

REGULAR STRENGTH ANTACID ANTIGAS- aluminum hydroxide, magnesium hydroxide, simethicone liquid
Chain Drug Marketing Association Inc.

QCH Regular Strength Antacid Antigas 503

ACTIVE INGREDIENTS (in each 10 mL)

Aluminum hydroxide (equiv. to dried gel, USP) 400 mg
Magnesium hydroxide 400 mg
Simethicone 40 mg

PURPOSE

Antacid
Antacid
Antigas

USE(S)

relieves:

- acid indigestion
- heartburn
- sour stomach
- upset stomach and gas associated with these symptoms

WARNINGS

Do not take more than 80 mL in a 24-hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a doctor.

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.

STOP USE AND ASK DOCTOR

if symptoms last more than two weeks

KEEP OUT OF REACH OF CHILDREN

DIRECTIONS

- **shake well before use**
- adults and children 12 years and older: take 10 - 20 mL four times a day, or as directed by a doctor
- children under 12 years: consult a doctor
- mL = milliliter

OTHER INFORMATION

- **each 10 mL contains:** magnesium 170 mg, sodium 5 mg
- store at 20°C-25°C (68°-77°F)
- do not freeze

INACTIVE INGREDIENTS

ethyl alcohol, flavor, glycerin, hydroxyethyl cellulose, methylparaben, propylene glycol, propylparaben, purified water, saccharin sodium, simethicone emulsion, sorbitol

PRINCIPAL DISPLAY PANEL

NDC 83324-125-12

Regular strength

Antacid Liquid

Antacid & Antigas

Aluminum Hydroxide 400mg

Magnesium Hydroxide 400mg

Simethicone 40mg

Relieves:

Acid Indigestion

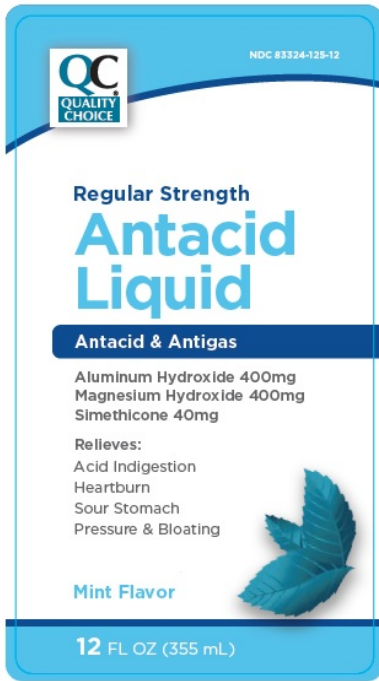
Heartburn

Sour Stomach

Pressure & Bloating

Mint Flavor

12 FL OZ (355 mL)



Drug Facts DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING

Active ingredients (in each 10 mL) Purpose
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 Magnesium hydroxide 400 mg.....Antacid
 Simethicone 40 mg.....Antigas

Uses relieves: ■ acid indigestion ■ heartburn ■ sour stomach
 ■ upset stomach and gas associated with these symptoms

Warnings
 Do not take more than 80 mL in a 24-hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a doctor.
 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.
 Stop use and ask a doctor if symptoms last more than two weeks.
 Keep out of reach of children.

Directions ■ shake well before use ■ adults and children 12 years and older: take 10 - 20 mL four times a day, or as directed by a doctor ■ children under 12 years: consult a doctor ■ mL = milliliter

Other information ■ each 10 mL contains: magnesium 170 mg, sodium 5 mg ■ store at 20°C-25°C (68°F-77°F) ■ do not freeze

Inactive ingredients ethyl alcohol, flavor, glycerin, hydroxyethyl cellulose, methylparaben, propylene glycol, propylparaben, purified water, saccharin sodium, simethicone emulsion, sorbitol.

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Distribute d by CDMA, Inc. Novi, MI 48375 www.quality-choice.com Questions: 800-935-2362

UNFINISHED AREA FOR LOT & EXP (LINE DOES NOT PRINT)

REGULAR STRENGTH ANTACID ANTIGAS

aluminum hydroxide, magnesium hydroxide, simethicone liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-125
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)	MAGNESIUM HYDROXIDE	400 mg in 10 mL
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	40 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) (UNII: S38J6RZN16)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics			
Color	WHITE	Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-125-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/27/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	08/27/2024	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Revised: 8/2024

Chain Drug Marketing Association Inc.