# QCH ULTRA STRENGTH ANTACID 171AF- calcium carbonate tablet, chewable Chain Drug Marketing Association Inc.

-----

# QCH Ultra Strength Antacid Assorted Fruit 171AF

## **ACTIVE INGREDIENT (in each tablet)**

Calcium carbonate 1000 mg

#### **PURPOSE**

**Antacid** 

# USE(S)

relieves:

- acid indigestion
- heartburn

#### 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 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**

.

#### **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 10mg
- 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

NDC83324-127-72

QC

**QUALITY CHOICE** 

\*Compare to the Active Ingredient in Ultra Strength Tums®

# Ultra Strength Antacid Tablets

Calcium Carbonate 1000mg

#### **RELIEVES:**

Heartburn

**Acid Indigestion** 

#### **Assorted Fruit**

**72** Chewable Tablets









# **QCH ULTRA STRENGTH ANTACID 171AF**

calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-127
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	
STARCH, CORN (UNII: O8232NY3SJ)		
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
SUCROSE (UNII: C151H8M554)		
TALC (UNII: 7SEV7J4R1U)		
CROSPOVIDONE (UNII: 2S7830E561)		
ADIPIC ACID (UNII: 76A0JE0FKJ)		
FD&C BLUE NO. 1 Aluminum lake (UNII: J9EQA3S2JM)		
FD&C YELLOW NO. 6 ALUMINUM LAKE (UNII: GYP6Z2JR6Q)		
D&C RED NO. 27 ALUMINUM LAKE (UNII: ZK64F7XSTX)		
D&C RED NO. 30 ALUMINUM LAKE (UNII: GE75M6YV5W)		
D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6)		

Product Characteristics			
Color	PINK, ORANGE, YELLOW, GREEN	Score	no score
Shape	ROUND	Size	18mm
Flavor	CHERRY, LEMON, LIME, ORANGE	Imprint Code	G171
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:83324-127- 72	72 in 1 BOTTLE; Type 0: Not a Combination Product	08/03/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	08/03/2024	

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Guardian Drug Company		119210276	MANUFACTURE(83324-127)	

Revised: 8/2024

Chain Drug Marketing Association Inc.