

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

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

OTC

D-Cal[®]

Calcium Supplement Antacid

Calcium 300 mg
Vitamin D₃ 100IU



60 Chewable Tablets

Supplement Facts	Adults	Children
■ Serving size	2	1
■ Servings per Container:	30	60
■ Amount per Serving	4	2
Calories:	600 mg (60%DV)	300 mg (30%DV)
Calcium: (as calcium carbonate)	200 IU (50%DV)	100 IU (25%DV)
Vitamin D ₃		

Drug Facts

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Manufactured by:
A&Z Pharmaceutical, Inc.
Hauppauge, NY 11788

LB122
R1014



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Lot:
Exp:

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: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6Z B)	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	
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	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: 9/2025

A&Z Pharmaceutical Inc.