REGULAR STRENGTH ANTACID- aluminum hydroxide, magnesium hydroxide, dimethicone suspension VALU MERCHANDISERS COMPANY

ALS (BC) ANTACID original

Active ingredients (in each 5 mL teaspoonful)

Aluminum hydroxide 200 mg (equivalent to dried gel, USP) Magnesium hydroxide 200 mg Simethicone 20mg

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 **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 and older: take 2 to 4 teaspoonfuls between meals, at bedtime, or as directed by a doctor
- do not take more than 24 teaspoonfuls in 24 hours or use the maximum dosage for more than 2 weeks
- children under 12 years: ask a doctor

Other information

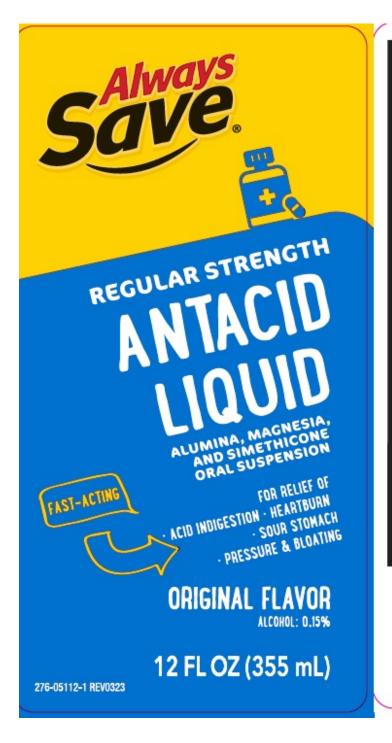
- each 5 mL teaspoonful contains: magnesium 85 mg, sodium 3 mg
- store at room temperature
- protect from freezing
- keep tightly closed

Inactive ingredients

benzyl alcohol, butylparaben, flavor (contains alcohol), hydroxyethylcellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments? 1-800-540-3765

package Label



Drug Facts

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

Active ingredients (in each 5 mL teaspoonful)

Purpose

Aluminum hydroxide 200 mg (equivalent to dried gel, USP) Magnesium hydroxide 200 mg Simethicone 20 mg

Uses relieves • heartburn

- sour stomach
- acid indigestion
 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 for more than 2 weeks 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 2 to 4 teaspoonfuls between meals, at bedtime or as directed by a doctor
- do not exceed 24 teaspoonfuls in 24 hours 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 85 mg, sodium 3 mg
- store at room temperature
- protect from freezing keep tightly closed

Inactive ingredients benzyl alcohol, butylparaben, flavor (contains alcohol), hydroxyethylcellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or comments? 1-800-540-3765

PROUDLY DISTRIBUTED BY: ASSOCIATED WHOLESALE GROCERS, INC. KANSAS CETY, KANSAS 66106

FOR PRODUCT INFORMATION PLEASE WISIT: ALWAYSSAVEBRAND_COM/66703





REGULAR STRENGTH ANTACID

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information HUMAN OTC DRUG NDC:63941-629 **Product Type** Item Code (Source) **Route of Administration** ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE -	ALUMINUM	200 mg	

UNII:5QB0T2IUN0)	HYDROXIDE	in 5 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	200 mg in 5 mL
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	20 mg in 5 mL

Inactive Ingredients				
Ingredient Name	Strength			
BENZYL ALCOHOL (UNII: LKG8494WBH)				
BUTYLPARABEN (UNII: 3QPI1U3FV8)				
HYDROXYETHYL CELLULOSE (2000 CPS AT 1%) (UNII: S38J6RZ N16)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
WATER (UNII: 059QF0KO0R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SORBITOL (UNII: 506T60A25R)				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1		355 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2017		
2		710 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2017	09/01/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	07/01/2017	

Labeler - VALU MERCHANDISERS COMPANY (868703513)

Registrant - GCP Laboratories (965480861)

Establishment				
Name	Address	ID/FEI	Business Operations	
GCP Laboratories		965480861	manufacture(63941-629), pack(63941-629)	

Revised: 11/2023 VALU MERCHANDISERS COMPANY