# NUTRALOX- calcium carbonate tablet, chewable HART Health

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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#### **NUTRALOX**

Active Ingredient (in each tablet): Calcium Carbonate 420 mg

Purpose: Antacid

**Uses:** Temporarily relieves

- heartburn
- sour stomach
- acid indigestion
- upset stomach

## **Warnings:**

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

Stop use and ask a doctor if symptoms last 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. 1-800-222-1222

#### **Directions:**

#### Do not take more than directed

Adults and children 12 years of age and over:

- thoroughly chew 2 tablets every 3 to 4 hours as symptoms occur
- do not take more than 16 tablets in 24 hours unless directed by a doctor
- do not use the maximum dose for more than 2 weeks

Children under 12 years of age: ask a doctor

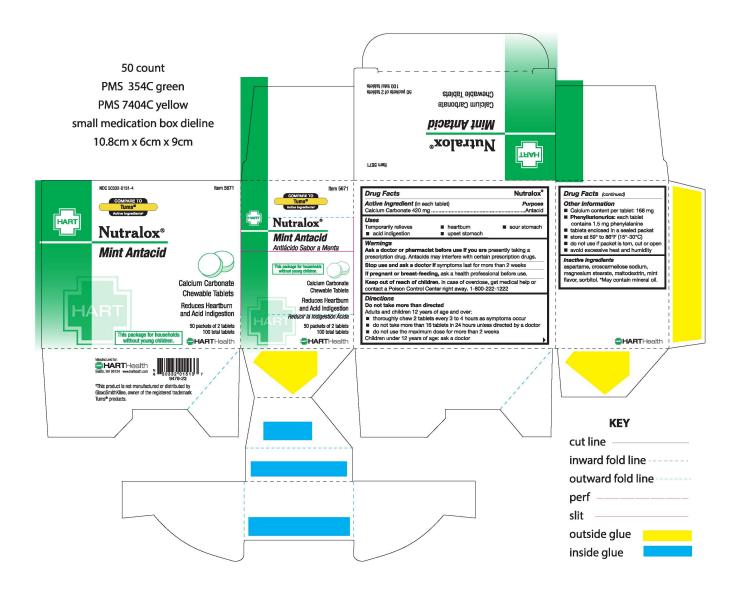
## **□Other information:**

- Calcium content per tablet: 168mg
- phenylketonurics: each tablet may contain 1.5 mg phenylalanine
- tablets enclosed in a sealed packet
- do not use if packet is torn, cut or open
- store at 59° to 86°F (15°-30°C)
- avoid excessive heat and humidity

# Inactive ingredients

aspartame, croscarmellose sodium, magnesium stearate, maltodextrin, mint flavor, sorbitol

May contain mineral oil



#### **NUTRALOX**

calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50332-0151
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII: 2M83C4R6ZB)	CALCIUM CARBONATE	420 mg	

Inactive Ingredients			
Ingredient Name	Strength		
ASPARTAME (UNII: Z0H242BBR1)			
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			
MINERAL OIL (UNII: T5L8T28FGP)			
SORBITOL (UNII: 506T60A25R)			

Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	11mm	
Flavor		Imprint Code	AZ;036	
Contains				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50332- 0151-3	20 in 1 BOX, UNIT-DOSE	01/20/1987	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:50332- 0151-4	50 in 1 BOX, UNIT-DOSE	01/20/1987	
2		2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:50332- 0151-7	125 in 1 BOX, UNIT-DOSE	01/20/1987	
3		2 in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:50332- 0151-8	250 in 1 BOX, UNIT-DOSE	01/20/1987	
4		2 in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:50332- 0151-1	500 in 1 BOX, UNIT-DOSE	01/20/1987	
5		2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part331	01/20/1987			

# Labeler - HART Health (069560969)

Revised: 8/2023 HART Health