DE LA CRUZ SODIUM BICARBONATE ANTACID- sodium bicarbonate powder
DLC Laboratories, Inc.

DE LA CRUZ ® SODIUM BICARBONATE ANTACID

Drug Facts

Active ingredient (in each dose = 1/2 teaspoon)

Sodium bicarbonate USP 2,616 mg

Purpose

Antacid

Uses

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

Warnings

FOR ORAL USE ONLY

STOMACH WARNING

TO AVOID SERIOUS INJURY, DO NOT TAKE UNTIL POWDER IS COMPLETELY DISSOLVED. IT IS VERY IMPORTANT NOT TO TAKE THIS PRODUCT WHEN OVERLY FULL FROM FOOD OR DRINK.

Do not use in children under 12 years of age.

Ask a doctor before use if you have
- a sodium-restricted diet.

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

Stop use and ask a doctor if
- you have used the maximum dosage for 2 weeks
- severe stomach pain occurs after using this product

If pregnant or breastfeeding, 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 immediately.

Directions

- take ½ level teaspoon in ½ glass (4 fl. oz.) of water every 2 hours up to maximum dosage or as directed by a doctor. Dissolve completely in water before drinking.

Age	Maximum Dosage
Adults 60 years and over	Do not exceed three doses of 1/2 teaspoon in a 24 hour period.
Adults and children 12 years and over	Do not exceed six doses of 1/2 teaspoon in a 24 hour period.
Children under 12 years	Do not use.

Do not exceed recommended dosage. See warnings.

Other information

- **each 1/2 teaspoon dose contains:** sodium 716 mg

Distributed by:
De La Cruz Products
A Division of DLC Laboratories, Inc.
Paramount, CA 90723 USA

Questions

1-800-858-3889

Inactive Ingredient

none

PRINCIPAL DISPLAY PANEL - 113 g Bottle Label

De La Cruz ®

OVER
40
DOSES

Sodium
Bicarbonate

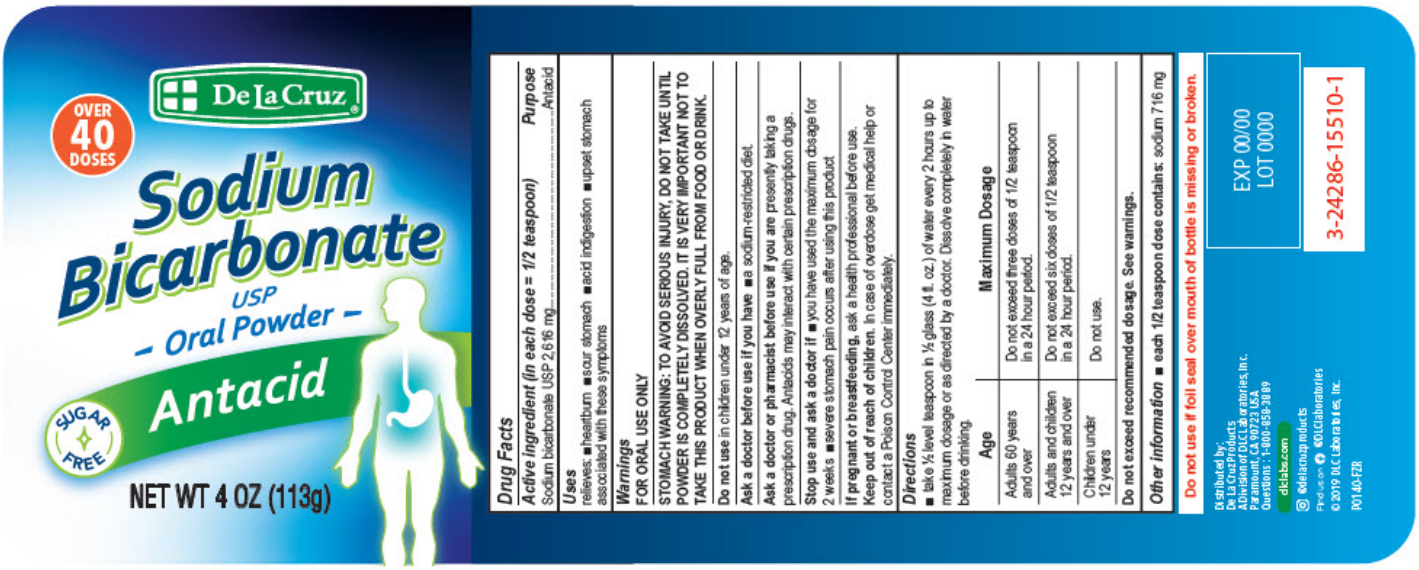
USP

- Oral Powder -

SUGAR
FREE

Antacid

NET WT 4 OZ (113g)



DE LA CRUZ SODIUM BICARBONATE ANTACID

sodium bicarbonate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1537
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (BICARBONATE ION - UNII:HN1ZRA3Q20)	SODIUM BICARBONATE	2.6 g in 2.6 g

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286-1537-7	113 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/25/2013	
2	NDC:24286-1537-8	226 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/22/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	09/25/2013	

Labeler - DLC Laboratories, Inc. (093351930)

Establishment

Name	Address	ID/FEI	Business Operations
DLC Laboratories, Inc.		093351930	label(24286-1537) , manufacture(24286-1537)

Revised: 10/2024 DLC Laboratories, Inc.