

ANTACID ANTIGAS NTA- aluminum hydroxide, magnesium hydroxide, simethicone liquid
Preferred Pharmaceuticals, Inc.

Good Neighbor Pharmacy Antacid Anti-Gas max cherry

Active ingredients (in each 5 mL, 1 teaspoon)

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

Magnesium hydroxide 400 mg

Simethicone 40 mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- upset stomach due to these symptoms
- symptoms of 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 presently 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. In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- shake well before use
- adults and children 12 years of age and older: take 2 to 4 teaspoonfuls two times a day or as directed by a doctor
- do not exceed 8 teaspoonfuls in a 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 5 mL teaspoonful contains:** magnesium 165 mg, sodium 5 mg
- store at room temperature tightly closed
- protect from freezing

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

Relabeled by Preferred Pharmaceuticals, Inc.

Compare To Maalox® Maximum Strength active ingredients*

NDC 68788-7641-03

MAXIMUM STRENGTH

Antacid

Anti-Gas

Alumina, Magnesia, and Simethicone Oral Suspension

Fast Relief of:

- Acid Indigestion
- Heartburn
- Sour Stomach
- Pressure and Bloating

Cherry Creme Flavor

12 FL OZ (355 mL)

Maximum Strength Mi-Acid



CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.

Maximum Strength Mi-Acid
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Log

Maximum Strength Mi-Acid
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Chart

Maximum Strength Mi-Acid
Qty: Ins:
Insurance NDC:
Lot: Bat:

Billing

Maximum Strength Mi-Acid
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Patient

Each 5 mL teaspoonful contains: Aluminum hydroxide 400mg (equivalent to dried gel, USP)...Antacid / Magnesium hydroxide 400mg...Antacid / Simethicone 40mg...Antigas

Pkg Size: Exp Date: ####/####/####
Lot#: Batch#:

Ins:
Mfg: Good Neighbor Pharmacy

Prod#:

Warning

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. Store at room temperature. Protect from freezing. Keep tightly closed. Tamper-Evident: do not use if breakaway band on bottle cap is missing or broken.



Directions English

Take as Directed



GTIN

SN #####
EXP #####

Instrucciones Espanol:
Tomelo como se indica

ANTACID ANTIGAS NTA

aluminum hydroxide, magnesium hydroxide, simethicone liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-7641(NDC:46122-432)
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
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	400 mg in 5 mL
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE, UNSPECIFIED	40 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	Score

Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7641-3	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/28/2020	

Labeler - Preferred Pharmaceuticals, Inc. (791119022)

Registrant - Preferred Pharmaceuticals, Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals, Inc.		791119022	RELABEL(68788-7641)

Revised: 8/2025

Preferred Pharmaceuticals, Inc.