## ACID REDUCER - omeprazole tablet, delayed release Best Choice

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## 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

#### **Drug Facts**

#### Active ingredient (in each tablet)

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

## 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

## 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:

- warfarin, clopidogrel or cilostazol (blood-thinning medicines)
- prescription antifungal or anti-yeast medicines
- diazepam (anxiety medicine)
- digoxin (heart medicine)
- tacrolimus or mycophenolate mofetil (immune system medicines)
- prescription antiretrovirals (medicines for HIV infection)
- methotrexate (arthritis medicine)

## 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 (1-800-222-1222) 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 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° to 25°C (68° to 77° F) and protect from moisture

## Inactive ingredients

crospovidone, glyceryl monostearate, hydroxypropyl cellulose, hypromellose, magnesium stearate, methacrylic acid copolymer dispersion, microcrystalline cellulose, polyethylene glycol, polysorbate 80, red iron oxide, silicified microcrystalline cellulose, sodium hydroxide, sodium stearyl fumarate, sugar spheres [which contains liquid glucose, starch (maize) and sucrose], talc, titanium dioxide, triethyl citrate and yellow iron oxide.

#### **Questions?**

Call 1-855-274-4122

PROUDLY DISTRIBUTED BY: VALU MERCHANDISERS, CO. **5000 KANSAS AVE. KANSAS CITY, KS 66106** MADE IN INDIA

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Code TS/DRUGS/22/2009

## PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg (14 Tablet Bottle)

NDC 63941-006-05 Acid Reducer **Best OMEPRAZOLE Choice® DELAYED-RELEASE TABLETS 20 mg** 

24 HR

14 TABLETS **Treats Frequent Heartburn** One 14-day course of treatment May take 1 to 4 days for full

effect



# **Top Ply**

# **Bottom Ply**

■ 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-leeding, 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 (1-800-222-1222) 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 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 warmings before use ■ keep the carrior, it contains important information. ■ store at 20° to 25° C (88° to 77°F) and protect from moisture inactive ingredients crospovidone, glyceryl monostearate, hydroxypropyl cellulose, hyprometiose, magnesium stearate, methacrylic acid copolymer dispersion, microcrystalline cellulose, polyethylene glycol, polysorbate 80, red iron oxide, silicified microcrystalline cellulose, sodium hydroxide, sodium stearyl fumarate, sugar spheres [which contains liquid glucose, starch (maize) and sucrose], talc, titanium dioxide, triethyl citrate and yellow iron oxide. Questions? Call 1-855-274-4122

## PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg Container Carton Label

NDC 63941-006-05

Best Choice®

COMPARE TO THE ACTIVE INGREDIENT IN PRILOSEC OTC®\* Acid Reducer OMEPRAZOLE

DELAYED-RELEASE
TABLETS 20 mg
24 HR
Treats FREQUENTS Heartburn
14 TABLETS
One 14-day course of treatment
May take 1 to 4 days for full effect



## **ACID REDUCER**

omeprazole tablet, delayed release

Product Information				
<b>Product Type</b>	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-006	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				

Ingredient Name	Basis of Strength	Strength
OMEPRAZOLE MAGNESIUM (UNII: 426QFE7XLK) (OMEPRAZOLE - UNII: KG60484QX9)	OMEPRAZOLE	20 mg

Inactive Ingredients	
Ingredient Name	Strength
CROSPOVIDONE (35 .MU.M) (UNII: 40UAA97IT9)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
HYDROXYPROPYL CELLULOSE (90000 WAMW) (UNII: UKE75GEA7F)	
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (1:1) TYPE A (UNII: NX76LV5T8J)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
HYDROXYPROPYL CELLULOSE (45000 WAMW) (UNII: 8VAB711C5E)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	

Product Characteristics			
Color	PINK	Score	no score
Shape	RECTANGLE (Oblong)	Size	14mm
Flavor		Imprint Code	Z;69
Contains			

Packaging			
Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:63941-006- 05	1 in 1 CARTON	06/06/2018	
	14 in 1 BOTTLE; Type 0: Not a Combination Product		
NDC:63941-006- 62	3 in 1 CARTON	06/06/2018	
	14 in 1 BOTTLE; Type 0: Not a Combination Product		
	Item Code NDC:63941-006-05 NDC:63941-006-	Item Code  Package Description  NDC:63941-006- 05  1 in 1 CARTON  14 in 1 BOTTLE; Type 0: Not a Combination Product  NDC:63941-006- 62  3 in 1 CARTON  14 in 1 BOTTLE; Type 0: Not a Combination	Item CodePackage DescriptionMarketing Start DateNDC:63941-006-051 in 1 CARTON06/06/201814 in 1 BOTTLE; Type 0: Not a Combination Product06/06/2018NDC:63941-006-623 in 1 CARTON06/06/201814 in 1 BOTTLE; Type 0: Not a Combination

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206877	06/06/2018	

## Labeler - Best Choice (868703513)

## Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(63941-006), MANUFACTURE(63941-006)

Revised: 12/2023 Best Choice