OMEPRAZOLE- omeprazole tablet, delayed release
Living Better Brands LLC

Omeprazole

Drug Facts

Active ingredient (in each tablet)

Omeprazole USP 20 mg

Purpose

Acid reducer

Use

- treats frequent heartburn (occurs ***2 or more*** 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 1-800-818-4555 weekdays.

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

PRINCIPAL DISPLAY PANEL - 20 mg Tablet Blister Pack Carton

COMPARE TO Prilosec OTC®*

NDC 21130-991-14

Signature
SELECT®

OMEPRAZOLE

24 HR | TREATS FREQUENT
HEARTBURN

Delayed-Release Tablets 20 mg

- Acid Reducer

SWALLOW-DO NOT CHEW

Actual Size

14 TABLETS

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

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Drug Facts (continued)

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KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

NOT FOR RESALE

5240692 DNH/DRUGS/NH/138

RD24127
S6792

Unvarnish Area: 50x20 mm

DISTRIBUTED BY:
BETTER LIVING BRANDS LLC
P.O. BOX 99
PLEASANTON, CA 94566-0009
1-888-723-3929
MADE IN INDIA

LOVE IT OR IT'S ON US!

BOX PAPER
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Safety Feature – Do not use if printed seal blister is open or torn.

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COMPARE TO *Prilosec OTC* ®*

NDC 21130-991-14

Signature SELECT

OMEPRAZOLE

24 HR | **TREATS FREQUENT HEARTBURN**

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• Acid Reducer

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OMEPRAZOLE
Delayed-Release Tablets
20 mg / Acid Reducer

Signature SELECT

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

omeprazole tablet, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-991
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9)	OMEPRAZOLE	20 mg

Inactive Ingredients	
Ingredient Name	**Strength**
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
HYPROMELLOSE ACETATE SUCCINATE 12070923 (3 MPA.S) (UNII: 36BGF0E889)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
2-AMINOETHANOL (UNII: 5KV86114PT)	
METHYLCELLULOSE (1500 MPA.S) (UNII: P0NTE48364)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
SODIUM STEARATE (UNII: QU7E2XA9TG)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
AMMONIA (UNII: 5138Q19F1X)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
BUTYL ALCOHOL (UNII: 8PJ61P6TS3)	
SHELLAC (UNII: 46N107B71O)	

Product Characteristics			
Color	BROWN (brownish pink)	**Score**	no score
Shape	OVAL (biconvex)	**Size**	12mm
Flavor		**Imprint Code**	20
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-991-44	1 in 1 CARTON	06/01/2019	
1		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:21130-991-28	2 in 1 CARTON	06/01/2019	
2		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA207891	06/01/2019	

Labeler - Living Better Brands LLC (009137209)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharmaceutical Industries Limited		650445203	ANALYSIS(21130-991) , MANUFACTURE(21130-991)

Revised: 8/2024 Living Better Brands LLC