MAGNESIUM OXIDE- magnesium oxide tablet
Westminster Pharmaceuticals, LLC

Magnesium Oxide

Drug Facts

Active Ingredient

Magnesium Oxide 400 mg (241.3 mg Elemental Magnesium)

Purpose

Antacid

Use

relieves:
- acid indigestion
- upset stomach

Warnings

Ask a doctor before use if
- you have kidney disease
- you are taking a prescription drug (antacids may interact with certain prescription drugs)
- you are pregnant or breast feeding.

Do not take more than 2 tablets in a 24 hour period.
- May have a laxative effect.

Keep out of reach of children.

Directions
- take one or two antacid tablets daily. Do not exceed two tablets unless directed by a physician.

Other Information
- Store at controlled room temperature 15° to 30°C (59° to 86°F).
- Tamper evident, do not use if imprinted safety seal under cap is broken or missing.

Inactive Ingredients

Corn Starch, Croscarmellose Sodium, Magnesium Stearate, Microcrystalline Cellulose,

Silicon Dioxide, Stearic Acid.

Questions?

Call 1-844-221-7294 M-F 9am - 5pm EST

PRINCIPAL DISPLAY PANEL - 400 mg Tablet Bottle Label

NDC 69367-298-20

Magnesium
Oxide 400 mg

241.3 mg Elemental Magnesium
Antacid

Sugar and Gluten Free

120 Tablets

Westminster
Pharmaceuticals

MAGNESIUM OXIDE

magnesium oxide tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69367-298
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
STARCH, CORN (UNII: O8232NY3SJ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	WHITE	**Score**	2 pieces
Shape	ROUND	**Size**	11mm
Flavor		**Imprint Code**	AM3
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69367-298-20	120 in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M001	10/15/2020	

Labeler - Westminster Pharmaceuticals, LLC (079516651)

Revised: 10/2020 Westminster Pharmaceuticals, LLC