# CALCIUM CARBONATE 1250 MG- calcium carbonate tablet CitraGen 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.

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#### Calcium Carbonate Tablets, USP 1250 mg

#### **Drug Facts**

#### Active ingredient (in each tablet)

Calcium carbonate 1250 mg

### **Purpose**

Antacid / Calcium Supplement

#### Uses

#### relieves:

- acid indigestion
- heartburn
- 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.

**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**

- take one to two tablets daily, or as directed by a doctor.
- 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 500 mg

store at room temperature 15° - 30° C (59° - 86° F)

#### **Inactive ingredients**

Hydrogenated Vegetable Oil, Maltodextrin and Microcrystalline Cellulose

#### Questions or comments?

Phone: +1-510-249-9066 (9AM-5PM PST, Mon-Fri); e-mail: info@citragenpharma.com

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

#### Manufactured by:

CitraGen Pharmaceuticals, Inc., Fremont, CA 94538.

www.citragenpharma.com

#### CitraGen Pharmaceuticals, Inc.

NDC: 70369-005-02 Rev. 08/18 R-00

Calcium Carbonate Tablets, USP 1250 mg

#### **ANTACID / Calcium Supplement**

500 Tablets

THIS PACKAGE IS FOR HOUSEHOLDS WITHOUT YOUNG CHILDREN



#### CitraGen Pharmaceuticals, Inc.

NDC: 70369-005-03

Rev. 08/18 R-00

Calcium Carbonate Tablets, USP 1250 mg

**ANTACID / Calcium Supplement** 

100 Tablets



NDC:70369-005-03

# Calcium Carbonate

Tablets, USP 1250 mg

ANTACID / CALCIUM SUPPLEMENT

100 Tablets

Drug Facts Active Ingredient (in each tablet) Purpose Calcium Carbonate 1250 mg .Antacid Uses: relieves: ●acid indigestion ●heartburn ●sour stomach ●upset stomach associated with these symptoms CALCIUM SUPPLEMENT USES: As a daily source of extra calcium DIRECTIONS: Take 1 tablets twice daily Ask a doctor or pharmacist before use if you are taking a prescription Supplement Facts drug. Antacids may interact with certain prescription drugs. Serving Size: 1 Tablets Stop use and ask a doctor if symptoms last more than 2 weeks. Servings Per Container: 100 If pregnant or breast-feeding, ask a health professional before use. Amount Per Serving %Daily Value\* Keep out of reach of children. Calories 1 Calcium 500 mg Directions Percent Daily Values are based on a 2000 calorie die •take one to two tablets daily, or as directed by a doctor. Ingredients: Calcium carbonate, hydrogenated do not take more than 4 tablets in 24 hours getable oil, maltodextrin, and microcrystalline do not use the maximum dosage for more than 2 weeks Other information: •each tablet contains: Calcium 500 mg •store at room temperature 15° - 30°C (59° - 86°F) Inactive ingredients: Hydrogenated Vegetable Oil, Maltodextrin, and Microcrystalline Cellulose. Questions or comments? Phone: +1-510-249-9066, 9AM-5PM PST, Mon-Fri; e-mail: info@citragenpharma.com

uticals, Inc.

#### CALCIUM CARBONATE 1250 MG

calcium carbonate tablet

#### **Product Information**

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:70369-005

Route of Administration ORAL

#### **Active Ingredient/Active Moiety**

Ingredient Name Basis of Strength CALCIUM CARBONATE (UNII: H0 G9379 FGK) (CALCIUM CATION - UNII: 2M83C4R6ZB) CALCIUM CARBONATE 1250 mg

## **Inactive Ingredients**

Ingredient Name Strength
HYDRO GENATED COTTONSEED O IL (UNII: Z82Y2C65EA)

MALTO DEXTRIN (UNII: 7CVR7L4A2D)

CELLULOSE, MICRO CRYSTALLINE (UNII: OP1R32D61U)

#### **Product Characteristics**

ColorwhiteScoreno scoreShapeCAPSULESize18 mmFlavorImprint CodeCG005ContainsContains

### **Packaging**

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#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>				
1	NDC:70369-005-02	500 in 1 BOTTLE; Type 0: Not a Combination Product	09/26/2018					
2	NDC:70369-005-03	100 in 1 BOTTLE; Type 0: Not a Combination Product	09/26/2018					

Marketing Information								
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
OTC monograph final	part331	09/26/2018						

# Labeler - CitraGen Pharmaceuticals, Inc. (024949457)

## **Registrant** - CitraGen Pharmaceuticals, Inc. (024949457)

Establishment								
Name	Address	ID/FEI	Business Operations					
CitraGen Pharmaceuticals, Inc.		024949457	manufacture(70369-005)					

Revised: 12/2020 CitraGen Pharmaceuticals, Inc.