# CALCIUM CARBONATE - calcium carbonate tablet Carilion Materials Management

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.

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#### Calcium Carbonate Tablets

#### **Active Ingredient**

(in each tablet)

Calcium carbonate 10 gr (648 mg)

#### **Purpose**

Antacid

#### Use

relieves

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

### **Warnings**

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

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

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

### Keep out of reach of children.

#### **Directions**

- take one to four tablets daily
- do not take more than 4 tablets in 24 hours
- do not use the maximum dosage for more than 2 weeks

#### Other Information

- Each tablet contains: calcium 260 mg
- store at room temperature 15°-30°C (59°-86°F)

#### **Inactive Ingredients**

croscarmellose sodium, magnesium sterate, maltodextrin, microcrystalline cellulose, stearic acid

#### **Questions or Comments**

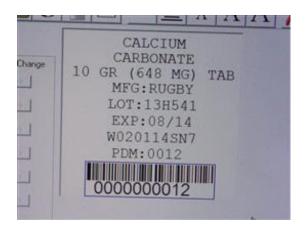
Call Monday – Friday, 9 am – 5 pm ET **1-800-645-2158**,

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL UNDER CAP IS BROKEN OR MISSING

This is a bulk pakage, dispense contents with a child-resistant closure in a tight, light-resistant container as defined in the USP.

Mfd for: Rugby Laboratories,Inc

#### **CALCIUM CARBONATE TABLET**



#### **CALCIUM CARBONATE**

calcium carbonate tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68151-0012(NDC:0536-3414)	

Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CALCIUM CARBONATE (UNII: H0 G9379 FGK) (CALCIUM CATION - UNII:2M83C4R6 ZB, CARBONATE ION - UNII:7UJQ5OPE7D)	CALCIUM CARBONATE	648 mg	

Inactive Ingredients		
Ingredient Name	Strength	
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MALTO DEXTRIN (UNII: 7CVR7L4A2D)		
CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)		
STEARIC ACID (UNII: 4ELV7Z65AP)		

Product Characteristics					
Color	WHITE (off white)	Score	no score		

Shape	ROUND	Size	11mm
Flavor		Imprint Code	AP;025
Contains			

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68151-0012-1	1 in 1 BOTTLE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part331	04/12/1999		

# Labeler - Carilion Materials Management (079239644)

## **Registrant -** Carilion Materials Management (079239644)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Carilion Materials Management		079239644	REPACK(68151-0012)	

Revised: 12/2012 Carilion Materials Management