ESOMEPRAZOLE- esomeprazole capsule, delayed release Safeway, Inc.

Esomeprazole

Drug Facts

Active ingredient (in each capsule)

*Esomeprazole 20 mg (Each delayed-release capsule contains 22.3 mg esomeprazole magnesium trihydrate)

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 esomeprazole
- Esomeprazole may cause severe skin reactions. Symptoms 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

unless directed by a doctor

- 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
- 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.
- Store at 20°C to 25°C (68°F to 77°F). [See USP controlled room temperature.]
- keep the carton. It contains important information.

Inactive Ingredients

FD & C blue no 1, FD & C red no 3, ferrosoferric oxide, gelatin, hydroxypropyl cellulose, hypromellose, magnesium stearate, methacrylic acid copolymer dispersion, mono and di glycerides, polysorbate 80, potassium hydroxide, propylene glycol, shellac, sugar spheres (corn starch and sucrose), talc, titanium dioxide and triethyl citrate.

Questions?

Call toll-free weekdays 8:30 AM to 5 PM EST at 1-800-818-4555.

TAMPER-EVIDENT FEATURES: Do not use if seal under bottle cap imprinted with "SEALED for YOUR PROTECTION" or blue band around center of each capsule is broken or missing.

Tips for Managing Heartburn

- Avoid foods or drinks that are more likely to cause heartburn, such as rich, spicy, fatty and fried foods, chocolate, caffeine, alcohol and even some acidic fruits and vegetables.
- Eat slowly and do not eat big meals.
- Do not eat late at night or just before bedtime.
- Do not lie flat or bend over soon after eating.
- Raise the head of your bed.
- Wear loose-fitting clothing around your stomach.
- If you are overweight, lose weight.
- If you smoke, quit smoking.

DISTRIBUTED BY:

BETTER LIVING BRANDS LLC P.O. BOX 99 PLEASANTON, CA 94566-0009

R0922

PRINCIPAL DISPLAY PANEL - 20 mg Capsule Bottle Carton

Compare to Nexium[®] 24HR active ingredient[†]

NDC 21130-926-14

See new warning information

Esomeprazole Magnesium Delayed-Release Capsules USP, 20 mg*

ACID REDUCER

May take 1 to 4 days for full effect One 14-day course of treatment 1 x 14 Count Bottle Inside

Signature care® Quality Guaranteed

24 HR Treats FREQUENT Heartburn

Actual Size

14 CAPSULES



ESOMEPRAZOLE

esomeprazole capsule, delayed release

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:21130-926 Route of Administration ORAL

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

Inactive Ingredients		
	Ingredient Name	Strength

FERROSOFERRIC OXIDE (UNII: XM0M87F357) **GELATIN, UNSPECIFIED** (UNII: 2G86QN327L) HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH) HYPROMELLOSE 2208 (100 MPA.S) (UNII: B1QE5P712K) MAGNESIUM STEARATE (UNII: 70097M6I30) METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER (UNII: NX76LV5T8J) POLYSORBATE 80 (UNII: 60ZP39ZG8H) POTASSIUM HYDROXIDE (UNII: WZH3C48M4T) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) **SHELLAC** (UNII: 46N107B710) TALC (UNII: 7SEV7J4R1U) TITANIUM DIOXIDE (UNII: 15FIX9V2JP) TRIETHYL CITRATE (UNII: 8Z96QXD6UM) STARCH, CORN (UNII: O8232NY3SJ) SUCROSE (UNII: C151H8M554) FD&C BLUE NO. 1 (UNII: H3R47K3TBD) FD&C RED NO. 3 (UNII: PN2ZH5LOQY) **GLYCERYL MONOSTEARATE** (UNII: 2300U9XXE4)

Product Characteristics			
Color	PINK	Score	no score
Shape	CAPSULE	Size	14mm
Flavor		Imprint Code	RG;50
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-926- 42	3 in 1 CARTON	12/07/2022	
1		14 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:21130-926- 28	2 in 1 CARTON	12/07/2022	
2		14 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:21130-926- 14	1 in 1 CARTON	12/07/2022	
3		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	ANDA212866	12/07/2022	

Labeler - Safeway, Inc. (009137209)

Establishment			
Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	Manufacture(21130-926)

Revised: 8/2023 Safeway, Inc.