#### BASIC CARE OMEPRAZOLE- omeprazole tablet, delayed release Amazon.com Services LLC

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#### **Amazon Omeprazole 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? 1-800-719-9260

### Package/Label Principal Display Panel

Acid Reducer Compare to Prilosec OTC® FDA Approved Omeprazole Delayed Release Tablets 20 mg Acid Reducer Treats Frequent Heartburn! 24 HR Wildberry Mint Coated Tablet See current Drug Facts SWALLOW - DO NOT CHEW actual size 14 TABLETS One 14-day course of treatment May take 1 to 4 days for full effect



BASIC CARE OMEPRA	AZOLE								
omeprazole tablet, delayed re	lease								
Product Information									
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:72288-401					
Route of Administration	ORAL								
Active Ingredient/Active Moiety									
Ingredient Name			<b>Basis of Strength</b>		Strength				
OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9)			OMEPRAZ OLE		20 mg				

Inactive Ingredients				
Ingredient Name	Strength			
BENZYL ALCOHOL (UNII: LKG8494WBH)				
CARNAUBA WAX (UNII: R12CBM0EIZ)				
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)				
MONOETHANOLAMINE (UNII: 5KV86114PT)				
POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
SODIUM STEARATE (UNII: QU7E2XA9TG)				
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
TALC (UNII: 7SEV7J4R1U)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
TRIACETIN (UNII: XHX3C3X673)				
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)				

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#### Packaging

#	ltem Code	Package Description	ackage Description Marketing Start Date					
1	NDC:72288-401- 03	3 in 1 CARTON	10/15/2020					
1		14 in 1 BOTTLE; Type 0: Not a Combination Product						
2	NDC:72288-401- 01	1 in 1 CARTON	04/12/2021					
2		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				
NE	A	NDA022032	10/15/2020					

# Labeler - Amazon.com Services LLC (128990418)