

CALCIUM ANTACID- calcium carbonate tablet, chewable
Allegiant Health

Drug Facts

Active ingredient (in each tablet)

Calcium Carbonate 500 mg

Purpose

Antacid

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- upset stomach associated with these symptoms

Warnings

Ask a doctor or pharmacist before use if you

- have kidney disease
- are taking a prescription drug. Antacids may interact with certain prescription drugs

When using this product

- do not take more than 15 tablets in 24 hours
- if pregnant, do not take more than 10 tablets in 24 hours
- do not use the maximum dosage for more than 2 weeks except under the advice and supervision of a doctor

Keep out of reach of children.

Directions

- **adults and children 12 years of age and over:** chew 2 - 4 tablets as symptoms occur, or as directed by a doctor
- do not take for symptoms that persist for more than 2 weeks unless advised by a doctor

Other information

- **each tablet contains:** calcium 200mg
- store at 15°-30°C (59°-86°F)

- do not use if imprinted safety seal under cap is broken or missing

Inactive ingredients

assorted flavors, D&C yellow #10 aluminum lake, dextrose, FD&C blue #1 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin

Package/Label Principal Display Panel

SODIUM FREE

Health A2Z®

CALCIUM ANTACID

REGULAR STRENGTH

500 mg

Calcium Supplement

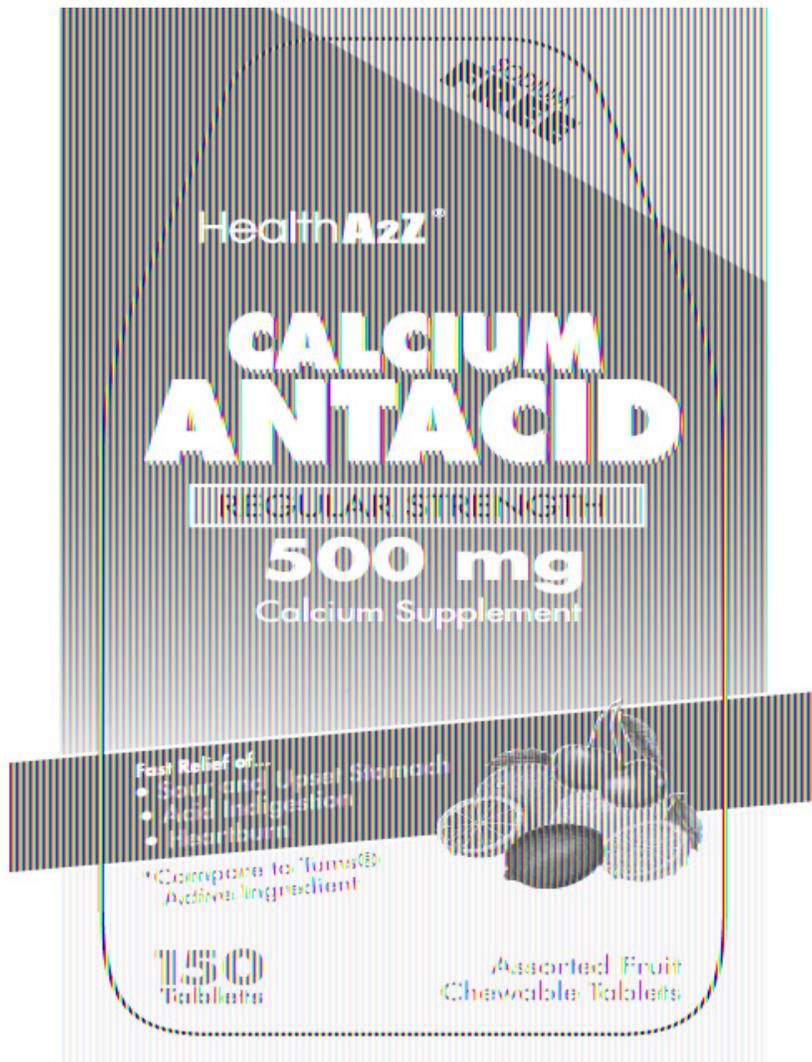
Fast Relief of...

- Sour & Upset Stomach
- Acid Indigestion
- Heartburn

*Compare to Tums® Active Ingredient

Assorted Fruit

Chewable Tablets



Antacid

Drug

Active Ingredient (in each tablet):
 Calcium carbonate
 USP 500mg

Uses: relieves heartburn, acid indigestion, and gas with these symptoms

Warnings:
 As a doctor or pharmacist may have kidney disease, antacids may interact with it.

When using this product in:
 in 24 hours; if pregnant, do not use for 24 hours; do not use the medicine for more than 2 weeks except under the advice of a doctor.

Keep out of reach of children.

Directions: in adults and over 12 years of age: chew 2 to 4 tablets as directed by a doctor; do not take for more than 2 weeks unless advised by a doctor.

Other information:
 in each tablet contains:
 calcium 200mg
 in store at 15° - 30°C (59° - 86°F)
 do not use if imprinted safety seal under cap is broken or missing.

Inactive ingredients:
 FD&C blue no. 1 aluminum lake, dextrose, FD&C red no. 40 aluminum lake, FD&C yellow no. 5 aluminum lake, magnesium stearate, polyethylene glycol 400, polyethylene glycol 600, polyethylene glycol 800, polyethylene glycol 1000, polyethylene glycol 1500, polyethylene glycol 2000, polyethylene glycol 3000, polyethylene glycol 4000, polyethylene glycol 6000, polyethylene glycol 8000, polyethylene glycol 10000, polyethylene glycol 15000, polyethylene glycol 20000, polyethylene glycol 30000, polyethylene glycol 40000, polyethylene glycol 60000, polyethylene glycol 80000, polyethylene glycol 100000.

This product is not manufactured by GSK Group, owned and registered trademark: Tums®.

Manufactured by:
 Allergiant Health
 Deer Park, NY 11729

CALCIUM ANTACID

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69168-219
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	500 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

DEXTROSE (UNII: IY9XDZ35W2)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Product Characteristics

Color	yellow, orange, pink, green	Score	no score
Shape	ROUND	Size	16mm
Flavor	LIME, CHERRY, ORANGE, LEMON	Imprint Code	AZ;024
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69168-219-50	50 in 1 BOTTLE; Type 0: Not a Combination Product	12/16/2014	05/15/2023
2	NDC:69168-219-02	150 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	12/16/2014	

Labeler - Allegiant Health (079501930)

Revised: 11/2017

Allegiant Health