MAGNESIUM OXIDE- magnesium oxide tablet
Bryant Ranch Prepack

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

HOW SUPPLIED

NDC: 71335-1960-1: 60 Tablets in a BOTTLE

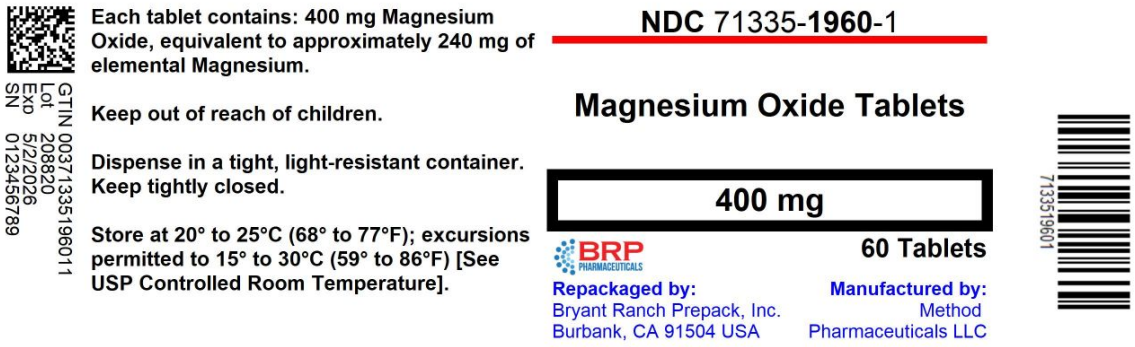
NDC: 71335-1960-2: 30 Tablets in a BOTTLE

NDC: 71335-1960-3: 90 Tablets in a BOTTLE

NDC: 71335-1960-4: 100 Tablets in a BOTTLE

NDC: 71335-1960-5: 120 Tablets in a BOTTLE

Magnesium Oxide 400 mg Tablet



MAGNESIUM OXIDE

magnesium oxide tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71335-1960(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)	
MICROCRYSTALLINE CELLULOSE (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:71335-1960-1	60 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2021	
2	NDC:71335-1960-2	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2021	
3	NDC:71335-1960-3	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2021	
4	NDC:71335-1960-4	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2021	
5	NDC:71335-1960-5	120 in 1 BOTTLE; Type 0: Not a Combination Product	10/06/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	07/25/2018	

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-1960) , RELABEL(71335-1960)