

SHOPRITE ANTACID- calcium carbonate tablet, chewable
Wakefern Food 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.

ShopRite Antacid Drug Facts

Active ingredient (in each 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:calcium 430mg
- store at 20-25°C (68-77°F)

- does not meet USP for assay

Inactive ingredients

citric acid, dextrose, FD&C blue #1 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #5 aluminum lake (tartrazine), FD&C yellow #6 aluminum lake, flavors, glycerin, magnesium stearate, maltodextrin, mineral oil

Questions?

1-800-SHOPRITE

Principal Display Panel

Compare to: Active Ingredient in Tums® Ultra Strength 1000

ULTRA STRENGTH

ANTACID

Calcium Carbonate 1000 mg

STARTS WORKING IN SECONDS

Chewable Tablets

Assorted Fruit Flavors

actual size

72 TABLETS



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DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING

*This product is not manufactured or distributed by GlaxoSmithKline, distributor of Tums® Ultra Strength 1000.

QUALITY GUARANTEE

Distributed By: Wakefern Food Corp.
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 Keasbey, NJ 08832 ©2016

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: 59523 88 B2

SHOPRITE ANTACID

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41190-595
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:7UJQ50PE7D)	CALCIUM CARBONATE	1000 mg

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
MINERAL OIL (UNII: T5L8T28FGP)	
DEXTROSE (UNII: IY9XDZ35W2)	
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)	

Product Characteristics

Color	YELLOW, GREEN, RED, ORANGE	Score	no score
Shape	ROUND	Size	20mm
Flavor	FRUIT (lime, orange, cherry, lemon)	Imprint Code	L595
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41190-595-23	72 in 1 BOTTLE; Type 0: Not a Combination Product	01/30/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	01/30/2017	

Labeler - Wakefern Food Corporation (069722418)

Revised: 12/2018

Wakefern Food Corporation