

SUPREME ANTACID- calcium carbonate, magnesium hydroxide suspension
FAMILY DOLLAR SERVICES INC

FD antacid supreme cherry

Active ingredients (in each 5 mL teaspoonful)

Calcium carbonate 400 mg
Magnesium hydroxide 135 mg

Purpose

Antacid

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- upset stomach associated with these symptoms
- overindulgence in food and drink

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 18 teaspoonfuls in a 24 hour period or use the maximum dosage for more than 2 weeks
- children under 12 years: ask a doctor

Other information

- **each 5 mL teaspoonful contains:** calcium 160mg, magnesium 55 mg
- refrigerate after opening to preserve freshness and purity
- do not freeze
- keep tightly closed

Inactive ingredients

benzyl alcohol, flavor, hydroxyethylcellulose, purified water, saccharin sodium, simethicone emulsion, sorbic acid, sorbitol solution, xanthan gum

Questions or comments?

1-800-540-3765

package Label



COMPARE TO
THE ACTIVE
INGREDIENTS
OF MYLANTA®
SUPREME*

ANTACID SUPREME

RELIEVES:

- Heartburn
- Acid indigestion
- Sour stomach



**CHERRY
FLAVOR**

12 FL OZ
(355 mL)

NDC 55319-624-12

Drug Facts

TAMPER EVIDENT: Do not use if the breakaway band on cap is broken or missing.

Active ingredients

(in each 5 mL teaspoonful)

Active ingredients	Purpose
Calcium carbonate 400 mg	Antacid
Magnesium hydroxide 135 mg	Antacid

Uses

- relieves • acid indigestion • sour stomach • heartburn
• overindulgence in food and drink
• upset stomach due to these symptoms

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 as needed, at bedtime or as directed by a doctor
- do not exceed 18 teaspoonfuls in a 24 hour period or use the maximum dosage for more than 2 weeks
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Questions or comments? 1-800-540-3765

*This product is not manufactured or distributed by the owner of the registered trademark Mylanta®.

DISTRIBUTED BY:
MIDWOOD BRANDS, LLC
500 VOLVO PKWY
CHESAPEAKE, VA 23320 USA

PRODUCT OF USA

NOT 100% SATISFIED?

Return within 30 days to any Family Dollar store for a refund (with receipt) or exchange.

939-07322-5 REV GC-0523



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SUPREME ANTACID

calcium carbonate, magnesium hydroxide suspension

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-624
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6Z B,	CALCIUM	400 mg

CARBONATE ION - UNII:7UJQ5OPE7D)	CARBONATE	in 5 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	135 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SORBIC ACID (UNII: X045WJ989B)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
XANTHAN GUM (UNII: TTV12P4NEE)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY (cherry)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-624-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	06/01/2017	

Labeler - FAMILY DOLLAR SERVICES INC (024472631)

Registrant - GCP Laboratories (965480861)