

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

Gluten-Free

Non-GMO

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

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Dist. by: GSK CH,

Warren, NJ 07059

Principal Display Panel (Black Cherry & Watermelon)

gsk

GOES TO WORK IN SECONDS!

CALCIUM CARBONATE

TUMS

ANTACID

NEW

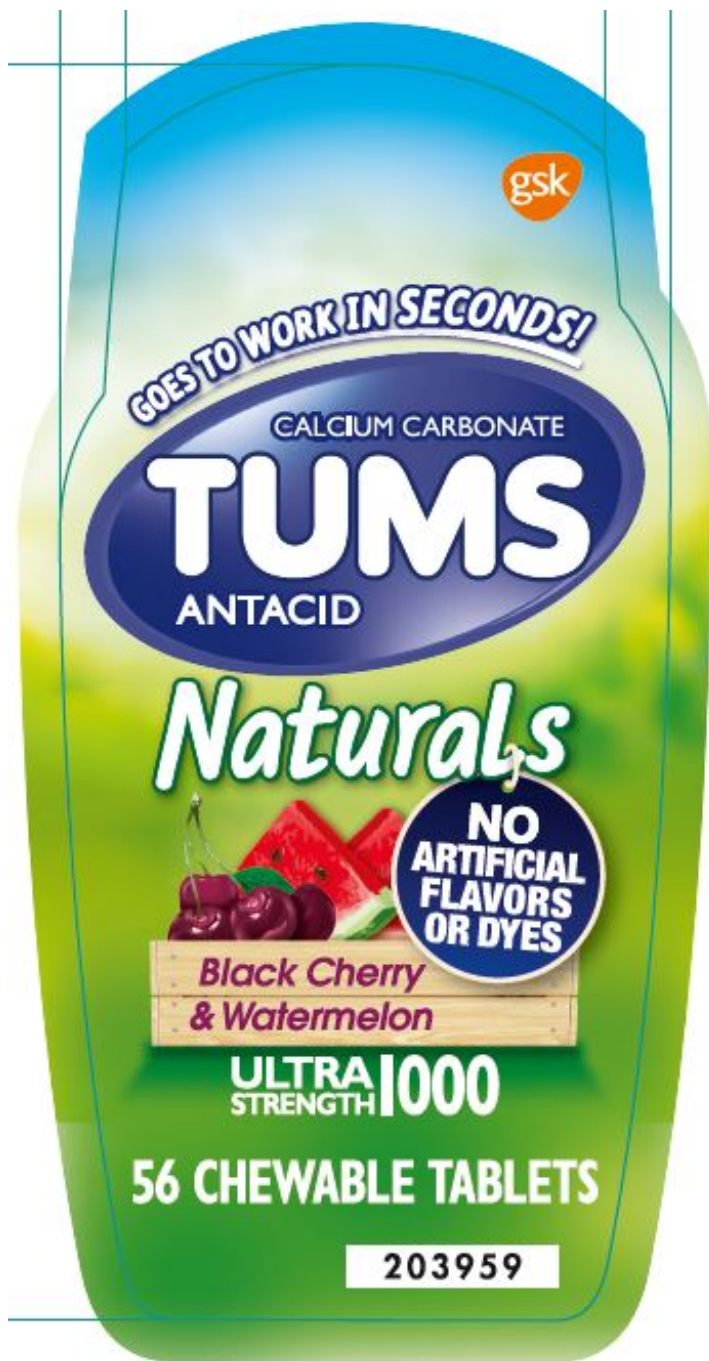
Naturals

NOARTIFICIAL FLAVORS OR DYES

Black Cherry & Watermelon

ULTRA STRENGTH **1000**

56 CHEWABLE TABLETS



Principal Display Panel (Coconut Pineapple)

gsk

GOES TO WORK IN SECONDS!

CALCIUM CARBONATE

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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, 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	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: 3/2024

Haleon US Holdings LLC