

**OMEPRAZOLE- omeprazole tablet, delayed release**  
**PLD Acquisitions LLC DBA Avéma Pharma Solutions**

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**OMEPRAZOLE MAGNESIUM DELAYED-RELEASE TABLETS**

***Active ingredient (in each tablet)***

Omeprazole delayed-release tablet 20 mg  
(equivalent to 20.6 mg omeprazole magnesium)

***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.

***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 everyday 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***

acetyl tributyl citrate, colloidal silicon dioxide, corn starch, croscarmellose sodium, dextrose, FD&C Blue #2, FD&C Red #40, flavor, hydroxypropyl cellulose, hypromellose 2910, magnesium stearate, methacrylic acid copolymer type C, microcrystalline cellulose, polyethylene glycol 400, polyethylene glycol 3350, polyethylene glycol 8000, polyvinyl alcohol, propylene glycol, sucrose, talc, titanium dioxide, triacetin, triethyl citrate

### ***Tips for Managing Heartburn***

- Do not lie flat or bend over after eating
- Do not wear tight-fitting clothing around the stomach
- Do not eat before bedtime
- Raise the head of your bed
- Avoid heartburn-causing foods such as rich, spicy, fatty or fried foods, chocolate, caffeine, alcohol and certain fruits and vegetables
- Eat slowly and avoid big meals
- If overweight, lose weight
- Quit smoking

### **Principal display panel**

†Compare to the active ingredient in Prilosec® OTC.

24-hour

Omeprazole

Delayed-release tablets 20 mg

Acid reducer

Treats FREQUENT heartburn 24 HR

May take 1 to 4 days for full effect

Tablets

One 14-day course of treatment

Coated with Wildberry Flavor

†This product is not affiliated with, manufactured by or produced by the makers or owners of Prilosec OTC®.

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.**

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**

Manufactured by:

Avema Pharma Solutions  
 10400 NW 29th Terrace  
 Miami, FL 33172

**Package label**

**Drug Facts (continued)**

■ store at 20° to 25°C (68° to 77°F)

**Inactive ingredients** acetyltrihutyl citrate, colloidal silicon dioxide, corn starch, croscarmellose sodium, hydroxypropyl cellulose, hypromellose, 50:10, magnesium stearate, methacrylic acid copolymer type C, microcrystalline cellulose, polyethylene glycol 400, polyethylene glycol 3350, polyethylene glycol 8000, polyvinyl alcohol, red iron oxide, sucrose, talc, titanium dioxide, triethyl citrate

**Questions or comments?**  
 Call 1-877-758-3838, Monday-Friday 9AM-5PM EST

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**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.**

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**


NDC 63548-297-14  
 (Compare to the active ingredient in Prilosec OTC\*)

See current Drug Facts

**24-hour**  
**Omeprazole**  
 Delayed-release tablets, 20 mg  
 Acid reducer

*Treats FREQUENT heartburn!*  
 May take 1 to 4 days for full effect

14 Tablets  
 One 14-day course of treatment



Actual Size

63548-297-14  
 (Compare to the active ingredient in Prilosec OTC\*)

See current Drug Facts

**24-hour**  
**Omeprazole**  
 Delayed-release tablets, 20 mg  
 Acid reducer

*Treats FREQUENT heartburn!*  
 May take 1 to 4 days for full effect  
 SWALLOW-DO NOT CHEW

14 Tablets  
 One 14-day course of treatment

**Drug Facts**

**Active ingredient (in each tablet)**  
 (Omeprazole delayed-release tablet 20 mg (equivalent to 20.6 mg omeprazole magnesium))

**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, ■ hives, ■ 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, back 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
- dizziness or weakness
- stomach pain

Manufactured by:  
 Avema Pharma Solutions  
 10400 NW 29th Terrace  
 Miami, FL 33172

1 BOTTLE INSIDE



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Rev. 12/23  
 PLD-8649A  
 FC036925

Lot No:  
 Exp. Date:

**Avema Pharma Solution Acid Reducer**

**OMEPRAZOLE**

omeprazole tablet, delayed release

**Product Information**

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:63548-7370 |
| <b>Route of Administration</b> | ORAL           |                           |                |

**Active Ingredient/Active Moiety**

| Ingredient Name   | Basis of Strength | Strength |
|---|-------------------|----------|
| <b>OMEPRAZOLE MAGNESIUM</b> (UNII: 426QFE7XLK) (OMEPRAZOLE - UNII:KG60484QX9) | OMEPRAZOLE        | 20.6 mg  |

**Inactive Ingredients**

| Ingredient Name                                  | Strength |
|--|----------|
| <b>ACETYLTRIBUTYL CITRATE</b> (UNII: OZBX0N59RZ) |          |
| <b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XB4)         |          |
| <b>STARCH, CORN</b> (UNII: O8232NY3S)            |          |
| <b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)   |          |
| <b>DEXTROSE</b> (UNII: IY9XDZ35W2)               |          |
| <b>FD&amp;C BLUE NO. 2</b> (UNII: L06K8R7DQK)    |          |

|   |
|---|
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)                           |
| <b>HYDROXYPROPYL CELLULOSE, UNSPECIFIED</b> (UNII: 9XZ8H6N6OH)          |
| <b>HYPROMELLOSE 2910 (15 MPA.S)</b> (UNII: 36SFW2JZ0W)                  |
| <b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)                            |
| <b>METHACRYLIC ACID AND ETHYL ACRYLATE COPOLYMER</b> (UNII: NX76LV5T8J) |
| <b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)                       |
| <b>POLYETHYLENE GLYCOL 3350</b> (UNII: G2M7P15E5P)                      |
| <b>POLYETHYLENE GLYCOL 8000</b> (UNII: Q662QK8M3B)                      |
| <b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)                |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)                              |
| <b>SUCROSE</b> (UNII: C151H8M554)                                       |
| <b>TALC</b> (UNII: 7SEV7J4R1U)  |
| <b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)                              |
| <b>TRIACETIN</b> (UNII: XHX3C3X673)                                     |
| <b>TRIETHYL CITRATE</b> (UNII: 8Z96QXD6UM)                              |
| <b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)                    |

### Product Characteristics

|                 |                 |                     |          |
|-----------------|-----------------|---------------------|----------|
| <b>Color</b>    | red (brown)     | <b>Score</b>        | no score |
| <b>Shape</b>    | OVAL            | <b>Size</b>         | 13mm     |
| <b>Flavor</b>   | BERRY (Wlberry) | <b>Imprint Code</b> | 0        |
| <b>Contains</b> |                 |                     |          |

### Packaging

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:63548-7370-4 | 1 in 1 BOX   | 10/31/2024           |                    |
| 1 |                  | 14 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product |                      |                    |
| 2 | NDC:63548-7370-8 | 1 in 1 BOX   | 10/31/2024           |                    |
| 2 |                  | 28 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product |                      |                    |
| 3 | NDC:63548-7370-2 | 1 in 1 BOX   | 10/31/2024           |                    |
| 3 |                  | 42 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| ANDA               | ANDA206582                               | 10/31/2024           |                    |

**Labeler** - PLD Acquisitions LLC DBA Avéma Pharma Solutions (804087794)

