QUALITY CHOICE MAXIMUM STRENGTH ANTACID - aluminum hydroxide, magnesium hydroxide, simethicone liquid Chain Drug Marketing Association Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Quality Choice Maximum Strength Antacid and Antigas

ACTIVE INGREDIENTS (in each 10 mL)

Aluminum hydroxide (equiv. to dried gel, USP) 800 mg Magnesium hydroxide 800 mg Simethicone 80 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 60 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 mL 20 mL two 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 340 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 63868-019-12

OC

QUALITY CHOICE

*Compare to the Active Ingredients in Maximum Strength Mylanta®

Maximum Strength

Antacid Liquid

Antacid & Antigas

Aluminum hydroxide 800 mg Magnesium hydroxide 800 mg Simethicone 80 mg

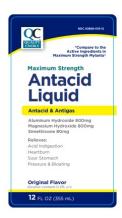
Relieves:

Acid Indigestion
Heartburn
Sour Stomach
Pressure & Bloating

Original Flavor

Alcohol content 0.2% v/v

12 FL OZ (355 mL)





QUALITY CHOICE MAXIMUM STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, simethicone liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-019
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	800 mg in 10 mL	
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM HYDROXIDE	800 mg in 10 mL	
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	80 mg in 10 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
GLYCERIN (UNII: PDC6A3C0OX)		
HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) (UNII: S38J6RZ N16)		
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 (Lemon mint)	Imprint Code	
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:63868-019- 12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/02/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part331	08/02/2022	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Revised: 8/2022