

**OMEPRAZOLE- omeprazole tablet, delayed release**  
**Rite Aid Corporation**

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**Rite Aid Corporation Omeprazole Delayed Release Tablets 20 mg Drug Facts**

**Active ingredient (in each tablet)**

Omeprazole 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 20-25°C (68-77°F) and protect from moisture

## Inactive ingredients

carnauba wax, ferric oxide red, ferric oxide yellow, hypromellose, hypromellose acetate succinate, lactose monohydrate, monoethanolamine, propylene glycol, sodium lauryl sulfate, sodium starch glycolate, sodium stearate, sodium stearyl fumarate, talc, titanium dioxide, triethyl citrate

## Questions or comments?

1-800-719-9260

## Principal Display Panel

TREATS FREQUENT HEARTBURN!

Compare to Prilosec OTC<sup>®</sup>

OMEPRAZOLE

Delayed Release Tablets, 20 mg

ACID REDUCER

ACTUAL SIZE

24HR

42 TABLETS

THREE 14-DAY COURSES OF TREATMENT

MAY TAKE 1 TO 4 DAYS FOR FULL EFFECT

**Drug Facts**

**Active ingredient (in each tablet)**  
Omeprazole 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 redness
  - 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, swelling, 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 vomiting, particularly with heartburn
- unexplained weight loss
- nausea or vomiting
- stomach pain

**Drug Facts (continued)**

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), everyday 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**

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- take every day for 14 days
- do not take more than 1 tablet a day
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**Repeated 14-Day Courses (if needed)**

- you may repeat a 14-day course every 4 months
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- 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 cap on. It contains important information.
- store at 20-25°C (68-77°F) and protect from moisture

**Inactive ingredients**

carnauba wax, ferric oxide red, ferric oxide yellow, hypromellose, hypromellose succinate succinate, lactose monohydrate, monoethanolamine, propylene glycol, sodium lauryl sulfate, sodium starch glycolate, sodium stearate, sodium stearyl fumarate, talc, titanium dioxide, triethyl citrate

**Questions or comments?**  
1-800-719-9260

\*This product is not manufacturer or distributed by Pfizer Inc. Camak, distributor of Prilosec OTC.

**3 BOTTLES INSIDE**

**Safety Feature** - Do not use if printed seal under cap is broken or missing.

**Drug Facts (continued)**

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:

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# OMEPRAZOLE

omeprazole tablet, delayed release

## Product Information

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:11822-0915 |
| <b>Route of Administration</b> | ORAL           |                           |                |

## Active Ingredient/Active Moiety

| <b>Ingredient Name</b>                                       | <b>Basis of Strength</b> | <b>Strength</b> |
|--|--------------------------|-----------------|
| OMEPRAZOLE (UNII: KG60484QX9) (OMEPRAZOLE - UNII:KG60484QX9) | OMEPRAZOLE               | 20 mg           |

## Inactive Ingredients

| <b>Ingredient Name</b>   | <b>Strength</b> |
|--|-----------------|
| CARNAUBA WAX (UNII: R12CBM0EIZ)                                      |                 |
| FERRIC OXIDE RED (UNII: 1K09F3G675)                                  |                 |
| FERRIC OXIDE YELLOW (UNII: EX438O2MRT)                               |                 |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)                         |                 |
| LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)                               |                 |
| MONOETHANOLAMINE (UNII: 5KV86114PT)                                  |                 |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3)                                  |                 |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J)                             |                 |
| SODIUM STEARATE (UNII: QU7E2XA9TG)                                   |                 |
| SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)                           |                 |
| TALC (UNII: 7SEV7J4R1U)  |                 |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP)                                  |                 |
| TRIETHYL CITRATE (UNII: 8Z96QXD6UM)                                  |                 |
| HYPROMELLOSE ACETATE SUCCINATE 12070923 (3 MM2/S) (UNII: 36BGF0E889) |                 |

## Product Characteristics

|                 |       |                     |          |
|-----------------|-------|---------------------|----------|
| <b>Color</b>    | BROWN | <b>Score</b>        | no score |
| <b>Shape</b>    | OVAL  | <b>Size</b>         | 12mm     |
| <b>Flavor</b>   |       | <b>Imprint Code</b> | 20       |
| <b>Contains</b> |       |                     |          |

## Packaging

| # | Item Code        | Package Description                                    | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:11822-0915-2 | 14 in 1 CARTON   | 02/25/2008           | 10/01/2018         |
| 1 |                  | 1 in 1 BLISTER PACK; Type 0: Not a Combination Product |                      |                    |
| 2 | NDC:11822-0915-1 | 2 in 1 CARTON  | 02/27/2008           | 10/01/2018         |
| 2 |                  | 14 in 1 CARTON   |                      |                    |

|   |                  |  |            |            |
|---|------------------|--|------------|------------|
| 2 |                  | 1 in 1 BLISTER PACK; Type 0: Not a Combination Product |            |            |
| 3 | NDC:11822-0915-3 | 3 in 1 CARTON  | 02/27/2008 | 12/01/2018 |
| 3 |                  | 14 in 1 CARTON   |            |            |
| 3 |                  | 1 in 1 BLISTER PACK; Type 0: Not a Combination Product |            |            |
| 4 | NDC:11822-0915-5 | 1 in 1 CARTON  | 12/27/2011 |            |
| 4 |                  | 14 in 1 BOTTLE; Type 0: Not a Combination Product      |            |            |
| 5 | NDC:11822-0915-4 | 2 in 1 CARTON  | 12/27/2011 |            |
| 5 |                  | 14 in 1 BOTTLE; Type 0: Not a Combination Product      |            |            |
| 6 | NDC:11822-0915-6 | 3 in 1 CARTON  | 12/27/2011 |            |
| 6 |                  | 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 |
|--------------------|--|----------------------|--------------------|
| NDA                | NDA022032                                | 02/25/2008           |                    |

**Labeler** - Rite Aid Corporation (014578892)

Revised: 11/2023

Rite Aid Corporation