ACID REDUCER - omeprazole tablet, delayed release Family Dollar, Inc.

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 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 (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**

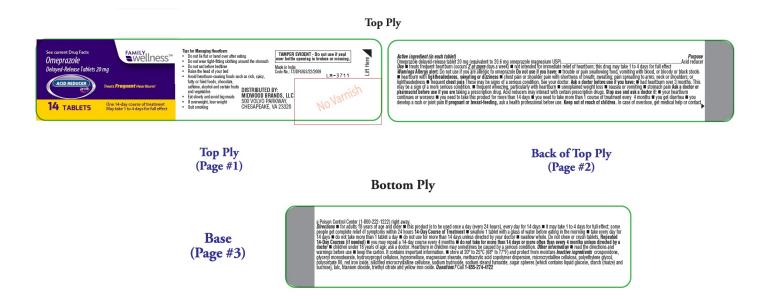
DISTRIBUTED BY:

MDWOOD BRANDS, LLC

500 VOLVO PARKWAY, CHESAPEAKE, VA 23320

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg (14 Tablets Container Label)

FAMILY
WellnessTM
See current Drug Facts
omeprazole
Delayed-Release Tablets 20 mg
ACID REDUCER
24 HR
Treats Frequent Heartburn!
One 14-day course of treatment
May take 1 to 4 days for full effect
14 TABLETS



PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg (42 Tablets Container Carton Label)

FAMILY
wellnessTM
COMPARE TO PRILOSEC OTC®
ACTIVE INGREDIENT*
ACID REDUCER
See current Drug Facts

Omeprazole Delayed-Release Tablets 20 mg

Treats **Frequent** Heartburn1! **24 HR OTC**

network

42 TABLETS

Three 14-day courses of treatment. May take 1 to 4 days for full effect



ACID REDUCER

omeprazole tablet, delayed release

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-037
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
OMEPRAZOLE MAGNESIUM (UNII: 4260FE7XLK) (OMEPRAZOLE - UNII:KG604840X9)	OMEPRAZOLE	20 mg		

Inactive Ingredients		
Ingredient Name	Strength	
CROSPO VIDO NE (35 .MU.M) (UNII: 40 UAA97IT9)		
GLYCERYL MONOSTEARATE (UNII: 230 O U9 XXE4)		
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) MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U) POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95) POLYSORBATE 80 (UNII: 6OZP39ZG8H) FERRIC OXIDE RED (UNII: 1K09F3G675) SILICON DIO XIDE (UNII: ETJ7Z6 XBU4) SODIUM HYDRO XIDE (UNII: 55X04QC32I) SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI) **DEXTROSE, UNSPECIFIED FORM** (UNII: IY9 XDZ35W2) STARCH, CORN (UNII: O8232NY3SJ) SUCROSE (UNII: C151H8M554) TALC (UNII: 7SEV7J4R1U) TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) TRIETHYL CITRATE (UNII: 8Z96QXD6UM) FERRIC OXIDE YELLOW (UNII: EX438O2MRT) HYDROXYPROPYL CELLULOSE (45000 WAMW) (UNII: 8VAB711C5E) HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0 WZ8 WG20 P6)

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

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:55319-037-05	1 in 1 CARTON	06/06/2018		
1		14 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:55319-037-62	3 in 1 CARTON	06/06/2018		
2		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
ANDA	ANDA206877	06/06/2018	

Labeler - Family Dollar, Inc. (024472631)

$Registrant - {\it Aurohealth LLC} \, (078728447)$

Establishment				
Name	Address	ID/FEI	Business Operations	
Aurobindo Pharma Limited		650381903	ANALYSIS(55319-037), MANUFACTURE(55319-037)	

Revised: 10/2019 Family Dollar, Inc.