

## **TARGET ANTACID FRUIT CHEWS- calcium carbonate tablet, chewable**

### **Target**

*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.*

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### **Calcium carbonate 750 mg**

### **Active ingredient (in each chewable tablet)**

Calcium carbonate 750 mg

### **Purpose**

Antacid

### **Uses Relieves**

- Heartburn
- Sour stomach
- Acid indigestion
- Upset stomach due to these symptoms

### **Warnings**

**Do not use** if you have ever had an allergic reaction to this product or any of its ingredients.

**Ask a doctor or pharmacist before use if you are** now taking a prescription drug. Antacids may interact with certain prescription drugs.

Do not take more than 5 chewable tablets in a 24-hour period, or use the maximum dosage for more than 2 weeks, except under the advice and supervision of a physician.

**When using this product** constipation may occur.

**If pregnant or breast-feeding**, ask a health professional before use.

**Keep out of reach of children.**

### **Directions**

Adults and children 12 years and older chew and swallow 1 to 2 chewable tablets every 2-4 hours as needed. Do not exceed 5 chewable tablets in 24 hours. Children under 12 years consult a doctor.

### **Other information**

- **Each chewable tablet contains:** calcium 300mg
- Contains soy
- Contains FD&C yellow No. 5 (tartrazine) as a color additive
- Store between 20° to 25°C (68° to 77 °F) in a dry place

### **Inactive ingredients**

Beeswax, carmine, carnauba wax, citric acid, corn starch, corn syrup, DL-alpha tocopherol, ethyl acetate, FD&C blue no. 1 aluminum lake, FD&C red no. 40 aluminum lake, FD&C yellow no. 5 lake (tartrazine), FD&C yellow no. 6, FD&C yellow no. 6 aluminum lake, gum arabic, hydrogenated coconut oil, maltodextrin, medium chain triglycerides, methyl paraben, modified corn starch, natural and artificial flavors, phosphoric acid, pregelatinized corn starch, propyl paraben, propylene glycol, purified water, shellac, sodium benzoate, sorbic acid, sorbitol, soy lecithin, soybean oil, sucrose and titanium dioxide.

**Questions?** Or to report an adverse event call **1-800-245-2898**, Monday - Friday, 9AM to 4PM EST

**Drug Facts (continued)****Inactive ingredients**

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\*This product is not manufactured or distributed by Bayer, owner of the registered trademark Alka-Seltzer® or any of its affiliates.

**GLUTEN FREE**

245 05 4859 R00 C-001227-01-007



Dist. by Target Corporation  
Minneapolis, MN 55403  
**Made in U.S.A. of U.S. and imported ingredients**  
TM & ©2020 Target Brands, Inc.

Compare to active ingredient in Alka-Seltzer® ReliefChews®\*

NDC 11673-045-90

extra strength  
**antacid chews**  
calcium carbonate, 750 mg

helps relieve heartburn, sour stomach and acid indigestion



90 CHEWABLE TABLETS

**Drug Facts**

**Active ingredient (in each chewable tablet)** Purpose  
Calcium carbonate 750 mg.....Antacid

**Uses** Relieves

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TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL, RUBBER CAP IS BROKEN OR MISSING. PN-218737

**TARGET ANTACID FRUIT CHEWS**

calcium carbonate tablet, chewable

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11673-051
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	750 mg

**Inactive Ingredients**

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

**Product Characteristics**

<b>Color</b>	orange (Orange, Pink, Yellow)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	14mm
<b>Flavor</b>	LEMON (Lemon, Orange, Strawberry)	<b>Imprint Code</b>	FC
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-051-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2019	



Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	11/01/2019	

**Labeler** - Target (006961700)

**Registrant** - Bestco LLC (002149136)

**Establishment**

Name	Address	ID/FEI	Business Operations
Bestco LLC		002149136	manufacture(11673-051)

Revised: 11/2019

Target