QC ANTACID CALCIUM ASSORTED BERRY- calcium carbonate tablet, chewable

CHAIN DRUG MARKETING ASSOCIATION INC.

Ultra Strength Assorted Berry Antacid Calcium Supplement Drug Facts

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

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
- do not use the maximum dosage for more than 2 weeks.

Keep out of reach of children.

Directions

chew 2-3 tablets as symptoms occur, or as directed by a doctor

Other Information

- Store below 30°C (86°F).
- Keep the container tightly closed

• For calcium and magnesium content see Supplement Facts.

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

Inactive ingredients

adipic acid, blue 1 lake, dextrose, flavor, flavors, magnesium stearate, maltodextrin, red 40 lake, starch, sucralose.

Package/Label Principal Display Panel

QUALITY CHOICE®

NDC# 63868-049-72

Compare To Tums® Ultra Strength active ingredient*

ULTRA STRENGTH

Antacid Tablets

Antacid Calcium Supplement

CALCIUM CARBONATE 1000 mg

Relieves:

- Upset Stomach
- Heartburn
- Acid Indigestion

Assorted Berry

GLUTEN FREE

72 CHEWABLE TABLETS

K PAREVE

QC SATISFACTION GURANTEED

© Distributed by: C.D.M.A., Inc.

43157 W Nine Mile

Novi, MI 48376-0995

www.qualitychoice.com

Questions: 248-449-9300

*This product is not manufactured or distributed by GlaxoSmithKline, the distributor of Tums $^{@}$.



Drug Facts

Active ingredient Purpose (per tablet)

Calcium Carbonate USP 1000mg.....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 completely as symptoms occur, or as directed by a doctor. Do not swallow tablets whole.

• 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 at room temperature. Keep the container tightly closed.

Inactive ingredients adipic acid, dextrose, FD&C Blue No. 1, FD&C Red No. 40, flavors, magnesium stearate, maltodextrin, starch, sucralose

Questions or comments? 1-866-467-2748 DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSIN

GLUTEN-FREE

"This product is not manufactured or distributed by GlaxoSmithKline LLC, owner of the registered trademark, Ultra Strength Tums".





EXP.: 103AB72QCLB

OC ANTACID CALCIUM ASSORTED BERRY

calcium carbonate tablet, chewable

Product Information

HUMAN OTC DRUG **Product Type** Item Code (Source) NDC:63868-049

ORAL Route of Administration

Active Ingredient/Active Moiety

Active ingredient/Active Molety		
Ingredient Name	Basis of Strength	Strength
CALCIUM CARRONATE (UNII: HOG9379EGK) (CALCIUM CATION - UNII: 2M83C4R67B	CALCIUM	

Inactive Ingredients		
Ingredient Name	Strength	
ADIPIC ACID (UNII: 76A0JE0FKJ)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ 35W2)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
STARCH, CORN (UNII: O8232NY3SJ)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		

Product Characteristics			
Color	PINK (blue-pink) , BLUE	Score	no score
Shape	ROUND	Size	17mm
Flavor	BERRY (assorted berry)	Imprint Code	RP103
Contains			

ı	Pa	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:63868-049- 72	72 in 1 BOTTLE; Type 0: Not a Combination Product	04/16/2019	

Marketing In	Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M001	04/16/2019		

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC. (011920774)

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