ANTACID- calcium carbonate tablet
Target Corporation

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.

Drug Facts

Active ingredient (per tablet)

Calcium Carbonate USP 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
- do not take for symptoms that persist for more than 2 weeks unless advised by a doctor

Other information

- **each tablet contains:** elemental calcium 400mg, sodium 2mg (Assorted Fruit)
- store below 30°C (86°F)

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 ingredients (Peppermint)

sucrose, calcium carbonate, corn starch, talc, mineral oil, natural flavor, sodium polyphosphate

Questions?

Call 1-888-367-7919 weekdays

Principal Display Panel

NDC 11673-481-41

ultra strength

antacid 1000

calcium carbonate

Compare to Tums Ultra®*

relief of heartburn

acid indigestion and

upset stomach

associated with these symptoms

up&up®

ASSORTED FRUIT FLAVOR

160 CHEWABLE TABLETS

naturally and artificially flavored

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

*TUMS and TUMS ULTRA are registered trademarks of the GSK group of companies.

245 05 0457 R00 ID285492

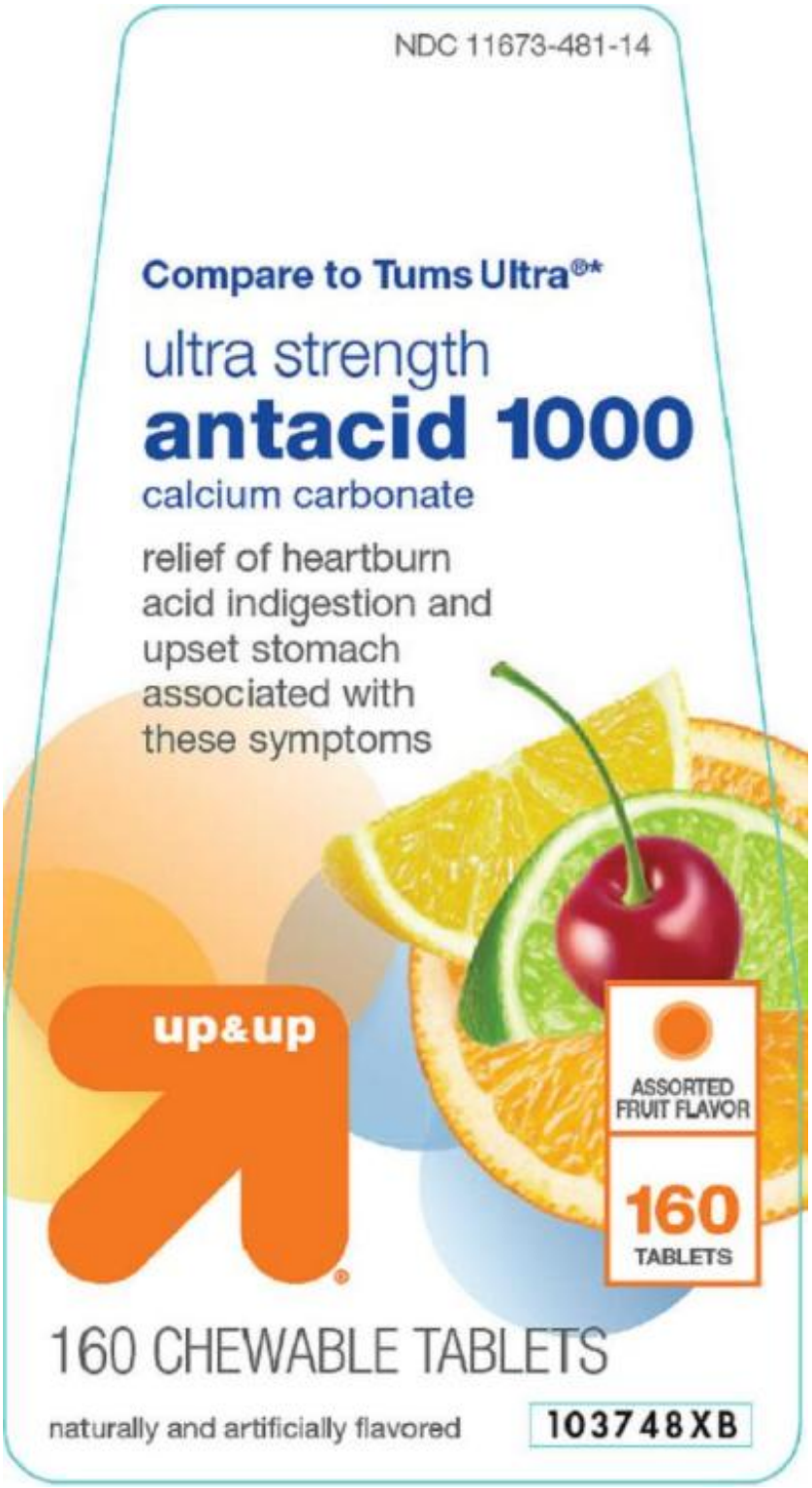
Dist. by Target Corp., Mpls., MN 55403

Shop Target.com

GLUTEN FREE

103748XB (front label)

103747XB (back label)



Principal Display Panel

NDC 11673-722-01

antacid tablets

ultra strength antacid/calcium supplement

calcium carbonate, 1,000 mg

Compare to active ingredient in TUMS® Ultra*

up & up®

160 CHEWABLE TABLETS

naturally flavored

peppermint flavor

CALCIUM SUPPLEMENT

USES: As a daily source of extra calcium.

