

**REGULAR STRENGTH ASSORTED FLAVORS- antacid calcium carbonate tablet,  
chewable  
AKRON PHARMA INC**

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**Regular Strength Antacid Calcium Carbonate 150 Chewable Tablets**

**Active ingredient**

Calcium Carbonate USP 500mg

**Purpose**

Antacid

**Uses**

relieves:

- heartburn
- sour stomach
- acid indigestion
- 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 15 tablets in 24 hours
- If pregnant do not take more than 10 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 2-4 tablets 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 200 mg**
- Store at room temperature. Keep the container tightly closed

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

### **Inactive ingredients**

adipic acid, D&C Red No.27 lake, D&C Yellow No.10 lake, dextrose, FD&C Blue No.1 lake, FD&C Yellow No.6 lake, flavors, magnesium stearate, maltodextrin, starch, sucralose.

**Questions or comments?** call toll-free 1-877-225-6999

### **Manufactured for:**

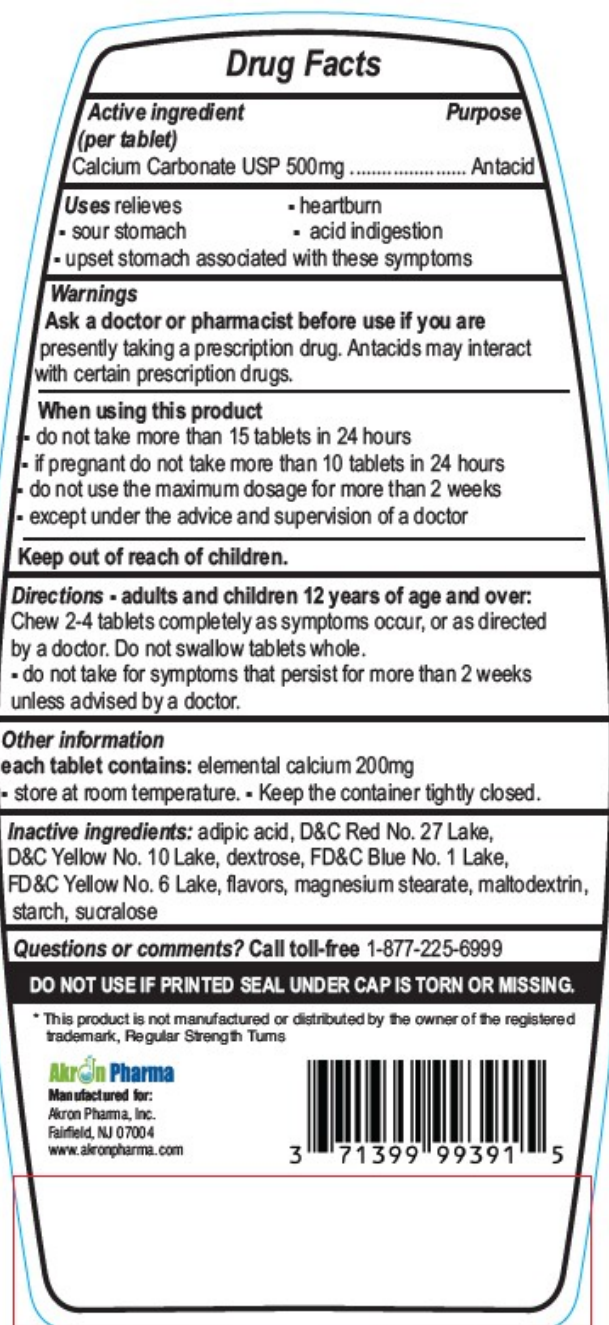
Akron Pharma Inc.,

Fairfield, NJ-07004

[www.akronpharma.com](http://www.akronpharma.com)

\*This product is not manufactured or distributed by GlaxoSmithKline LLC, owner of the registered trademark, Regular Strength Tums®.

### **Package/Label Principal Display Panel**



## REGULAR STRENGTH ASSORTED FLAVORS

antacid calcium carbonate tablet, chewable

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-9939
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION -	CALCIUM CARBONATE	500 mg

UNII:2M83C4R6Z B)			CALCIUM CARBONATE	500 mg
Inactive Ingredients				
Ingredient Name			Strength	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MALTODEXTRIN (UNII: 7CVR7L4A2D)				
STARCH, CORN (UNII: O8232NY3SJ)				
SUCRALOSE (UNII: 96K6UQ3ZD4)				
ADIPIC ACID (UNII: 76A0JE0FKJ)				
D&C RED NO. 27 (UNII: 2LRS185U6K)				
Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	16mm	
Flavor	FRUIT	Imprint Code	AP101	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-9939-1	150 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/23/2025	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001		09/23/2025	

**Labeler** - AKRON PHARMA INC (067878881)

Revised: 9/2025

AKRON PHARMA INC