

TUMS NATURALS- calcium carbonate tablet, chewable
Haleon US Holdings LLC

Drug Facts

Active ingredient (per tablet)

Calcium carbonate 1000 mg

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

presently taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product

- do not take more than 7 tablets in 24 hours
- If pregnant do not take more than 5 tablets in 24 hours
- do not use the maximum dosage for more than 2 weeks, except under the advice and supervision of a doctor

Keep out of reach of children.

Directions

- **adults and children 12 years of age and over:** chew 2-3 tablets as symptoms occur, or as directed by a doctor. Chew or crush tablets completely before swallowing.
- do not take for symptoms that persist for more than 2 weeks unless advised by a doctor

Other information

- **each tablet contains:** elemental calcium 400 mg
- do not store above 25°C (77°F)

Inactive ingredients (Black Cherry & Watermelon)

calcium stearate, carmine, dextrose, flavors, gum arabic

Inactive ingredients (Coconut Pineapple)

calcium stearate, dextrose, flavor, gum arabic

Questions or comments?

1-800-897-7535

Generic Section

Do not use if printed inner safety seal under cap is broken or missing.

Gluten-Free

Non-GMO

Principal Display Panel (Black Cherry & Watermelon)

GOES TO WORK IN SECONDS!

CALCIUM CARBONATE

TUMS

ANTACID

Naturals

NOARTIFICIAL FLAVORS OR DYES

Black Cherry & Watermelon

ULTRA STRENGTH **1000**

56 CHEWABLE TABLETS

HALEON

GOES TO WORK IN SECONDS!

CALCIUM CARBONATE

TUMS

ANTACID

Naturals



*Black Cherry
& Watermelon*

**NO
ARTIFICIAL
FLAVORS
OR DYES**

**ULTRA 1000
STRENGTH**

56 CHEWABLE TABLETS

214177

Principal Display Panel (Coconut Pineapple)

gsk

GOES TO WORK IN SECONDS!

CALCIUM CARBONATE

TUMS

ANTACID

NEW

Naturals

NO ARTIFICIAL FLAVORS OR DYES

Coconut Pineapple

ULTRA STRENGTH **1000**

56 CHEWABLE TABLETS



TUMS NATURALS

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-1326
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB,	CALCIUM	1000 mg

CARBONATE ION - UNII:7UJQ5OPE7D)		CARBONATE	1000 mg	
Inactive Ingredients				
Ingredient Name		Strength		
CALCIUM STEARATE (UNII: 776XM7047L)				
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)				
ACACIA (UNII: 5C5403N26O)				
Product Characteristics				
Color	PINK (Reddish-pink) , PINK (Light pink)	Score	no score	
Shape	ROUND	Size	16mm	
Flavor	CHERRY (Black Cherry) , WATERMELON	Imprint Code	TUMS;N	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-1326-01	56 in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
2	NDC:0135-1326-03	190 in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	
3	NDC:0135-1326-02	10 in 1 BOTTLE; Type 0: Not a Combination Product	03/12/2021	
4	NDC:0135-1326-05	2 in 1 POUCH; Type 0: Not a Combination Product	06/10/2021	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M001	10/15/2020		

TUMS NATURALS			
calcium carbonate tablet, chewable			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0135-1328
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	1000 mg

Inactive Ingredients

Ingredient Name	Strength
CALCIUM STEARATE (UNII: 776XM7047L)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
ACACIA (UNII: 5C5403N26O)	

Product Characteristics

Color	WHITE	Score	no score
Shape	ROUND	Size	16mm
Flavor	COCONUT, PINEAPPLE	Imprint Code	TUMS;N
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-1328-01	56 in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	10/15/2020	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 2/2025

Haleon US Holdings LLC