QUALITY CHOICE ULTRA STRENGTH ANTACID ASSORTED BERRY ULTRA STRENGTH ANTACID ASSORTED BERRY- 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.

Quality Choice Ultra Strength Antacid Assorted Berry

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 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, dextrose, FD&C blue 1 lake, FD&C red 40 lake, flavors, magnesium stearate, maltodextrin, sucrose, talc.

PRINCIPAL DISPLAY PANEL

NDC 63868-127-72 **QC**

Quality Choice *Compare to the Active ingredient in Ultra Strength Tums®

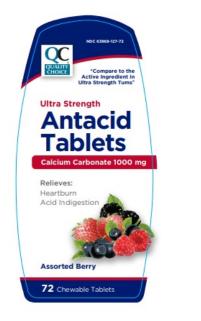
Ultra Strength **Antacid Tablets** Calcium Carbonate 1000 mg

Relieves:

Heartburn Acid Indigestion

Assorted Berry

72 Chewable Tablets





QUALITY CHOICE ULTRA STRENGTH ANTACID ASSORTED BERRY ULTRA STRENGTH ANTACID ASSORTED BERRY

calcium carbonate tablet, chewable

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Sour	ce)	NDC:6386	8-127	
Route of Administration	ORAL					
Active Ingredient/Active	Moiety					
Ing	Basis of Strength		Strength			
CALCIUM CARBONATE (UNII: H0 UNII:2M83C4R6ZB)	G9379FGK) (CALCIUM CATIOI	N -	CALCIUM CA	RBONATE	1000 mg	
Inactive Ingredients						
Ingredient Name					Strength	
ADIPIC ACID (UNII: 76A0JE0FKJ)						
STARCH, CORN (UNII: 08232NY3	SJ)					

		TIFIED FORM (UNII: IY9XDZ 35W2)					
		luminum lake (UNII: J9EQA3S2JM)					
FD&C RED NO.	40 (l	JNII: WZ B9127XOA)					
MAGNESIUM S	FEAR	ATE (UNII: 70097M6I30)					
MALTODEXTRI	N (UNI	II: 7CVR7L4A2D)					
SUCROSE (UNII:		· ·					
TALC (UNII: 7SE	V7J4R	10)					
Product Ch	arac	teristics					
Color	BLUE	, PINK, PINK (PINKISH BLUE)			Score		no score
Shape	ROUI	ND		Size			17mm
Flavor	DEDD					da	G171
FIAVUI	DEKR	RY (STRAWBERRY, RASPBERRY, WDBERRY)			Imprint Co	ue	61/1
	DERF	(Y (STRAWBERRY, RASPBERRY, WIDBERRY)			Imprint Co	ue	6171
Contains	DERF			Markoti			
Contains Packaging		Package Description		Marketiı Da	ng Start	Mark	eting End Date
Contains Packaging # Item Cod	e 27- 7				ng Start	Mark	eting End
Contains Packaging # Item Cod 1 NDC:63868-1	e 27- 7	Package Description 22 in 1 BOTTLE; Type 0: Not a Combination		Da	ng Start	Mark	eting End
Contains Packaging # Item Cod 1 NDC:63868-1 72	e 27- 7 F	Package Description 22 in 1 BOTTLE; Type 0: Not a Combination		Da	ng Start	Mark	eting End
Contains Packaging # Item Cod 1 NDC:63868-1 72	e 27- 7 F	Package Description 22 in 1 BOTTLE; Type 0: Not a Combination Product	07	Da /19/2022 Marke	ng Start	Mark	eting End Date
Contains Packaging # Item Cod 1 NDC:63868-1 72 Marketin Marketin	e 27- 7 g Ir	Package Description 22 in 1 BOTTLE; Type 0: Not a Combination Product Information Application Number or Monogra	07	Da /19/2022 Marke	ng Start te ting Start Date	Mark	eting End Date keting End

Labeler - Chain Drug Marketing Association Inc. (011920774)

Revised: 7/2022

Chain Drug Marketing Association Inc.