#### OMEPRAZOLE- omeprazole tablet, orally disintegrating, delayed release Rite Aid Corporation

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# **Rite Aid Corporation 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 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

- take 1 tablet before eating in the morning
- do not crush or chew tablets
- place the tablet on tongue; tablet disintegrates, with or without water. The tablets can also be swallowed whole with water.
- 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
- do not take this medicine with alcohol

# Repeated 14-Day Course (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 before use. 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); keep product out of high heat and moisture

#### Inactive ingredients

amino methacrylate copolymer, ascorbic acid, cetyl alcohol, colloidal silicon dioxide, crospovidone, ferric oxide, flavor, hypromellose, hypromellose phthalate, maize maltodextrin, mannitol, microcrystalline cellulose, propylene glycol, silicon dioxide, sodium stearate, sodium stearyl fumarate, sorbitol, sucralose, sugar spheres, talc, titanium dioxide, triethyl citrate

#### **Questions or comments?**

1-800-719-9260: weekdays 7:30 AM to 5:00 PM EST

# Package/Label Principal Display Panel MELTS IN YOUR MOUTH DISSLOVES WITHOUT WATER Compare to Prilosec OTC<sup>®</sup> MELTech<sup>™</sup> Melts In Your Mouth OMEPRAZOLE Delayed Release Orally Disintegrating Tablets 20 mg ACID REDUCER TREATS FREQUENT HEARTBURN! Strawberry Flavor ACTUAL SIZE 24HR 42 TABLETS Three 14-day courses of treatment May take 1 to 4 days for full effect





If you're not satisfied, we'll happily refund your money

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1Y955 83 C3

MADE IN ISRAEL

#### OMEPRAZOLE omeprazole tablet, orally disintegrating, delayed release **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:11822-1040 **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 DIMETHYLAMINOETHYL METHACRYLATE - BUTYL METHACRYLATE - METHYL METHACRYLATE COPOLYMER (UNII: 905HNO1SIH) ASCORBIC ACID (UNII: PQ6CK8PD0R) CETYL ALCOHOL (UNII: 936JST6JCN) SILICON DIOXIDE (UNII: ETJ7Z6XBU4) CROSPOVIDONE (15 MPA.S AT 5%) (UNII: 68401960MK) FERRIC OXIDE RED (UNII: 1K09F3G675) HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) MANNITOL (UNII: 30WL53L36A) MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) **SODIUM STEARATE** (UNII: QU7E2XA9TG) SODIUM STEARYL FUMARATE (UNII: 7CV7WK4UI) SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3ZD4) TALC (UNII: 7SEV7J4R1U) TITANIUM DIOXIDE (UNII: 15FIX9V2JP) TRIETHYL CITRATE (UNII: 8Z96QXD6UM) **Product Characteristics** Color RED (reddish) Score no score ROUND 9mm Shape Size **STRAWBERRY** 20 Flavor Imprint Code Contains Packaging **Marketing Start Marketing End Item Code Package Description** # Date Date - NDC:11822- 2 := 1 CARTON 01/00/2010

1	1040-1	S III I CARTON	02/23/2010	
1		14 in 1 CARTON		
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822- 1040-2	14 in 1 CARTON	03/13/2019	
2		1 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
NE	DA	NDA209400	03/29/2018	

Labeler - Rite Aid Corporation (014578892)

Revised: 7/2023

Rite Aid Corporation