ADVANCED ANTACID MINT- aluminum hydroxide, magnesium hydroxide, dimethicone suspension
GOODSENSE

GS ant mnt

Active ingredients (in each 10 mL dose)

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

Magnesium hydroxide 400 mg

Simethicone 40mg

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 each use

• adults and children 12 years and older: take 10 mL to 20 mL (1-2 doses) four times a day or as directed by a doctor
• do not exceed 80 mL (8 doses) in a 24 hour period or use the maximum dosage for more than 2 weeks
• children under 12 years: ask a doctor

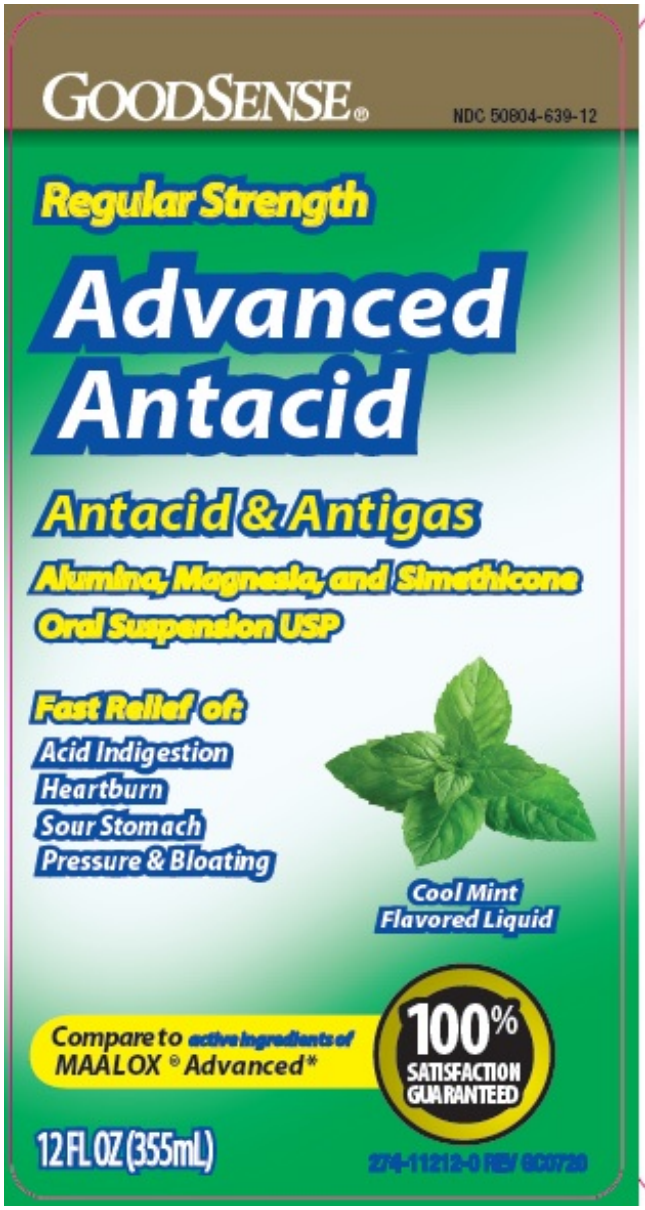
Other information

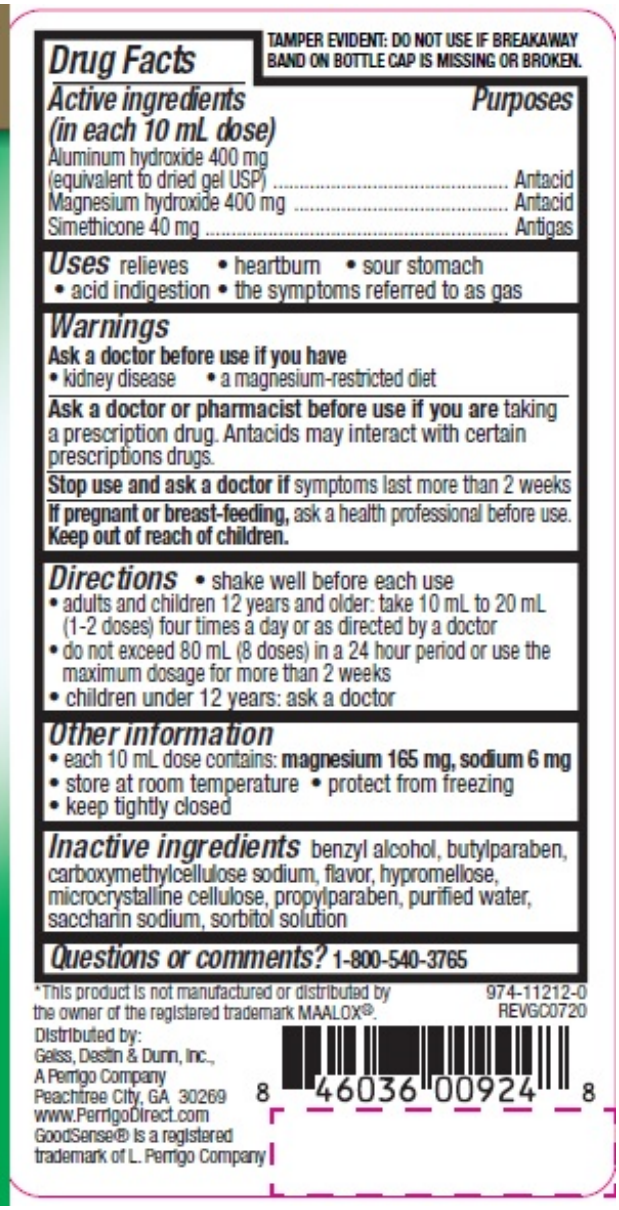
• each 10 mL dose contains: **magnesium 165 mg, sodium 6 mg**
• store at room temperature
• protect from freezing
• keep tightly closed

Inactive ingredients

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

package Label





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 400 mg (equivalent to dried gel USP) ..... Antacid
Magnesium hydroxide 400 mg ..... Antacid
Simethicone 40 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 • a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are taking a prescription drug. Antacids may interact with certain prescriptions 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 each use
• adults and children 12 years and older: take 10 mL to 20 mL (1-2 doses) four times a day or as directed by a doctor
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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, 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 the owner of the registered trademark MAALOX®.

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REVGC0720

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ADVANCED ANTACID MINT

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-639
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
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)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	MINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-639-12	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(50804-639)

Revised: 11/2023 GOODSENSE