

**MAXIMUM STRENGTH ANTACID AND GASRELIEF- aluminum hydroxide,
magnesium hydroxide, dimethicone suspension
GOODSENSE**

GS 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

Do not take more than 60 mL (6 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 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 of age 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 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

GOODSENSE[®]

NDC 50804-619-12

Maximum Strength

Antacid & Gas Relief

Alumina, Magnesia, and Simethicone
Oral Suspension USP

Fast Acting
Soothing Relief of:

Acid Indigestion

Heartburn

Sour Stomach

Pressure & Bloating

Original
Flavored Liquid

Compare to **active ingredients of**
MYLANTA[®] Maximum Strength*

Contains less than
0.5% Alcohol

12 FL OZ (355 mL)

274-06122-1 REV GC0821

Drug Facts

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

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
Do not take more than 60 mL (6 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 of age and older: take 10 mL to 20 mL (1 to 2 doses) between meals as needed, at bedtime, or as directed by a doctor
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Other information
• each 10 mL dose contains: magnesium 340 mg, sodium 10 mg
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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

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

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Perrigo Direct, Inc.
Peachtree City, GA 30269
www.PerrigoDirect.com
1-800-540-3765
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974-06122-1 REV GC0821



MAXIMUM STRENGTH ANTACID AND GASRELIEF

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50804-619
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

UNII:5QB0T2IUN0)	HYDROXIDE	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:50804-619-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	07/01/2020	

Labeler - GOODSENSE (076059836)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(50804-619)

Revised: 11/2023

GOODSENSE