DIRECTIONS: Chew 2 tablets once or twice daily with a meal.

Supplement Facts

Serving Size: 2 Tablets

Servings Per Container: 80

Amount Per Serving	**% Daily Value**		
Calories 10			
Sugars 3g			
Calcium 800mg	80%		
Sodium 10mg	Less than 1%		

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

*TUMS and TUMS ULTRA are registered trademarks of the GlaxoSmithKline group of companies.

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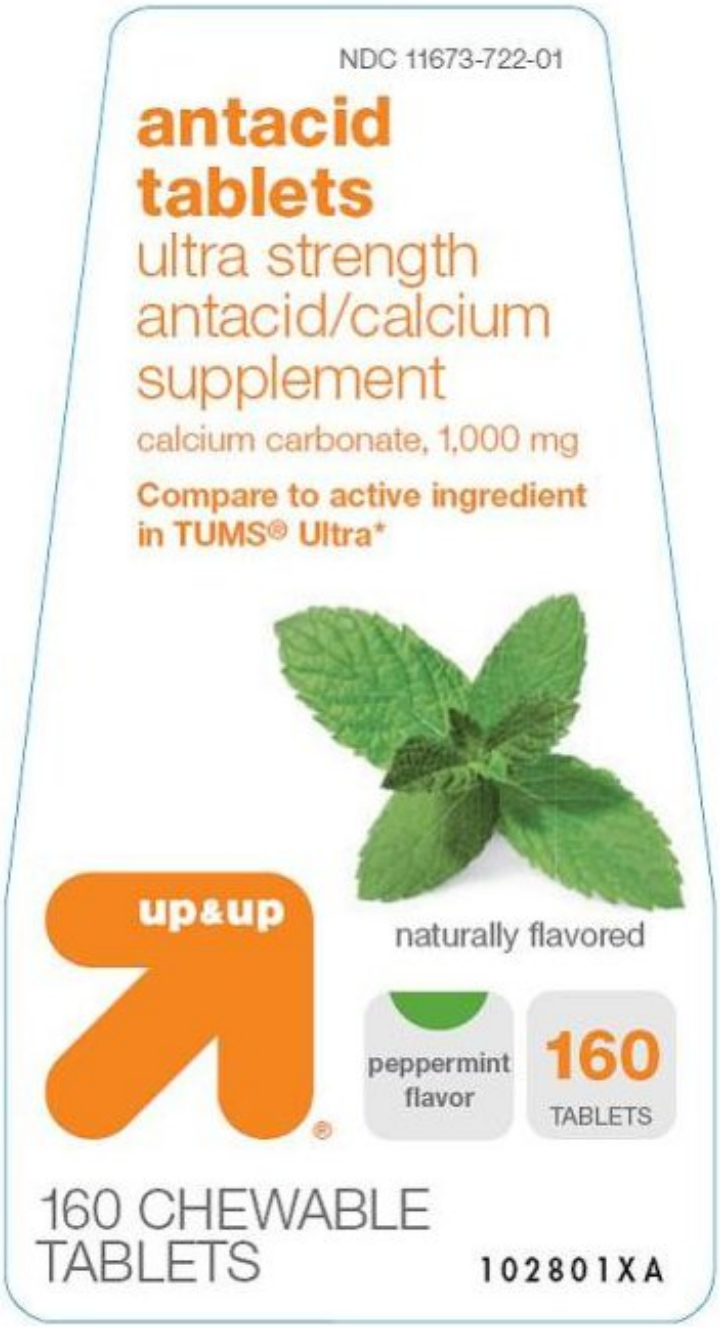
Front Label: 102801XA

Back Label: 102802XA



ANTACID

calcium carbonate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-481
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**
SUCROSE (UNII: C151H8M554)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
MINERAL OIL (UNII: T5L8T28FGP)	
ADIPIC ACID (UNII: 76A0JE0FKJ)	
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALUMINUM OXIDE (UNII: LMI26O6933)	

Product Characteristics			
Color	PINK (orange, yellow, green)	**Score**	no score
Shape	ROUND	**Size**	19mm
Flavor	CHERRY (assorted fruit, orange, lemon, lime)	**Imprint Code**	LH20
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-481-14	160 in 1 BOTTLE; Type 0: Not a Combination Product	02/15/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	02/15/2010	

ANTACID

calcium carbonate tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-722
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**
SUCROSE (UNII: C151H8M554)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
MINERAL OIL (UNII: T5L8T28FGP)	
SODIUM POLYMETAPHOSPHATE (UNII: P1BM4ZH95L)	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	ROUND	**Size**	19mm
Flavor	PEPPERMINT	**Imprint Code**	LH20
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-722-01	160 in 1 BOTTLE; Type 0: Not a Combination Product	03/08/2013	06/30/2018

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	03/08/2013	06/30/2018

Labeler - Target Corporation (006961700)

Revised: 10/2016 Target Corporation