

MAGNESIUM OXIDE- magnesium oxide tablet
Method Pharmaceuticals, LLC

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

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

Croscarmellose Sodium, Microcrystalline Cellulose, Silicon Dioxide, and Stearic Acid.

Questions?

Call Method at 1-877-250-3427 or FDA at 1-800-FDA-1088

Manufactured for:
 Method Pharmaceuticals, LLC
 Fort Worth, Texas 76118
 Rev. 04/18

Principal Display Panel

NDC 58657-120-12
 Magnesium Oxide
 400 mg
 241.3 Elemental Magnesium
 Antacid
 120 Tablets



MAGNESIUM OXIDE

magnesium oxide tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58657-120
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	11mm
Flavor		Imprint Code	120
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58657-120-12	120 in 1 BOTTLE; Type 0: Not a Combination Product	07/25/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	07/25/2018	

Labeler - Method Pharmaceuticals, LLC (060216698)

Revised: 2/2020

Method Pharmaceuticals, LLC