QUALITY CHOICE ULTRA STRENGTH ANTACID ASSORTED FRUIT ULTRA STRENGTH ASSORTED FRUIT ANTACID- calcium carbonate tablet, chewable
Chain Drug Marketing Association Inc.

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.

QC Ultra Strength Assorted Fruit Antacid

ACTIVE INGREDIENT (in each tablet)

Calcium carbonate 1000 mg

PURPOSE

Antacid

USE(S)

relieves:
- acid indigestion
- heartburn

WARNINGS

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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 a 24-hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a doctor.

IF PREGNANT OR/BREASTFEEDING,

ask a health professional before use.

KEEP OUT OF REACH OF CHILDREN

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DIRECTIONS

- **adults and children 12 years of age and over:**
- wet in mouth before chewing
- chew 2-3 tablets as symptoms occur, or as directed by a doctor.

OTHER INFORMATION

- **each tablet contains:** elemental calcium 400 mg, magnesium 10 mg
- do not use if printed seal under cap is torn or missing.
- store below 30°C (86°F).

INACTIVE INGREDIENTS

adipic acid, corn starch, crospovidone, D&C red 27 lake, D&C red 30 lake, D&C yellow 10 lake, dextrose, FD&C blue 1 lake, FD&C yellow 6 lake, flavors, magnesium stearate, maltodextrin, sucrose, talc.

PRINCIPAL DISPLAY PANEL

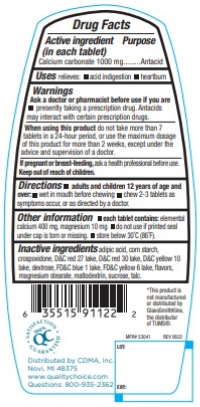
NDC 63868-161-72
QC

Quality Choice
***Compare to the active ingredient in Ultra Strength Tums®**
Ultra Strength
Antacid Tablets
Calcium Carbonate 1000 mg

Relieves:
Heartburn

Acid Indigestion
Assorted Fruit

72 Chewable Tablets



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calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-161
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1000 mg

Inactive Ingredients	
Ingredient Name	**Strength**
ADIPIC ACID (UNII: 76A0JE0FKJ)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSPOVIDONE (UNII: 68401960MK)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SUCROSE (UNII: C151H8M554)	

TALC (UNII: 7SEV7J4R1U)	
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)	
D&C RED NO. 30 (UNII: 2S42T2808B)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics

Color	ORANGE, YELLOW, GREEN, PINK (reddish pink)	**Score**	no score
Shape	ROUND	**Size**	17mm
Flavor	CHERRY, LEMON, LIME, ORANGE	**Imprint Code**	G171
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-161-72	72 in 1 BOTTLE; Type 0: Not a Combination Product	08/16/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part331	08/16/2022	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Revised: 8/2022 Chain Drug Marketing Association Inc.