

MAGNESIUM OXIDE- magnesium oxide tablet
Richmond Pharmaceuticals, Inc.

Magnesium Oxide Tablets

ACTIVE INGREDIENT (In each tablet)

Magnesium Oxide 400mg (241.2mg elemental magnesium)

Purpose

Antacid

USES

Relieves: ■ acid indigestion ■ upset stomach

WARNINGS

Ask a doctor if you have:

kidney disease

Ask a doctor or pharmacist before use if you are:

taking a prescription drug. Antacids may interact with certain prescription drugs.

Do not take

more than 2 tablets in a 24 hour period or use the maximum dosage of this product for more than two weeks, except under the advise and supervision of a physician. May have a laxative effect.

If pregnant or breast-feeding,

ask a health professional before use.

Keep Out of Reach of Children.

DIRECTIONS

Antacid Directions: ■ take 1 tablet twice a day or as directed by a physician

Magnesium Supplement Directions: ■ take 1 to 2 tablets daily or as directed by a physician

OTHER INFORMATION

■ store at room temperature 59°-86° F (15°-30°C) ■ do not use if imprinted safety seal

under cap is broken or missing

■ Magnesium content per tablet: 253 mg

INACTIVE INGREDIENTS

croscarmellose sodium, microcrystalline cellulose, magnesium stearate

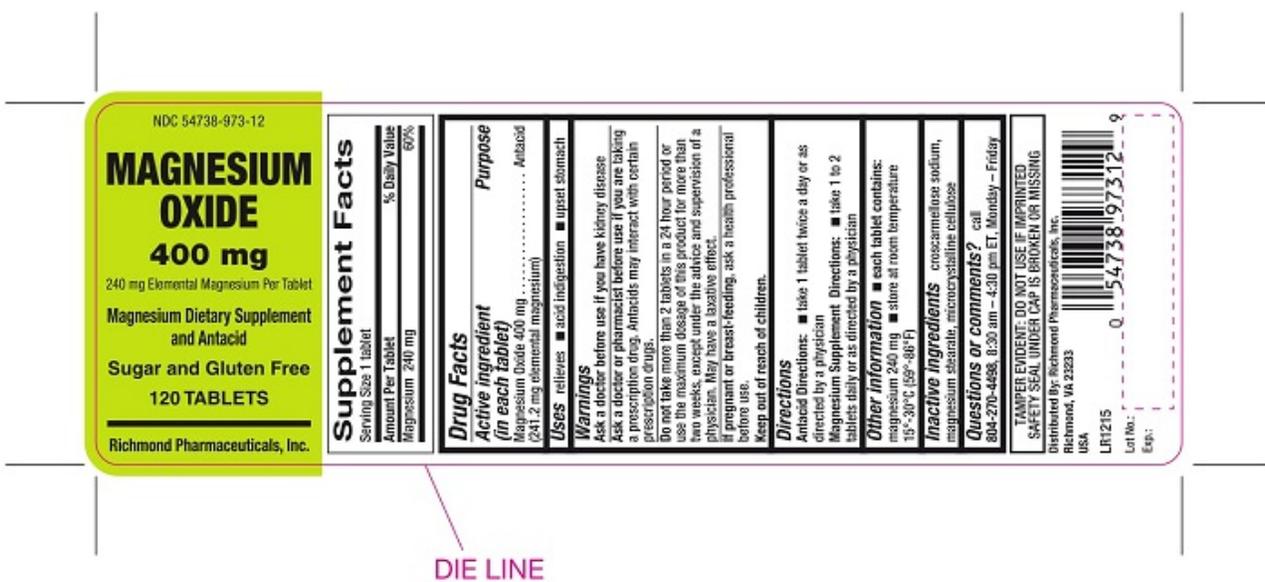
Questions or Comments

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

call 804-270-4498, 8.30 am-4.30 pm ET, Monday - Friday

PRINCIPAL DISPLAY PANEL

NDC 54738-973-12- 120 tablets



MAGNESIUM OXIDE

magnesium oxide tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54738-973
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MAGNESIUM OXIDE (UNII: 3A3U0G171G) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM OXIDE	400 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	white (Off)	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	AP;136
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54738-973-12	120 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/01/2016	

Labeler - Richmond Pharmaceuticals, Inc. (043569607)

Registrant - Advance Pharmaceutical, Inc. (078301063)

Establishment

Name	Address	ID/FEI	Business Operations
Advance Pharmaceutical, Inc.		078301063	manufacture(54738-973)

Revised: 12/2024

Richmond Pharmaceuticals, Inc.