

ANTACID- calcium carbonate tablet, chewable
Advance Pharmaceutical Inc.

Antacid

Active Ingredient

(in each tablet)

Calcium Carbonate 500 mg

Purpose

Antacid

Uses

relieves

- acid indigestion
- heartburn
- sour stomach

Warnings

Ask a doctor before use if you have

- kidney stones
- a calcium-restricted diet

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

Stop use and ask a doctor if symptoms last more than 2 weeks.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- chew 1-2 tablets as symptoms occurs.
- do not take more than 8 tablets in 24 hours
- do not use the maximum dosage for more than 2 weeks

Other Information

- **each tablet contains:** calcium 200 mg
- store at room temperature 15-30 °C (59-86 °F)

Inactive Ingredients

Al-lakes (D&C red #27, D&C yellow # 10, FD&C blue #1, FD&C yellow # 6), dextrose, flavors (cherry, lemon, lime, orange), magnesium stearate, maltodextrin

Questions or Comments

Call 631-981-4600, Monday-Friday, 8.30 am - 4.30 pm ET

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Manufactured by: Advance Pharmaceutical Inc. Holtsville, NY 11742

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

COMPARE TO
ACTIVE INGREDIENT IN
TUMS®*

NDC 17714-043-15

Calcium Rich ANTACID

Calcium Carbonate 500 mg

ASSORTED
FLAVORS

For fast relief of-

- Acid Indigestion
- Heartburn
- Sour & Upset Stomach



150
CHEWABLE TABLETS

SODIUM FREE

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY
SEAL UNDER CAP IS BROKEN OR MISSING.**

*Advance Pharmaceutical Inc. is not affiliated with the owner of the trademark Tums®.

Manufactured & Distributed by: Advance Pharmaceutical Inc., Holtsville, NY 11742



**Advance
Pharmaceutical Inc.**

Drug Facts

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AD0309



0.64"

0.375"



Calcium Carbonate 500 mg
Antacid tablets
NDC: 17714-043-15 - 150 COUNT

ANTACID

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17714-043
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:2M83C4R6ZB)		CALCIUM CARBONATE	500 mg
Inactive Ingredients			
Ingredient Name		Strength	
DEXTROSE (UNII: IY9XDZ35W2)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			
CHERRY (UNII: BUC5I9595W)			
Product Characteristics			
Color	yellow, pink, green	Score	score with uneven pieces
Shape	ROUND	Size	16mm
Flavor	LIME (Cherry,Lemon,Orange)	Imprint Code	AP;043
Contains			
Packaging			
#	Item Code	Package Description	Marketing Start Date
1	NDC:17714-043-15	150 in 1 BOTTLE; Type 0: Not a Combination Product	01/26/1996
2	NDC:17714-043-55	55 in 1 BOTTLE; Type 0: Not a Combination Product	01/26/1996
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	01/26/1996	

Labeler - Advance Pharmaceutical Inc. (078301063)

Registrant - Advance Pharmaceutical Inc. (078301063)

Establishment

Name	Address	ID/FEI	Business Operations
Advance Pharmaceutical Inc.		078301063	manufacture(17714-043)

Revised: 12/2024

Advance Pharmaceutical Inc.