TUMS ULTRA- 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, magnesium 10 mg
- store below 30°C (86°F)
- contains FD&C Yellow No. 5 (tartrazine) as a color additive

Inactive ingredients

corn starch, crospovidone, dextrose, FD&C red #40 lake, FD&C yellow #5 (tartrazine) lake, FD&C yellow #6 lake, flavor, magnesium stearate, maltodextrin, sucrose, talc

Questions?

1-800-897-7535

Additional Information

Safety sealed – Do not use if printed inner seal beneath cap is missing or broken.

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Dist. by: **GSK CH**
Warren, NJ 07059

Gluten-Free

Principal Display Panel

GOES TO WORK IN SECONDS!

CALCIUM CARBONATE
TUMS
ANTACID

Tropical
Fruit

72 CHEWABLE TABLETS

ULTRA
STRENGTH ***1000***

202113 Back Label
202112 Front Label


GOES TO WORK IN SECONDS!
gsk
CALCIUM CARBONATE
TUMS
ANTACID
Tropical Fruit
72 CHEWABLE TABLETS
ULTRA STRENGTH 1000
202112

TUMS ULTRA

calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0135-0553
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CALCIUM CARBONATE (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ5OPE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN (UNII: O8232NY3SJ)	
CROSPOVIDONE, UNSPECIFIED (UNII: 2S7830E561)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FD&C YELLOW NO. 5 ALUMINUM LAKE (UNII: JQ6BLH9FR7)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics			
Color	WHITE (Off-white) , YELLOW, ORANGE, RED	**Score**	no score
Shape	ROUND	**Size**	19mm
Flavor	TROPICAL FRUIT PUNCH, STRAWBERRY (Strawberry-banana) , PINEAPPLE (Pineapple-orange) , ORANGE (Mandarin orange)	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:0135-0553-01	72 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	03/01/2023	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 2/2024 Haleon US Holdings LLC