ACID REDUCER - omeprazole tablet, delayed release WALGREEN CO.

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 *<u>2</u> 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 and ethyl acrylate 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**

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MADE IN INDIA

Code: TS/DRUGS/22/2009

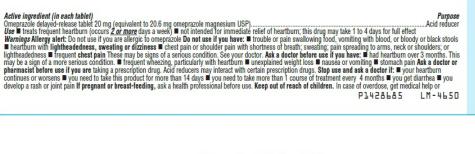
PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 20 mg (14 Tablet Bottle)

Walgreens NDC 0363-1008-05 Omeprazole DELAYED-RELEASE TABLETS 20 mg / ACID REDUCER • 24 HR • Treats frequent heartburn! 14 TABLETS ONE 14-DAY COURSE OF TREATMENT

MAY TAKE 1 TO 4 DAYS FOR FULL EFFECT







Back of Top Ply (Page #2)

Base (Page #3) contact a Poison Control Center (1-800-222-1222) right away. Directions T for adults 18 years of age and older T this product is to be used once a day (every 24 hours), every day for 14 days T in may take 1 to 4 days for full effect; some people get complete relief of symptoms within 24 hours 14-Day Course of Treatment S wallow 1 tablet with a glass of water before eating in the moming T take every day for 14 days T on take more than 1 tablet ad 9 do not use for more than 14 days and so for more than 14 days under software before eating in the moming T take every day for 14 days T on take more than 1 tablet ad 40 do not acke more than 1 tablet ad a dotor. Restruction in children may sometimes be caused by a endous condition. Other information T read the directions and warnings before use I keep the carton. It contains important information. Store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important information. Store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important information. Store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important information. Store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important information. Store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important information. A store at 20 to 25°C (68° T 27°F) and protect from moister dependency interpreteints crospovidone, gloveny important information. A store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important information. A store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important indicate componer dispersion, microsystaline cellulose, store at 20 to 25°C (68° T 27°F) and protect from moisture Inactive ingredients crospovidone, gloveny important indicate componer dispersio

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

MAY TAKE 1 to 4 DAYS FOR FULL EFFECT **NDC 0363-1008-05 Walgreens** Compare to the active

ingredient in Prilsec OTC®†

Omeprazole DELAYED-RELEASE TABLETS 20 mg / ACID REDUCER 24 HR

Treats frequent heartburn!

14 TABLETS ONE 14-DAY COURSE OF TREATMENT ACTUAL SIZE



ACID REDUCER

omeprazole tablet, delayed release

Product Information

Product Type		HUMAN OTC DRUG	Item Code (Sour	ce)	NDC:036	3-1008				
Route of Administ	ration	ORAL								
Active Ingredie	nt/Active	Moiety								
	Ing	redient Name		Basis Stren		Strength				
	ESIUM (UNII	426QFE7XLK) (OMEPRAZ	OLE -	OMEPRAZOLI		20 mg				
UNII:KG60484QX9)										
Inactive Ingredi	ionts									
mactive mgrea	ents	Ingredient Nan	ne			Strength				
CROSPOVIDONE (35	.MU.M) (UN	•								
GLYCERYL MONOST										
	HYDROXYPROPYL CELLULOSE (90000 WAMW) (UNII: UKE75GEA7F)									
HYPROMELLOSE 291										
MAGNESIUM STEARA	-									
		RYLATE COPOLYMER (1	:1) TYPE A (UNII: NX76	6LV5T8J)						
MICROCRYSTALLINE										
POLYETHYLENE GLY										
POLYSORBATE 80 (L										
FERRIC OXIDE RED (
SILICON DIOXIDE (UI										
SODIUM HYDROXIDE	-									
SODIUM STEARYL FU										
DEXTROSE, UNSPEC	-	•								
STARCH, CORN (UNII:										
SUCROSE (UNII: C151		,,								
TALC (UNII: 7SEV7J4R	-									
TITANIUM DIOXIDE (2IP)								
TRIETHYL CITRATE (-								
FERRIC OXIDE YELLO										
	-	15000 WAMW) (UNII: 8V/	NB711C5E)							
HYPROMELLOSE 291										
	LU (U MPA.3									
Product Charac	torictics									
Color	PINK		Score		no sc					
Shape	RECTANGL	E (Oblong)	Size		14mn	า				
Flavor			Imprint Code		Z;69					
Contains										
Packaging				.						
# Item Code	Pa	kage Description	Marketing Date	-		ting End ate				
1 NDC:0363-1008-	in 1 CARTO	I	05/06/2021							
- 05 -										

	IDA	ANDA206877	05/06/2021					
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
Marketing Information								
3		14 in 1 BOTTLE; Type 0: Not a Combination Product						
3	NDC:0363-1008- 62	3 in 1 CARTON	05/06/2021					
2		14 in 1 BOTTLE; Type 0: Not a Combination Product						
2	NDC:0363-1008- 61	2 in 1 CARTON	05/06/2021					
1		Product						

Labeler - WALGREEN CO. (008965063)

Registrant - Aurobindo Pharma Limited (650082092)

Establishment								
Name	Address	ID/FEI	Business Operations					
Aurobindo Pharma Limited		650381903	ANALYSIS(0363-1008), MANUFACTURE(0363-1008)					

Revised: 12/2023

WALGREEN CO.