MAGNESIUM OXIDE- magnesium oxide tablet Richmond Pharmaceuticals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Magnesium Oxide Tablets

ACTIVE INGREDIENT (In each tablet)

Magnesium Oxide 420 mg

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

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 54738-974-01- 100 tablets



magnesium oxide tablet

Product Information							
Product Type	HUMAN OTC DRUG		Item Code (Source)		NDC:54738	-974	
Route of Administration	ORAL						
Active Ingredient/Active M	loiety						
Ingredient Name Basis of Str						Strength	
MAGNESIUM O XIDE (UNII: 3A3U0 GI71G) (MAGNESIUM CATION - UNII:T6 V3LHY838)				MAGNESI	JM O XIDE	420 mg	
Inactive Ingredients							
Ingredient Name					St	Strength	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)							
CELLULOSE, MICROCRYSTALL	INE (UNII: OP1R32D61U)						
MAGNESIUM STEARATE (UNII: 70	097M6I30)						
Product Characteristics							
	(Off)	Score		1	no score		

Flavor	In	mprint Code		AP;137		
Contains						
D 1 1						
Packaging						
# Item Code	Package Description		Marketing Start Date	Marketing End Date		
1 NDC:54738-974-01	00 in 1 BOTTLE; Type 0: Not a Combination Product		0 2/0 1/20 16			
Marketing Information						
Marketing Category	Application Number or Monogra	ph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part331		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(54738-974)

Revised: 8/2019

Richmond Pharmaceuticals, Inc.