REGULAR STRENGTH ANTACID- aluminum hydroxide, magnesium hydroxide, dimethicone suspension Chain Drug Marketing Association

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Active ingredients (in each 5 mL teaspoonful)

Aluminum hydroxide 200 mg (equivalent to dried gel, USP)
Magnesium hydroxide 200 mg

Simethicone 20mg

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.

Directions

shake well before use

- adults and children 12 years and older: take 2 to 4 teaspoonfuls between meals, at bedtime, or as directed by a doctor
- do not take more than 24 teaspoonfuls in 24 hours or use the maximum dosage for more than 2 weeks
- children under 12 years: ask a doctor

Other information

- each 5 mL teaspoonful contains: magnesium 85 mg, sodium 3 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?

1-800-540-3765

package Label

NDC 63868-712-57



Compare to the Active Ingredients in Mylanta Regular Strength

Regular Strength Antacid

Antigas, Fast Acting

Alumina, Magnesia, and Simethicone Oral Suspension USP

Fast Soothing Relief of:

Acid Indigestion Heartburn Sour Stomach Pressure & Bloating

Alcohol: 0.15%

Original Flavor

12 FL OZ (355 mL)

REV GC-0818 231-05112-0

Drug Facts

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

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

Warnings

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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 for more than 2 weeks

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Questions or comments? 1-800-540-3765

*This product is not manufactured or distributed by the owner of the registered trademark MYLANTA®.



REV GC-0818 931-05112-0

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REGULAR STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63868-712

Route of Administration ORAL

Active Ingredient/Active Moiety

ı	Active ingredient/Active Molety					
	Ingredient Name	Basis of Strength	Strength			
	ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE -	ALUMINUM	200 mg			

UNII:5QB0T2IUN0)	HYDROXIDE	in 5 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	200 mg in 5 mL
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	20 mg in 5 mL

Inactive Ingredients				
Ingredient Name	Strength			
BENZYL ALCOHOL (UNII: LKG8494WBH)				
BUTYLPARABEN (UNII: 3QPI1U3FV8)				
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
WATER (UNII: 059QF0KO0R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SORBITOL SOLUTION (UNII: 8KW3E207O2)				

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	LEMON (citrus mint)	Imprint Code	
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:63868-712- 57	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2018	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	art Marketing End Date	
OTC Monograph Drug	M001	03/01/2014		

Labeler - Chain Drug Marketing Association (011920774)

Registrant - GCP Laboratories (965480861)

Establishment				
Name	Address	ID/FEI	Business Operations	
GCP Laboratories		965480861	manufacture(63868-712)	

Revised: 11/2023 Chain Drug Marketing Association