ORANGE ANTACID SOFTCHEW- calcium carbonate tablet, chewable
BestCo 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.

Calcium carbonate 1177 mg

Active ingredient (in each chew)

Calcium Carbonate 1177 mg

Purpose

Antacid

Uses: relieves

heartburn

sour stomach

acid indigestion

upset stomach due to these symptoms

Warnings

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

Do not take more than 6 chews in a 34-hour period, or use the maximum dosage for more than 2 weeks, except under the advice and supervision of a physician.

If you ae pregnant or nursing a baby, ask a doctor before using this product.

Keep out of reach of children.

Directions

Chew and swallow 2 to 3 chews, as symptoms occur or as directed by a physician.

Other information

Each chew contains: calcium 470 mg, mangesium 5mg

Contains milk and soy

Store between 20° to 25ºC (68º to 77ºF) in a dry place

inactive ingredients: Corn syrup, corn syrup solids, FD&C yellow #6 aluminum lake, glycerin, hydrogenated coconut oil, natural and artificial flavors, non-fat dairy milk powder, soy lecithin, and sucrose. Soybean oil and corn starch used as processing aids.

Questions? Or to report an adverse event call 1-800-245-2898, Monday - Friday, 8AM -4PM EST

ORANGE ANTACID SOFTCHEW

calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52642-021
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	1177 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SUCROSE (UNII: C151H8M554)	

Product Characteristics			
Color	orange	**Score**	no score

Shape	RECTANGLE	**Size**	22mm
Flavor	ORANGE	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52642-021-32	32 in 1 BAG; Type 0: Not a Combination Product	03/14/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	03/14/2014	

Labeler - BestCo Inc. (002149136)

Registrant - BestCo Inc. (002149136)

Establishment

Name	Address	ID/FEI	Business Operations
BestCo Inc.		002149136	manufacture(52642-021)