MAGNESIUM OXIDE- magnesium oxide tablet Marlex Pharmaceuticals Inc.

MAGNESIUM OXIDE- magnesium oxide tablet

ACTIVE INGREDIENT (In each tablet)

Magnesium Oxide 420 mg

Purpose

Dietary Supplement

Supplement Facts

Serving Size	1 Tablet	
Magnesium (as Magnesium Oxide		
Amount per Serving	253mg	
% Daily value	63%	

Other Ingredients

Colloidal silicon dioxide, Croscarmellose sodium, microcrystalline cellulose, stearic acid Magnesium oxide may help with magnesium deficiencies.*

SUGGESTED USE

As a dietary supplements, adult take one (1) tablet daily preferably with food or as directed by your healthcare practitioner.

CAUTIONS

If you are pregnant, nursing, taking any medications or have any medical condition, consult your doctor before use. If you have any kidney disease, take only under the supervision of a doctor. Discontinue use and consult your doctor if any adverse reactions occur.

Keep out of reach of children.

Store in a cool, dry place.

DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

^{*} These statement has not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

Manufactured for/ Distributed by:

Marlex Pharmaceuticals, Inc.,

New Castle, DE 19720

Rev.1 10/23 AA

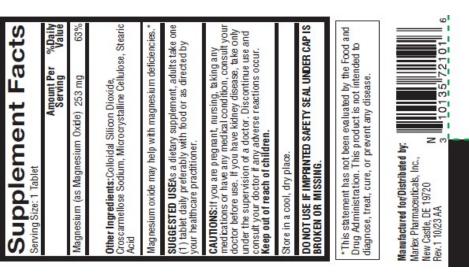
PRINCIPAL DISPLAY PANEL

NDC 10135-0721-01

Magnesium Oxide 420 mg

100 TABLETS





MAGNESIUM OXIDE

magnesium oxide tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10135-721
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength Strength
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MAGNESIUM OXIDE (UNII: 3A3U0GI71G) (MAGNESIUM CATION - UNII:T6V3LHY838) MAGNESIUM OXIDE 420 mg

Inactive Ingredients		
Ingredient Name	Strength	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		

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

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:10135-721- 01	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	11/01/2023	

Labeler - Marlex Pharmaceuticals Inc. (782540215)

Revised: 12/2023 Marlex Pharmaceuticals Inc.