GOOD SENSE OMEPRAZOLE- omeprazole tablet, delayed release Praxis, LLC

Perrigo 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.
- 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, 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

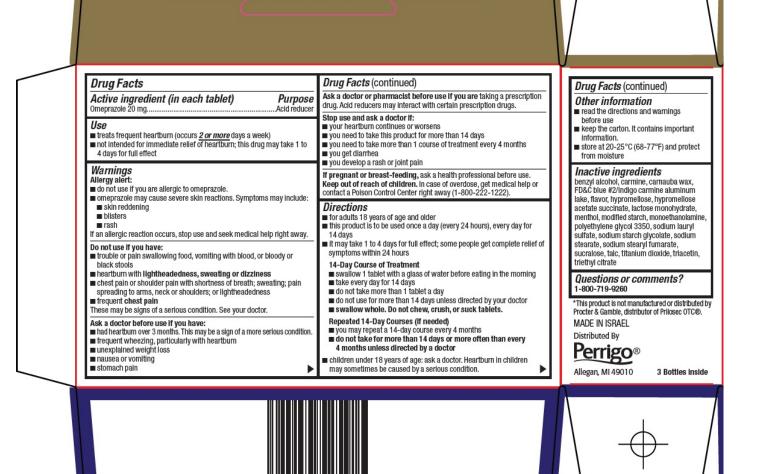
Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

GOODSENSE ® FDA APPROVED Treats Frequent Heartburn! 24 HR Omeprazole Delayed Release Tablets 20 mg Acid Reducer 20 Wildberry Mint Coated Tablet Compare to Prilosec OTC® Actual Size SWALLOW - DO NOT CHEW 42 Tablets Three 14-Day Courses of Treatment May Take 1 to 4 Days For Full Effect 3 Bottles Inside





| GOOD SENSE OMEPRAZOLE | | | | | |
|--|----------------|--------------------|--------------------------|---------------|----------|
| omeprazole tablet, delayed re | elease | | | | |
| | | | | | |
| Product Information | | | | | |
| Product Type | HUMAN OTC DRUG | Item Code (Source) | | NDC:59368-320 | |
| Route of Administration | ORAL | | | | |
| | | | | | |
| | | | | | |
| Active Ingredient/Active Moiety | | | | | |
| Ingredient Name | | | Basis of Strength | | 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) | | | | |
| HYPROMELLOSE ACETATE SUCCINATE, UNSPECIFIED (UNII: A7ZHS2RJ34) | | | | |
| 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) | | | | |

| Product Characteristics | | | | |
|-------------------------|--------|--------------|----------|--|
| Color | purple | Score | no score | |
| Shape | OVAL | Size | 12mm | |
| Flavor | BERRY | Imprint Code | 20 | |
| Contains | | | | |

Packaging

| # | ltem Code | | | Marketing End Date | |
|-----------------------|----------------------|---|-----------------|-----------------------|--|
| 1 | NDC:59368-320- 02 | 3 in 1 CARTON 05/22/2019 | | | |
| 1 | | 14 in 1 BOTTLE; Type 0: Not a Combination Product | | | |
| 2 | NDC:59368-320- 01 | 14 in 1 BOTTLE; Type 0: Not a Combination Product 05/22/2019 | | | |
| | | | | | |
| Marketing Information | | | | | |
| | Marketing | Application Number or Monograph | Marketing Start | Marketing End | |

| Marketing | Application Number or Monograph | Marketing Start | Marketing End |
|-----------|---------------------------------|-----------------|---------------|
| Category | Citation | Date | Date |
| NDA | NDA022032 | 05/22/2019 | |

Labeler - Praxis, LLC (016329513)

| Establishment | | | |
|---------------|---------|-----------|---|
| Name | Address | ID/FEI | Business Operations |
| Praxis, LLC | | 016329513 | manufacture(59368-320) , label(59368-320) , pack(59368-320) |

Revised: 1/2023