

CALCIUM CARBONATE- calcium carbonate suspension

Atlantic Biologicals Corps

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.

Calcium Carbonate

Drug Facts

Active ingredient (in each 5 mL teaspoonful)

Calcium Carbonate 1250 mg (Equivalent to 500 mg elemental Calcium)

Purpose

Antacid

Uses

relieves:

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

Warnings

Ask a doctor or pharmacist before use if you are taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product, do not take more than 6 teaspoonfuls (30 mL) in a 24-hour period or use the maximum dosage for more than 2 weeks.

Keep out of reach of children.

Directions

- **Shake well before using.**
- Take 1 to 2 teaspoonfuls (5 to 10 mL) as symptoms occur, or as directed by a doctor.

Other information

- store at 20° to 25°C (68° to 77°F)
- do not freeze
- packaged with tamper evident seal

Inactive ingredients

calcium saccharin, citric acid, D&C Red No. 33, FD&C Red No. 40, flavoring, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sorbitol and xanthan gum.

Questions?

Call 1-800-845-8210.

CALCIUM CARBONATE SUSPENSION

17856-3117-30
CALCIUM CARBONATE
ORAL SUSPENSION 1250
MG/5 ML



See package insert for indications and dosage schedule



Store at 20° to 25°C (68° to 77°F) Do not freeze
Sugar free/ Alcohol FREE/ Sodium free
SHAKE WELL
KEEP OUT OF THE REACH OF CHILDREN

17856-3117-30

Dosage: 5 ML

CALCIUM CARBONATE

Qty: 72 Cups



GTIN: 00117856311706

S/N: 01288001

Exp: 07/27/21

Lot: 012880

OTC

Packaged by: Unit Dose Solutions
Morrsville, NC 27560

Distributed by: Atlantic Biologicals Corp.
Miami FL 33179

Rev. 09/19

Call to Reorder: 800.509.7592

CALCIUM CARBONATE

calcium carbonate suspension

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:17856-3117(NDC:0121-0766)

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	1250 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
Propylparaben (UNII: Z8IX2SC1OH)	
Methylparaben (UNII: A2I8C7HI9T)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Sorbitol (UNII: 506T60A25R)	
Glycerin (UNII: PDC6A3C0OX)	
Xanthan Gum (UNII: TTV12P4NEE)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
FD&C Red No. 40 (UNII: WZB9127XOA)	
D&C Red No. 33 (UNII: 9DBA0SBB0L)	
Saccharin Calcium (UNII: 5101OP7P2I)	
Water (UNII: 059QF0K00R)	

Product Characteristics

Color	PINK	Score	
Shape		Size	
Flavor	BUBBLE GUM	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17856-3117-3	5 mL in 1 CUP; Type 0: Not a Combination Product	01/28/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part331	12/01/2004	

Labeler - Atlantic Biologicals Corps (047437707)

Establishment

Name	Address	ID/FEI	Business Operations
Atlantic Biologicals Corps		047437707	RELABEL(17856-3117) , REPACK(17856-3117)

