

**MAXIMUM STRENGTH ANTACID- aluminum hydroxide, magnesium hydroxide,
dimethicone suspension
WAL-MART STORES, INC.**

walmart ant max

Active ingredients (in each 10 mL dose)

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

Magnesium hydroxide 800 mg

Simethicone 80mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- 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

Keep out of the reach of children.

Directions

- shake well before use
- **adults and children 12 years and over:** 10 mL-20 mL (1-2 doses) between meals, at bedtime, or as directed by a doctor
- do not take more than 60 mL (6 doses) in any 24 hour period

- do not use the maximum dosage for more than 2 weeks
- children under 12 years: ask a doctor
- measure with dosing cup provided
- mL=milliliter

Other information

- each 10 mL dose contains: **magnesium 340 mg; sodium 10 mg**
- keep tightly closed
- store at room temperature and avoid freezing

Inactive ingredients

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

Questions or comments?

1-888-287-1915

package Label

NDC 49035-681-12

equate™

MAXIMUM STRENGTH

Antacid

Alumina, Magnesia, and
Simethicone Oral Suspension USP

Antacid/Anti-Gas
Liquid



Fast Acting Soothing Relief Of:

- Heartburn
- Acid indigestion
- Sour stomach
- Pressure and bloating



**Original
Flavor**

12 FL OZ (355 mL) ①

Contains Alcohol 0.5% 228-06122-1 REV GC-0419

Drug Facts

TAMPER-EVIDENT: Do not use if imprinted neckband is missing or broken.

Active ingredients (in each 10 mL dose)

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

Purposes

Uses relieves • heartburn • sour stomach
• acid indigestion • 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
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.

Keep out of the reach of children.

Directions

- shake well before use
- **adults and children 12 years and over:** 10 mL – 20 mL (1-2 doses) between meals, at bedtime or as directed by a doctor
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hypromellose, microcrystalline cellulose, propylparaben,
purified water, saccharin sodium, sorbitol solution

Questions? 1-888-287-1915

DISTRIBUTED BY:
Walmart Inc.,
Bentonville, AR 72716

*This product is not manufactured
or distributed by Johnson &
Johnson, owner of the registered
trademark Maximum Strength
Mylanta®.

REV GC-0219 928-06122-1

545902



MAXIMUM STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-681
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE -	ALUMINUM	800 mg

UNII:5QB0T2IUN0)	HYDROXIDE	in 10 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	800 mg in 10 mL
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	80 mg in 10 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-681-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	05/01/2011	

Labeler - WAL-MART STORES, INC. (051957769)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(49035-681)

Revised: 11/2023

WAL-MART STORES, INC.