and yellow iron oxide.

Questions or comments?

call **1-855-274-4122** (Monday – Friday 8:30 AM to 5:00 PM EST)

Distributed by: **AUROHEALTH LLC** 279 Princeton-Hightstown Road East Windsor, NJ 08520

Made in India

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg (14 Capsule Container Label)

Healthy Living

NDC 58602-896-05

Esomeprazole Magnesium Delayed-Release Capsules USP 20 mg* Acid Reducer 24 HR

May take 1 to 4 days for full effect **Treats Frequent Heartburn**

Capsules

14 Capsules
One 14-day course of treatment

Top Ply



Top Ply (Page #1)

*Lot: XXXXXXXXX Exp.: YYYY-MMM Prefix, Variables of Lot, Exp and Neutral code shall be printed online during packing.

Warnings

Allergy a left: ■ Do not use if you are allergic to esomeprazole. ■ Esomeprazole 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 = hearburn with lightheaddeness, sweating or dizziness ■ chest pain 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. ■ fequent 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. Sci deducers may interact with certain prescription drug. Sci 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 diarhtae ■ you develop a rask 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
■ adults 18 years of age and older ■ this product is to be used once a day (every 24 hours), every day for 14 days ■ may take 1 to 4 days for full effect. 14-Day Course of Treatment = swallow1 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

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Back of Top Ply (Page #2)

Bottom Ply

Base (Page #3)

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Inactive ingredients colloidal silicon dioxide, FD&C blue no.1, gelatin, hydroxypropyl cellulose, hypromellose, magnesium carbonate, magnesium oxide, methacrylic acid and ethyl acrylate copolymer dispersion, mono and di glycerides, polysorbate 80, propylene glycol, shellac, sodium lauryl sulfate, strong ammonia solution, sugar spheres (which contains starch (maize) and sucrose), talc, titanium dioxide, triethyl citrate and yellow iron oxide.

Questions or comments? call 1-855-274-4122 (Monday - Friday 8:30 AM to 5:00 PM EST)

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg (14 Capsule Container Carton)

Healthy Living

NDC 58602-896-62

HR Active ingredient

See new warning information
Esomeprazole Magnesium
Delayed-Release Capsules USP 20 mg*
Acid Reducer Capsules

24 HR Treats Frequent Heartburn

May take 1 to 4 days for full effect

42 Capsules

(3 bottles of 14 each)
Three 14-day courses of treatment



ESOMEPRAZOLE MAGNESIUM

esomeprazole magnesium capsule, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58602-896
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ESOMEPRAZOLE MAGNESIUM TRIHYDRATE (UNII: R6DXU4WAY9) (ESOMEPRAZOLE - UNII:N3PA6559FT)	ESOMEPRAZ OLE	20 mg	

Inactive Ingredients	
Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM CARBONATE (UNII: 0E53J927NA)	
MAGNESIUM OXIDE (UNII: 3A3U0GI71G)	
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B710)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
AMMONIA (UNII: 5138Q19F1X)	
STARCH, CORN (UNII: 08232NY3SJ)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	

Product Characteristics			
Color	WHITE	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	181
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58602-896- 62	3 in 1 CARTON	11/14/2025	
1		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
ANDA	ANDA209339	11/14/2025	

Labeler - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(58602-896), MANUFACTURE(58602-896)

Revised: 11/2025 Aurohealth LLC