CALCIUM CARBONATE CHEWABLE ANTACID- calcium carbonate tablet, chewable
NuCare Pharmaceuticals,Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active ingredient (in each tablet)

Calcium Carbonate 500 mg

Purpose

Antacid

Uses

- temporarily relieves: acid indigestion, heartburn, sour stomach

Warnings

Ask a doctor before use if you
• have kidney disease
• are taking prescription drugs; antacids may interact with certain prescription drugs

Stop use and ask a doctor if • symptoms last more than 2 weeks.

When using this product • do not use the maximum dosage for more than 2 weeks

If pregnant or breast feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

• do not exceed recommended dose
• adults and children 12 years of age and older: chew 2-4 tablets as symptoms occur
• repeat hourly if symptoms return, or as directed by a physician
• do not take more than 15 tablets in a 24 hour period
• children under 12 years: consult a doctor

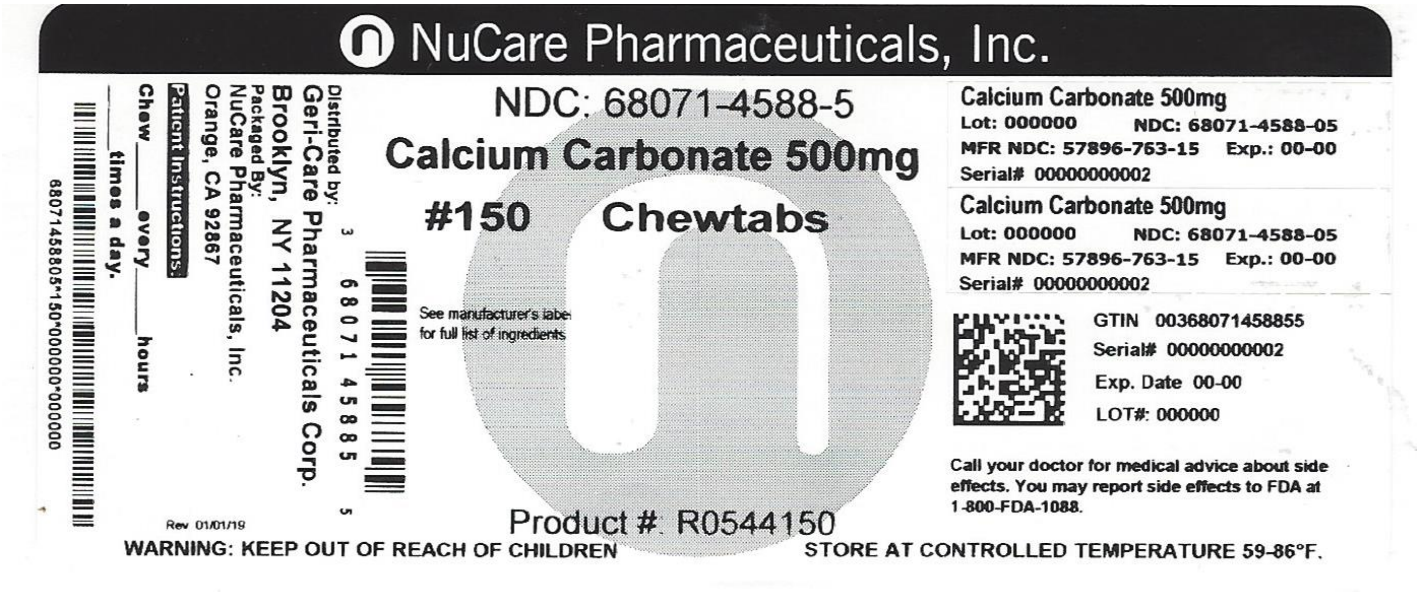
Other information

• **each tablet contains:** calcium 200 mg
• store at room temperature
• for institutional use only
• **Tamper Evident:** Do not use if imprinted seal under cap is missing or broken.

Inactive ingredients

assorted flavors, dextrose, D and C Yellow no. 10 lake, FD and C Blue no. 1 lake, FD and C Red no. 40 lake, FD and C Yellow no. 6 lake, magnesium stearate, maltodextrin. May also contain cellulose, FD and C Yellow no. 5 lake (tartrazine), stearic acid, sugar.

Package label



CALCIUM CARBONATE CHEWABLE ANTACID

calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68071-4588(NDC:57896-763)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB, CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	500 mg

Inactive Ingredients	
Ingredient Name	**Strength**
DEXTROSE (UNII: IY9XDZ35W2)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	red, green, orange, yellow	**Score**	no score
Shape	ROUND	**Size**	14mm
Flavor	CHERRY, LIME, ORANGE, LEMON	**Imprint Code**	AZ;024

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4588-5	150 in 1 BOTTLE; Type 0: Not a Combination Product	10/05/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	01/01/2004	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-4588)

Revised: 10/2019 NuCare Pharmaceuticals,Inc.