

UP AND UP ULTRA STRENGTH ANTACID ASSORTED FRUIT- antacid tablets tablet, chewable

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.

up and up ultra strength antacid 1000 chewable tablets

Active ingredient (per tablet)

Calcium Carbonate USP 1000mg

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

store below 30°C (86°F).

Inactive ingredients

adipic acid, blue 1, dextrose, flavors, magnesium stearate, maltodextrin, red 27, starch, sucralose, yellow 6 lake, yellow 10 lake.

Questions?

1-866-467-2748

Package/Label Principal Display Panel

NDC 11673-903-17

Compare to Tums Ultra®*

ultra strength

antacid 1000

calcium carbonate

relief of heartburn, acid indigestion, sour stomach and upset stomach associated with these symptoms

ASSORTED FRUIT FLAVOR

160 TABLETS

160 CHEWABLE TABLETS

naturally and artificially flavored

245 05 0457 R00

IDC-000274-01-007

Distributed by

Target Corporation

Minneapolis, MN 55403

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GLUTEN FREE

Safety sealed: Do not use if printed seal under cap is torn or missing

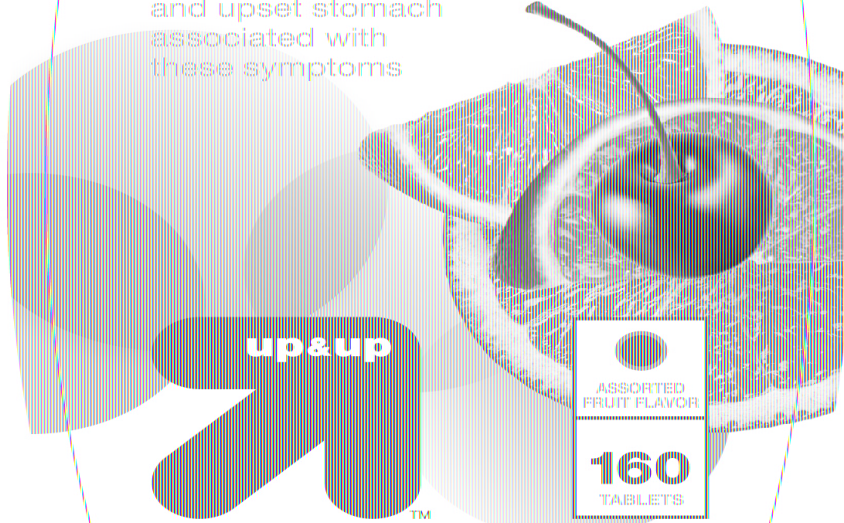
100% satisfaction guaranteed or your money back.

NDC 11673-903-17

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160 CHEWABLE TABLETS
naturally and artificially flavored



103AF160TGTLF

**Active ingredi
(per tablet)**

Calcium Carbonate

Uses relieves
• acid indigestion
• upset stomach

Warnings

Ask a doctor or pharmacist
taking a prescription drug
prescription drugs.

When using this product

- do not take more than
 - if pregnant do not take
 - do not use the maximum
- except under the advice

Keep out of reach of children

Directions

- adults and children 12 years
 - symptoms occur, or as directed
 - do not take for symptoms
- advised by a doctor

Other information

- each tablet contains 1000 mg
- store below 30°C (86°F).

Inactive ingredients:
magnesium stearate, malted
sorghum, yellow 10 lake.

Questions? 1-866-467-7233

*This product is not manufactured by the registered trademark owner of the registered trademark.

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**100% satisfaction
guaranteed or
your money back.**

103AF160TGTLB

UP AND UP ULTRA STRENGTH ANTACID ASSORTED FRUIT

antacid tablets tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-903
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CARBONATE ION - UNII:7UJQ50PE7D, CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

Inactive Ingredients

Ingredient Name	Strength
ADIPIIC ACID (UNII: 76A0JE0FKJ)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DEXTROSE (UNII: IY9XDZ35W2)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

Product Characteristics

Color	RED, GREEN, YELLOW, ORANGE	Score	no score
Shape	ROUND	Size	19mm
Flavor	FRUIT (assorted fruit)	Imprint Code	RP103
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-903-17	160 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	10/22/2015	

Labeler - Target Corporation (006961700)

Revised: 1/2015

Target Corporation