TUMS- calcium carbonate tablet
Haleon US Holdings LLC

Drug Facts

Active ingredient (per tablet)

Calcium Carbonate USP 500 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

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

When using this product

- do not take more than 15 tablets in 24 hours
- if pregnant do not take more than 10 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 2-4 tablets as symptoms occur, or as directed by a doctor
- do not take for symptoms that persist for more than 2 weeks unless advised by a doctor

Other information

- **each tablet contains:** elemental calcium 200mg
- store below 30ºC (86ºF)

Inactive ingredients (assorted fruit)

Adipic acid, corn starch, FD&C blue #1 lake, FD&C red #40 lake, FD&C yellow #5 (tartrazine) lake, FD&C yellow #6 lake, flavors, mineral oil, sodium polyphosphate, sucrose, talc

Inactive ingredient (peppermint)

corn starch, flavor, mineral oil, sodium polyphosphate, sucrose, talc

Questions?

1-800-897-7535

weekdays

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

Gluten-Free

www.tums.com

GlaxoSmithKline

Moon Twp, PA 15108

PRINCIPAL DISPLAY PANEL

NDC 0135-0071-27

TUMS®

CALCIUM CARBONATE

ANTACID

REGULAR STRENGTH 500

Goes to Work in Seconds!

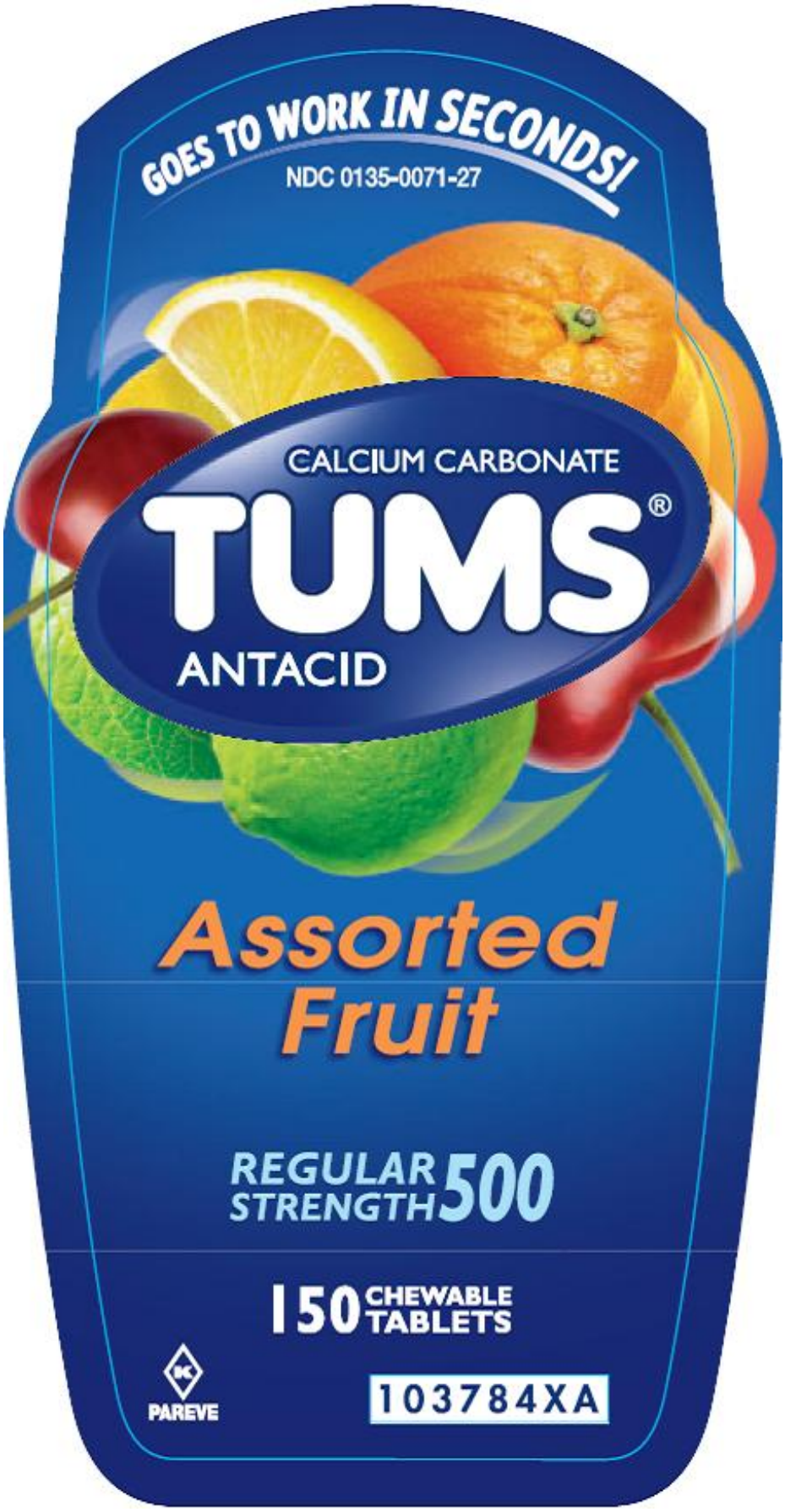
Assorted Fruit

150 CHEWABLE TABLETS

PAREVE

Front Label: 103784XA

Back Label: 103783XA



Principal Display Panel

NDC 0135-0070-27

TUMS®

CALCIUM CARBONATE

ANTACID

REGULAR STRENGTH 500

Goes to Work in Seconds!

Peppermint

150 CHEWABLE TABLETS

PAREVE

Front Label: 103781XA

Back Label: 103780XA


GOES TO WORK IN SECONDS!
NDC 0135-0070-27
CALCIUM CARBONATE
TUMS®
ANTACID
Peppermint
REGULAR STRENGTH 500
150 CHEWABLE TABLETS
PAREVE
103781XA

TUMS

calcium carbonate tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0135-0071
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	500 mg

Inactive Ingredients

Ingredient Name	Strength
ADIPIC ACID (UNII: 76A0JE0FKJ)	
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MINERAL OIL (UNII: T5L8T28FGP)	
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	PINK (orange, yellow, green)	**Score**	no score
Shape	ROUND	**Size**	16mm
Flavor	CHERRY (assorted fruit, orange, lemon, lime)	**Imprint Code**	TUMS
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0071-27	150 in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2010	
2	NDC:0135-0071-48	12 in 1 PACKAGE; Type 0: Not a Combination Product	03/10/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	03/10/2010	

TUMS

calcium carbonate tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0135-0070
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	500 mg

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Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
MINERAL OIL (UNII: T5L8T28FGP)	
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	
SUCROSE (UNII: C151H8M554)	
TALC (UNII: 7SEV7J4R1U)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	ROUND	**Size**	16mm
Flavor	PEPPERMINT	**Imprint Code**	TUMS
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0070-03	3 in 1 CELLO PACK	03/10/2010	
1		12 in 1 PACKAGE; Type 0: Not a Combination Product		
2	NDC:0135-0070-27	150 in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2010	
3	NDC:0135-0070-48	12 in 1 PACKAGE; Type 0: Not a Combination Product	03/10/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC Monograph Drug	M001	03/10/2010	

Labeler - Haleon US Holdings LLC (079944263)

Revised: 3/2024 Haleon US Holdings LLC