

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

**Chain Drug Marketing Association**

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**QC ant max**

**Active ingredients (in each 5 mL teaspoonful)**

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
- upset stomach due to these symptoms
- 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 this and all drugs out of the 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 12 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 165 mg, sodium 5 mg
- store at room temperature and avoid freezing
- close cap tightly after use

### **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**

NDC 63868-715-57



\*Compare to the  
Active Ingredients  
in Mylanta®  
Maximum Strength

# Maximum Strength Antacid

## Antigas

Alumina, Magnesia, and  
Simethicone  
Oral Suspension USP

Alcohol: 0.5%

Fast Soothing Relief of:  
Acid Indigestion  
Heartburn  
Sour Stomach  
Pressure & Bloating



Original Flavor

12 FL OZ (355 mL)

REV GC-0618  
231-06122-2

### Drug Facts

TAMPER EVIDENT: DO NOT USE IF THE BREAKAWAY BAND ON CAP IS BROKEN OR MISSING.

#### Active ingredients (in each 5 mL teaspoonful) Purposes

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

**Uses** relieves: • heartburn • sour stomach • acid indigestion  
• upset stomach due to these symptoms  
• the symptoms referred to as gas

#### Warnings

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

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Stop use and ask a doctor if symptoms last for more than 2 weeks  
If pregnant or breast-feeding, ask a health professional before use.  
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#### Directions

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\*This product is not manufactured or distributed by Johnson & Johnson Merck, distributor of Mylanta®.



REV GC-0618  
931-06122-1



Distributed by C.D.M.A., Inc.®  
43157 W. Nine Mile  
Novi, MI 48375-0995  
www.qualitychoice.com  
Questions: 248-449-9300

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## MAXIMUM STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-715
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 5 mL

<b>MAGNESIUM HYDROXIDE</b> (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	400 mg in 5 mL
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	40 mg in 5 mL

### Inactive Ingredients

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

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-715-57	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2014	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	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(63868-715)

Revised: 11/2023

Chain Drug Marketing Association