

**MAXIMUM STRENGTH ANTACID- aluminum hydroxide, magnesium hydroxide,
dimethicone suspension
CARDINAL HEALTH**

ldr max antacid

Active ingredients (in each 10 mL dose)

Aluminum hydroxide 800 mg (equivalent to dried gel, USP)

Magnesium hydroxide 800 mg

Simethicone 80 mg

Purposes

Antacid

Antigas

Uses

relieves

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

Warnings

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are taking a prescription drug.

Antacids may interact with certain prescription drugs.

Stop use and ask a doctor if symptoms last more than 2 weeks

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

Keep out of reach of children. In case of overdose, get medical help or contact Poison Control Center (1-800-222-1222) right away.

Directions

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

- do not exceed 60 mL (6 doses) in a 24 hour period or use the maximum dosage for more than 2 weeks
- children under 12 years of age: ask a doctor

Other information

- each 10 mL teaspoonful contains: **magnesium 340 mg, sodium 10 mg**
- do not freeze
- store at room temperature tightly closed

Inactive ingredients

benzyl alcohol, butylparaben, carboxymethylcellulose sodium, flavor (contains alcohol), hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments?

1-800-540-3765

package Label

LEADER²™

NDC 70000-0062-1

Maximum Strength

Antacid Liquid

Aluminum Hydroxide
Magnesium Hydroxide
Simethicone
Antacid | Antigas

Original Flavor

Fast Relief Of:
Acid Indigestion
Heartburn
Sour Stomach
Contains Alcohol 0.5%

COMPARE TO MYLANTA[®] MAXIMUM STRENGTH active ingredients*

100% Money Back Guarantee

12 FL OZ (355 mL)

295-06122-0 REV GC0920

Drug Facts

TAMPER-EVIDENT: Do not use if the breakaway band on cap is broken or missing.

Active ingredients (in each 10 mL dose)	Purposes
Aluminum hydroxide (equivalent to dried gel, USP) 800 mg	Antacid
Magnesium hydroxide 800 mg	Antacid
Simethicone 80 mg	Antigas

Uses relieves • heartburn • sour stomach • acid indigestion • the symptoms referred to as gas

Warnings

Ask a doctor before use if you have

• kidney disease • magnesium-restricted diet

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

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Other information

• each 10 mL dose contains: magnesium 340 mg, sodium 10 mg
• do not freeze • store at room temperature tightly closed

Inactive ingredients benzyl alcohol, butylparaben, carboxymethylcellulose sodium, flavor, hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments? 1-800-540-3765

*This product is not manufactured or distributed by Johnson & Johnson, owner of the registered trademark Maximum Strength Mylanta[®].

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www.myleader.com 1-800-200-6313
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995-06122-0 REV GC0920

100% Money Back Guarantee
Return to place of purchase if not satisfied.

CIN 5667936 REV. 9/20



MAXIMUM STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0062
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	800 mg in 10 mL

MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	800 mg in 10 mL
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	80 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0062-1	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	09/01/2020	

Labeler - CARDINAL HEALTH (063997360)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(70000-0062)

Revised: 11/2023

CARDINAL HEALTH