OMEPRAZOLE- omeprazole tablet, delayed release GERI-CARE PHARMACEUTICALS, CORP

Omeprazole

Drug Facts

Active ingredient (in each tablet)

Omeprazole USP 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.
- 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 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 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

anhydrous lactose, hypromellose, hypromellose acetate succinate, iron oxide red, iron oxide yellow, lactose monohydrate, methyl cellulose, monoethanolamine, propylene glycol, sodium lauryl sulfate, sodium starch glycolate, sodium stearate, sodium stearyl fumarate, talc, triethyl citrate and titanium dioxide.

The imprinting ink contains ammonium hydroxide, black iron oxide, n-butyl alcohol,

propylene glycol and shellac.

Questions or Comments?

Call toll free 1-800-540-3765 weekdays.

Distributed by: **Geri-Care** Pharmaceuticals Corp. 1295 Towbin Avenue Lakewood, NJ 08701

PRINCIPAL DISPLAY PANEL - 20 mg Tablet Bottle Carton

NDC 57896-659-42

GERI CARE[®] LIVE LIFE WELL

Compare To Prilosec OTC[®]*

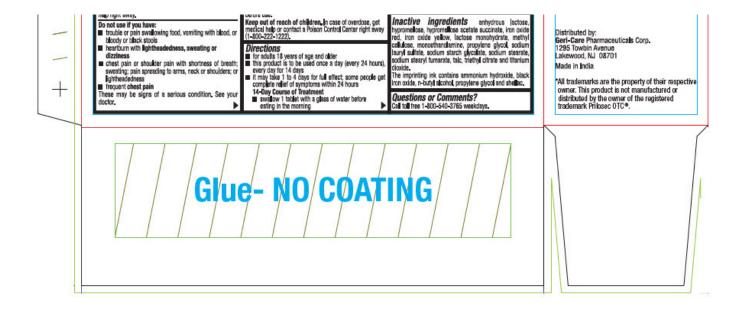
Omeprazole Delayed-release Tablets 20 mg / Acid Reducer SWALLOW- DO NOT CHEW

42 TABLETS Three 14-day courses of treatment May take 1 to 4 days for full effect

Treats FREQUENT Heartburn!

24 HR





omeprazole tablet, delayed n	elease							
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:57896-659						
Route of Administration	ORAL							
Active Ingredient/Active	Moiety							
Ingr	edient Name		Basis of Strength		Strengt			
OMEPRAZOLE (UNII: KG60484QX) (OMEPRAZOLE - UNII:KG60	484QX9)	OMEPRAZ OLE		20 mg			
Inactive Ingredients								
	Ingredient Name	•			Strength			
ANHYDROUS LACTOSE (UNII: 3S	Y5LH9PMK)							
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)								
HYPROMELLOSE ACETATE SUC	CINATE 12070923 (3 MPA	.S) (UNII: 36BGF	DE889)					
FERRIC OXIDE RED (UNII: 1K09F3G675)								
FERRIC OXIDE YELLOW (UNII: EX43802MRT)								
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)								
2-AMINOETHANOL (UNII: 5KV861	14PT)							
METHYLCELLULOSE (1500 MPA								
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)								
SODIUM STARCH GLYCOLATE T	YPE A POTATO (UNII: 5856J	3G2A2)						
SODIUM STEARATE (UNII: QU7E2XA9TG)								
SODIUM STEARYL FUMARATE (U	JNII: 7CV7WJK4UI)							
SODIUM LAURYL SULFATE (UNII	368GB5141J)							
TRIETHYL CITRATE (UNII: 8Z96Q	XD6UM)							
TALC (UNII: 7SEV7J4R1U)								

FERROSOFERRIC OXIDE (UNII: XM0M87F357)							
BU	TYL ALCOHOL (UNII: 8PJ61P6TS3)					
SH	ELLAC (UNII: 46N	I107B71O)					
D							
۲r	oduct Chara	icteristics					
Color		BROWN (brownish pink)		ore		no score	
Shape		OVAL (biconvex)		e		12mm	
Flavor			Imprint Code			20	
Со	ntains						
Pa	ackaging						
	ackaging Item Code	Package Description	ſ	Marketing Start Date	M	arketing End Date	
#		Package Description 3 in 1 CARTON		-	M		
# 1 1	Item Code NDC:57896-659- 42			Date	M		
# 1	Item Code NDC:57896-659- 42 NDC:57896-659-	3 in 1 CARTON 40 in 1 BOTTLE; Type 0: Not a Combination		Date	M		
# 1	Item Code NDC:57896-659- 42 NDC:57896-659-	3 in 1 CARTON 40 in 1 BOTTLE; Type 0: Not a Combination		Date	M		
# 1	Item Code NDC:57896-659- 42 NDC:57896-659- 14	3 in 1 CARTON 40 in 1 BOTTLE; Type 0: Not a Combination		Date	M		
# 1	Item Code NDC:57896-659- 42 NDC:57896-659- 14	3 in 1 CARTON 40 in 1 BOTTLE; Type 0: Not a Combination Product	05/	Date			

Labeler - GERI-CARE PHARMACEUTICALS, CORP (611196254)

Establishment						
Name	Address	ID/FEI	Business Operations			
Sun Pharmaceutical Industries Limited		650445203	ANALYSIS(57896-659), MANUFACTURE(57896-659)			

Revised: 5/2024

GERI-CARE PHARMACEUTICALS, CORP