ANTACID ANTIGAS NTA- aluminum hydroxide, magnesium hydroxide, simethicone liquid AMERISOURCEBERGEN DRUG CORPORATION

GNP Antacid Anti-Gas max

Active ingredients (in each 10 mL dose)

Aluminum hydroxide (equivalent to dried gel, USP) 800 mg
Magnesium hydroxide 800 mg
Simethicone 80 mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- symptoms of 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
- do not take more than 60 mL (6 doses) in any 24 hour period or use the maximum dosage for more than 2 weeks
- children under 12 years of age: ask a doctor

Other information

- each 10 mL dose contains: magnesium 340 mg, sodium 10 mg
- do not freeze
- store at room temperature 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

package label



Compare to Mylanta® Maximum Strength active ingredients*

NDC 46122-431-40

MAXIMUM STRENGTH

Antacid and Anti-Gas Liquid

Alumina, Magnesia, and Simethicone Oral Suspension

Fast Relief of:

- Acid Indigestion
- Heartburn

297-06122-0 REV0917

- Sour Stomach
- Pressure and Bloating

ORIGINAL FLAVOR

Contains: Alcohol 0.5%

12 fl oz (355 mL)

Drug Facts

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

Active ingredients (in each 10 mL dose)

Purposes

Àluminum hydroxide

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

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Other information

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Inactive ingredients benzyl alcohol, butylparaben, carboxymethylcellulose sodium, flavor (contains alcohol), hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

C Questions or comments? 1-800-540-3765



Distributed By Amerisource Bergen 1 West First Avenue Conshehooken, PA 19428 Questions or Concerns?



ANTACID ANTIGAS NTA

aluminum hydroxide, magnesium hydroxide, simethicone liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:46122-431

Route of Administration ORAL

Active Ingredient/Active Moiety

Active ingredient/Active Molecy				
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, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	800 mg in 10 mL
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	80 mg in 10 mL

Inactive Ingredients		
Ingredient Name	Strength	
BENZYL ALCOHOL (UNII: LKG8494WBH)		
BUTYLPARABEN (UNII: 3QPI1U3FV8)		
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)		
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:46122-431- 40	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	09/01/2017	

Labeler - AMERISOURCEBERGEN DRUG CORPORATION (007914906)

Registrant - GCP Laboratories (965480861)

Establishment				
Name	Address	ID/FEI	Business Operations	
GCP Laboratories		965480861	manufacture(46122-431)	

Revised: 11/2023 AMERISOURCEBERGEN DRUG CORPORATION