OMEPRAZOLE- omeprazole tablet, delayed release BI-MART

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 1-800-935-6737 weekdays.

Distributed by: BI-MART Eugene, OR 97402

PRINCIPAL DISPLAY PANEL - 20 mg Tablet Bottle Carton

[†]Compare to the active ingredient in Prilosec OTC[®]

NDC 37835-001-42

BI-MART

See new warning information

Treats FREQUENT Heartburn!

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HR

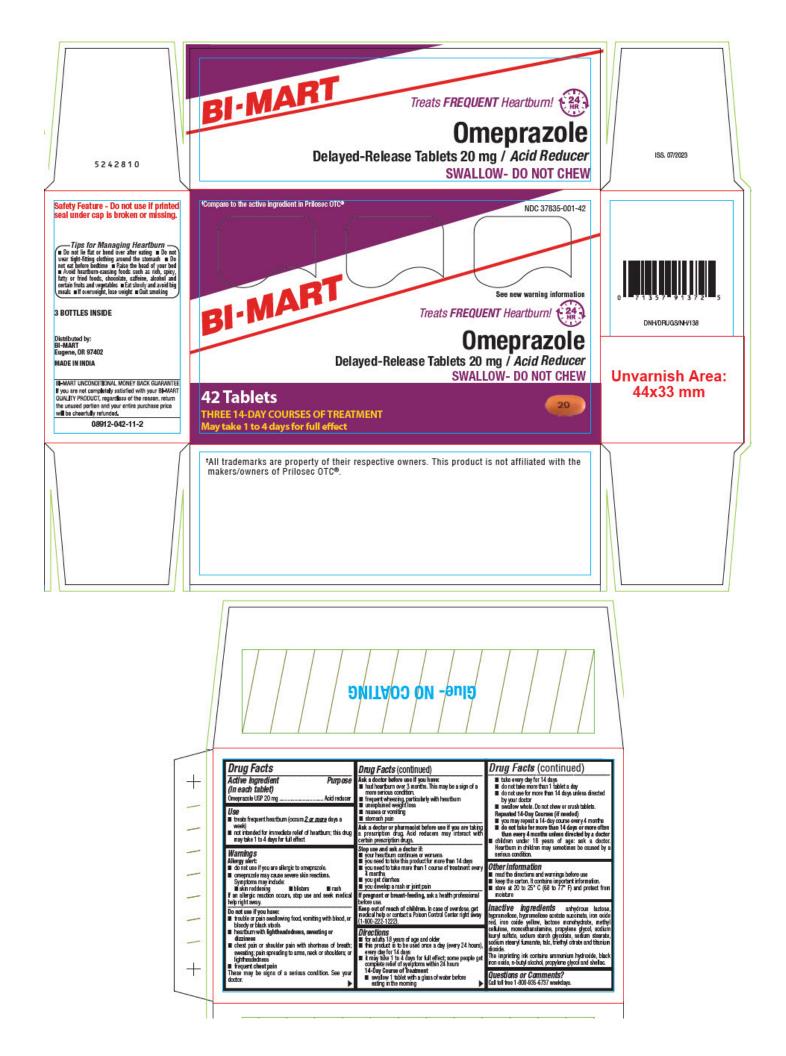
Omeprazole Delayed-Release Tablets 20 mg / Acid Reducer

SWALLOW- DO NOT CHEW

42 Tablets

THREE 14-DAY COURSES OF TREATMENT

May take 1 to 4 days for full effect



Glue-NO COATING

omeprazole tablet,	aciayca ie						
Product Informa	ation						
Product Type		HUMAN OTC DRUG	Item Code (S	ource)	IDC:3783	7835-001	
Route of Administ	ration	ORAL					
Active Ingredier	t/Active	Moiety					
-	Ingre	dient Name		Basis of Stre	ngth	Strength	
OMEPRAZOLE (UNII: K	G60484QX9)	OMEPRAZ OLE	-	20 mg			
Inactive Ingredi	ents						
mactive mgreat	ciics	Ingredient Na	me			Strength	
ANHYDROUS LACTOS	F (UNII: 35Y	•				Strength	
HYPROMELLOSE, UN							
-		INATE 12070923 (3 I	MM2/S) (UNII: 36BGI	F0E889)			
FERRIC OXIDE RED (U				,			
FERRIC OXIDE YELLO		· ·					
LACTOSE MONOHYD	RATE (UNII:	EWQ57Q8I5X)					
MONOETHANOLAMIN	IE (UNII: 5KV	36114PT)					
METHYLCELLULOSE	(1500 MPA.	S) (UNII: PONTE48364)					
PROPYLENE GLYCOL	(UNII: 6DC90	Q167V3)					
SODIUM STARCH GL	COLATE T	PE A POTATO (UNII: 5	856J3G2A2)				
SODIUM STEARATE (UNII: QU7E2>	(A9TG)					
SODIUM STEARYL FU	MARATE (U	NII: 7CV7WJK4UI)					
SODIUM LAURYL SUL	FATE (UNII:	368GB5141J)					
TRIETHYL CITRATE (U	JNII: 8Z96Q>	D6UM)					
TALC (UNII: 7SEV7J4R1	U)						
TITANIUM DIOXIDE (U	JNII: 15FIX9V	2JP)					
AMMONIA (UNII: 5138	Q19F1X)						
FERROSOFERRIC OX	IDE (UNII: XM	IOM87F357)					
BUTYL ALCOHOL (UN	II: 8PJ61P6TS	3)					
SHELLAC (UNII: 46N10	7B710)						
Product Charact	teristics						

Shape		OVAL (biconvex)	Size	12mm					
Flavor			Imprint Code	20					
Contains									
_									
Packaging									
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date					
1	NDC:37835-001- 42	3 in 1 CARTON	09/30/2019						
1		14 in 1 BOTTLE; Type 0: Not a Combination Product							
2	NDC:37835-001- 14	1 in 1 CARTON	09/30/2019						
2		14 in 1 BOTTLE; Type 0: Not a Combination Product							
Μ	larketing l	nformation							
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
	IDA	ANDA207891	09/30/2019						

Labeler - BI-MART (027630078)

Establishment									
Name	Address	ID/FEI	Business Operations						
Sun Pharmaceutical Industries Limited		650445203	ANALYSIS(37835-001), MANUFACTURE(37835-001)						

Revised: 8/2023

BI-MART