D-CAL- calcium carbonate tablet, chewable A&Z Pharmaceutical 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.

D-CAL (calcium carbonate) tablet, chewable

Active ingredient (in each tablet)

Calcium Carbonate 750 mg

Purpose

Antacid

Uses

relieves

- heartburn
- acid indigestion
- sour stomach
- upset stomach due to these symptoms

Warnings

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

When using this product

• do not take more than 10 tablets for adults and 5 tablets for children in a 24 hour period.

Keep out of reach of children.

Directions

- **Adults**: chew 2 tablets daily.
- **Children**: chew 1 tablet daily. If symptoms persist, ask a doctor.

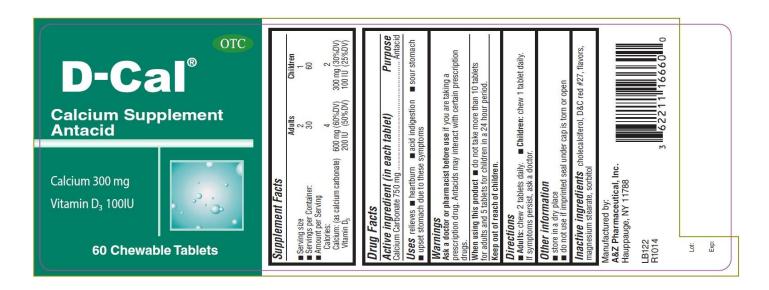
Other information

- Store in a dry place
- do not use if imprinted seal under cap is torn or open

Inactive ingredients

Cholecalciferol, D&C red #27, flavors, magnesium stearate, sorbitol

PRINCIPAL DISPLAY PANEL



D-CAL

calcium carbonate tablet, chewable

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62211-166
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CALCIUM CARBONATE (UNII: H0 G9379 FGK) (CALCIUM CATION - UNII:2M83C4R6 ZB)	CALCIUM CARBONATE	750 mg		

Inactive Ingredients		
Ingredient Name	Strength	
CHOLECALCIFEROL (UNII: 1C6V77QF41)		
D&C RED NO. 27 (UNII: 2LRS185U6K)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
SORBITOL (UNII: 506T60A25R)		

Product Characteristics			
Color	pink	Score	no score
Shape	RECTANGLE	Size	17mm
Flavor	TROPICAL FRUIT PUNCH	Imprint Code	DCAL;AZ
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:62211-166-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	09/17/1997	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	09/17/1997	

Labeler - A&Z Pharmaceutical Inc. (080225262)

Registrant - A&Z Pharmaceutical Inc. (080225262)

Establishment			
Name	Address	ID/FEI	Business Operations
A&Z Pharmaceutical Inc.		080225262	manufacture(62211-166)

Revised: 10/2019 A&Z Pharmaceutical Inc.