EQUATE OMEPRAZOLE- omeprazole tablet, delayed release Praxis, LLC

Wal-Mart Omeprazole Delayed Release Tablets 20 mg Drug Facts

Active ingredient (in each tablet)

Omeprazole 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

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, crush, or suck 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 20-25°C (68-77°F) and protect from moisture

Inactive ingredients

benzyl alcohol, carmine, carnauba wax, FD&C blue #2/indigo carmine aluminum lake, flavor, hypromellose, hypromellose acetate succinate, lactose monohydrate, menthol, modified starch, monoethanolamine, polyethylene glycol 3350, sodium lauryl sulfate, sodium starch glycolate, sodium stearate, sodium stearyl fumarate, sucralose, talc, titanium dioxide, triacetin, triethyl citrate

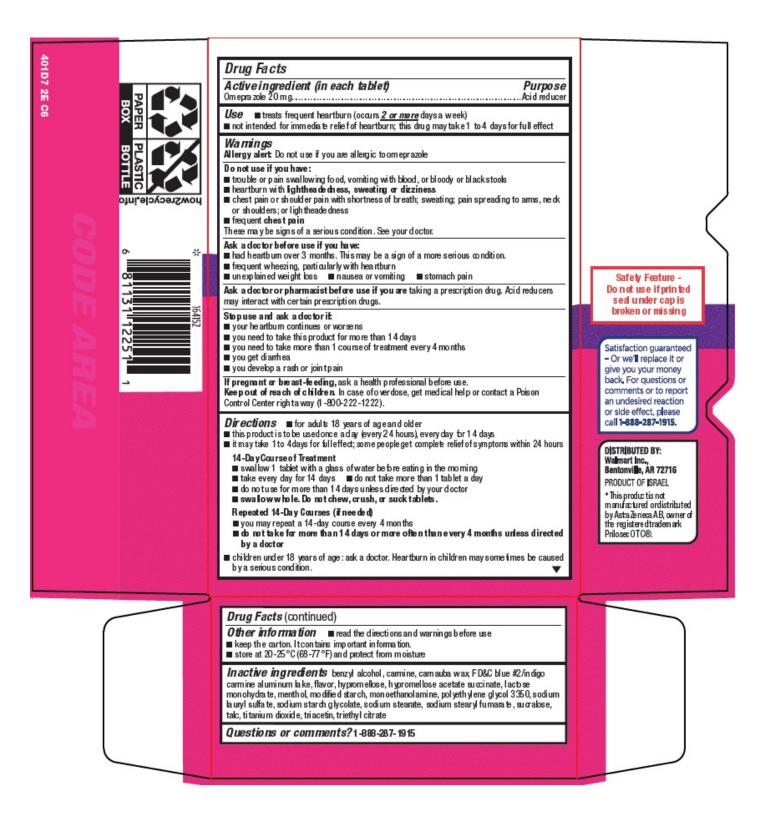
Questions or comments?

1-888-287-1915

Package/Label Principal Display Panel

3 PACK THREE 14-DAY COURSES OF TREATMENT equate [™] Compare to Prilosec OTC[®] Omeprazole Delayed Release Tablets 20 mg Acid Reducer **Treats Frequent Heartburn!** 24 HR Actual Size SWALLOW - DO NOT CHEW WILDBERRY MINT COATED TABLET 20 mg 42 TABLETS Three 14-day courses of treatment May take 1 to 4 days for full effect 3 Bottles Inside See Current Drug Facts





EQUATE OMEPRAZOLE

omeprazole tablet, delayed release

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59368-327		
Route of Administration	ORAL				

Active Ingredient/A	ctive Moiety				
	Ingredient Na	ame	Basis of Strengt	th Strength	
OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9) OMEPRAZOLE			OMEPRAZ OLE	20 mg	
Inactive Ingredient	s				
	Ingred	lient Name		Strength	
BENZYL ALCOHOL (UNII: L	KG8494WBH)				
CARNAUBA WAX (UNII: R12	2CBM0EIZ)				
FD&C BLUE NO. 2 (UNII: L	.06K8R7DQK)				
HYPROMELLOSE, UNSPEC	CIFIED (UNII: 3NX)	N29V3WO)			
LACTOSE MONOHYDRATI	E (UNII: EWQ57Q8I	5X)			
MENTHOL, UNSPECIFIED	FORM (UNII: L7T)	LOEIP3A)			
MONOETHANOLAMINE (U	NII: 5KV86114PT)				
POLYETHYLENE GLYCOL	3350 (UNII: G2M7	P15E5P)			
SODIUM LAURYL SULFAT	E (UNII: 368GB514	1J)			
SODIUM STEARATE (UNII:					
SODIUM STEARYL FUMAR		J K4UI)			
SUCRALOSE (UNII: 96K6UQ	(3Z D4)				
TALC (UNII: 7SEV7J4R1U)					
TITANIUM DIOXIDE (UNII:	15FIX9V2JP)				
TRIACETIN (UNII: XHX3C3X	673)				
TRIETHYL CITRATE (UNII:	8Z96QXD6UM)				
Product Characteri	stics				
Color	purple	Score	no scor	e	
Shape	OVAL	Size	12mm	12mm	
Flavor	BERRY	Imprint Code	20		
Contains					
Packaging					

	3 3				
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:59368-327- 01	1 in 1 CARTON	09/02/2015		
1		14 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:59368-327- 02	3 in 1 CARTON	09/02/2015		
2		14 in 1 BOTTLE; Type 0: Not a Combination Product			
Marketing Information					
	Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
NE	DA	NDA022032	09/02/2015		

Labeler - Praxis, LLC (016329513)

Establishment				
Name	Address	ID/FEI	Business Operations	
Praxis, LLC		016329513	manufacture(59368-327) , label(59368-327) , pack(59368-327)	

Revised: 1/2023

Praxis, LLC