

**REGULAR STRENGTH ANTACID- aluminum hydroxide, magnesium hydroxide,
dimethicone suspension
KROGER COMPANY**

KRO ANTACID org

Active ingredients (in each 10 mL dose)

Aluminum hydroxide 400 mg (equivalent to dried gel, USP)
Magnesium hydroxide 400 mg
Simethicone 40 mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- the symptoms referred to as gas

Warnings

Do not take more than 120 mL (12 doses) in a 24 hour period or use the maximum dosage for more than 2 weeks .

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are presently taking a prescription drug. Antacids may interact with certain prescription drugs.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children.

Directions

- shake well before use
- adults and children 12 years and older: take 10 mL to 20 mL (1 to 2 doses) between meals as needed, at bedtime, or as directed by a doctor.
- children under 12 years: ask a doctor

- measure with dosing cup provided
- mL=milliliter

Other information

- **each 10 mL dose contains:** magnesium 165 mg, sodium 6 mg
- protect from freezing
- store at room temperature
- keep tightly closed

Inactive ingredients

benzyl alcohol, butylparaben, flavor (contains alcohol), hydroxyethylcellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments?

1-800-632-6900

package label

NDC 30142-637-12



Regular Strength

Antacid & Gas Relief

Alumina, Magnesia & Simethicone Oral Suspension USP

Alcohol 0.15%

FAST RELIEF OF:

Heartburn & Acid Indigestion
Pressure & Bloating
Sour Stomach

Original Flavored Liquid

12 FL OZ (355 mL)

215-05112-0 REV 0021

Drug Facts

TAMPER-EVIDENT: Do not use if imprinted neckband is missing or broken.

Active ingredients (in each 10 mL dose)

Aluminum hydroxide 400 mg (equivalent to dried gel USP)	Antacid
Magnesium hydroxide 400 mg	Antacid
Simethicone 40 mg	Antigas

Purposes

Uses relieves

- heartburn
- sour stomach
- acid indigestion
- the symptoms referred to as gas

Warnings

Do not take more than 120 mL (12 doses) in a 24 hour period or use the maximum dosage for more than 2 weeks.

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are presently taking a prescription drug. Antacids may interact with certain prescriptions drugs.

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children.

Directions

- shake well before each use
- adults and children 12 years and older: take 10 mL to 20 mL (1 to 2 doses) between meals as needed, at bedtime, or as directed by a doctor
- children under 12 years: ask a doctor
- measure with dosing cup provided
- mL = milliliter

Other information

- each 10 mL dose contains: magnesium 165 mg, sodium 6 mg
- store at room temperature
- protect from freezing
- keep tightly closed

Inactive ingredients benzyl alcohol, butylparaben, flavor (contains alcohol), hydroxyethylcellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments? 1-800-632-6900

DISTRIBUTED BY THE KROGER CO. CINCINNATI, OH 45202

Our Brands. Our Guarantee. Love It or Your Money Back. www.kroger.com

915-05112-0 REV 0021



REGULAR STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-637
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	400 mg in 10 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	400 mg in 10 mL

DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)		DIMETHICONE	40 mg in 10 mL	
Inactive Ingredients				
Ingredient Name			Strength	
BUTYLPARABEN (UNII: 3QPI1U3FV8)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
WATER (UNII: 059QF0K00R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SORBITOL SOLUTION (UNII: 8KW3E207O2)				
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-637-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M001	03/01/2021		

Labeler - KROGER COMPANY (006999528)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(30142-637)

Revised: 11/2023

KROGER COMPANY