MAGNESIUM OXIDE- magnesium oxide tablet
PAR Pharmaceuticals

Magnesium Oxide Tablets

ACTIVE INGREDIENT (In each tablet)

Magnesium Oxide 400mg (240mg 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 controlled room temperature 59°-86° F (15°-30°C) ■ do not use if imprinted

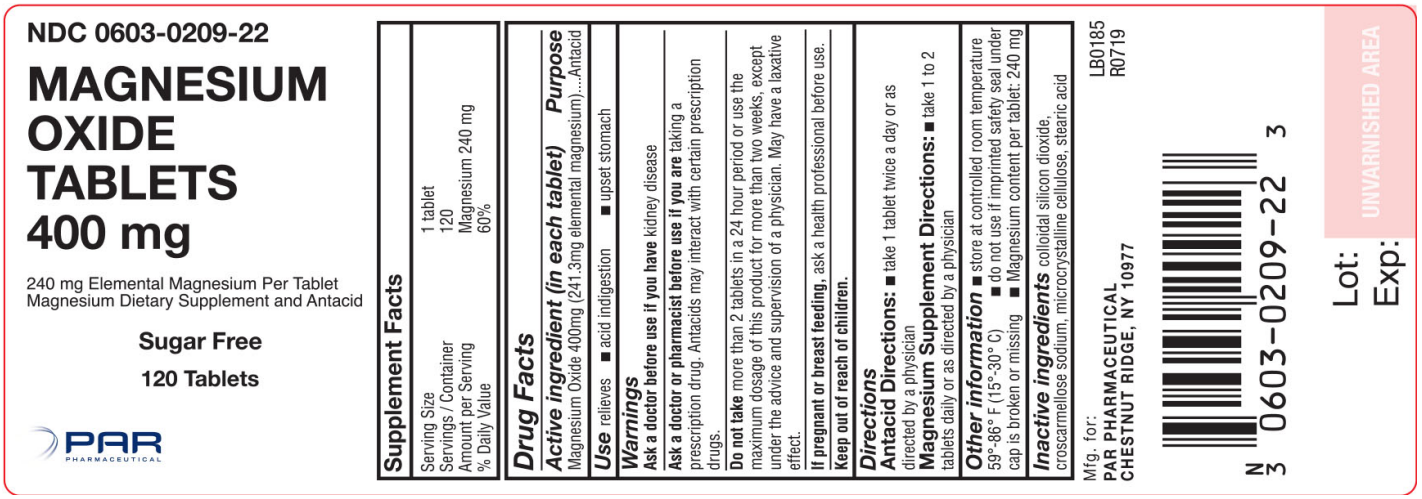
safety seal under cap is broken or missing

■ Magnesium content per tablet: 240 mg

INACTIVE INGREDIENTS

colloidal silicon dioxide, croscarmellose sodium, microcrystalline cellulose, stearic acid

PRINCIPAL DISPLAY PANEL



Magnesium Oxide Tablets 420mg

MAGNESIUM OXIDE

magnesium oxide tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0603-0209
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MAGNESIUM OXIDE (UNII: 3A3U0GI71G) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM OXIDE	400 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	11mm
Flavor		**Imprint Code**	174
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0603-0209-22	120 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	07/01/2014	

Labeler - PAR Pharmaceuticals (092733690)

Revised: 6/2020 PAR Pharmaceuticals