## MAGNESIUM OXIDE (MG SUPPLEMENT)- magnesium oxide tablet Nationwide Pharmaceutical LLC

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#### Magnesium Oxide

#### WARNINGS

Not recommended for use in amounts over the recommended daily intake of 420mg per day (1 tablet). Ask a doctor or pharmacist before use if you have a known allergy to this product. Stop use and contact a doctor if allergic reaction occurs. Consult a doctor before using this product if you have kidney disease or are pregnant or nursing a baby. May have a laxative effect.

# DIRECTIONS

Adults: Take one tablet daily with a meal as a dietary supplement or as directed by a doctor.

### **OTHER INFORMATION**

- Store at Controlled Room Temperature 15-30°C (58-86°F)
- Store in original container with lid tightly closed in a dry place
- Keep out of reach of children
- Do not use if imprinted safety seal under cap is broken or missing
- Use by expiration date on package

Supplement Facts Serving Size: 1 Tablet				
Amount Per Tablet	% Daily Value $^*$			
Magnesium 252 mg	63%			
(From Magnesium oxide 420 mg)				
<ul> <li>Percent Daily Values are b calorie diet.</li> </ul>	ased on a 2,000			

**Other ingredients:** Microcrystalline, Cellulose, Croscarmellose Sodium, Silicon Dioxide, Stearic Acid, Magnesium Stearate, Titanium dioxide, Hypromellose.

### Manufactured for/Distributed by:

Nationwide Pharmaceutical LLC San Antonio, TX 78216 Rev 4/2024

# **PRINCIPAL DISPLAY PANEL - 420 mg Tablet Bottle Label**

Nationwide Pharmaceutical

NDC 69375-0008-10
Magnesium Oxide
420 mg
Dietary Supplement
100 TABLETS
Made in the USA



MAGNESIUM OXIDE magnesium oxide tablet	(MG SUPPLEMEI	NT)			
Product Information					
Product Type	DIETARY SUPPLEMENT	Item Code (Source)		NHRIC:69375-008	
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ing	redient Name		Basis of S	trength	Strengt
Magnesium Oxide (UNII: 3A3U0GI71G) (Magnesium Cation - UNII:T6V3LHY838)			Magnesium Oxide		420 mg
Inactive Ingredients					
Inactive Ingredients	Ingredient Name			Sti	rength
-	•			Sti	rength
MICROCRYSTALLINE CELLULOS	<b>SE</b> (UNII: OP1R32D61U)			Sti	rength
MICROCRYSTALLINE CELLULOS SILICON DIOXIDE (UNII: ETJ7Z6X	SE (UNII: OP1R32D61U) BU4)			Sti	rength
MICROCRYSTALLINE CELLULOS SILICON DIOXIDE (UNII: ETJ7Z6X CROSCARMELLOSE SODIUM (U	<b>SE</b> (UNII: OP1R32D61U) BU4) NII: M28OL1HH48)			Sti	rength
MICROCRYSTALLINE CELLULOS SILICON DIOXIDE (UNII: ETJ7Z6X CROSCARMELLOSE SODIUM (U STEARIC ACID (UNII: 4ELV7Z65A)	<b>SE</b> (UNII: OP1R32D61U) BU4) NII: M28OL1HH48) P)			Sti	rength
Inactive Ingredients MICROCRYSTALLINE CELLULOS SILICON DIOXIDE (UNII: ETJ7Z6X CROSCARMELLOSE SODIUM (U STEARIC ACID (UNII: 4ELV7Z65AI MAGNESIUM STEARATE (UNII: 7 Titanium Dioxide (UNII: 15FIX9V	<b>SE</b> (UNII: OP1R32D61U) BU4) NII: M28OL1HH48) P) 0097M6I30)			Sti	rength

Packaging					
# Item Code	Package Description	Marketing	Start Date	Marketin	g End Date
<b>1</b> NHRIC:69375-008-10	0 100 in 1 BOTTLE				
Marketing In	formation				
Marketing In Marketing Category	<b>formation</b> Application Number or Mo Citation	nograph	Marketing St Date	tart Ma	rketing End Date
•	Application Number or Mo	nograph		tart Ma	

Supplement Facts				
Serving Size :		Serving per Container :		
	Amount Per Serving	% Daily Value		
color				
scoring	1			
shape				
size (solid drugs)	11 mm			

Labeler - Nationwide Pharmaceutical LLC (079265801)

Revised: 4/2024

Nationwide Pharmaceutical LLC