

ANTACID- calcium carbonate tablet, chewable
Advance 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.

Calcium Rich

Active Ingredient
(in each tablet)

Calcium Carbonate 500 mg

Purpose

Antacid

Uses

relieves

- acid indigestion
- heartburn
- sour stomach
- upset stomach associated with these symptoms

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.

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

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

Cherry flavor, dextrose, 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


Advance
Pharmaceutical Inc.

Calcium Carbonate 500 mg

SODIUM FREE

CHEWABLE TABLETS

100



CHERRY
FLAVOR

For fast relief of:
• Acid Indigestion
• Heartburn
• Sour & Upset Stomach

Calcium Rich
ANTACID

NDC 17714-041-01

Drug Facts

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(in each tablet)**

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Manufactured by: Advance Pharmaceutical Inc.
Holtsville, NY 11742, USA



Lot No.:

Exp. Date:

NON-VARNISH AREA

LA1112

Calcium Carbonate 500 mg

Antacid tablets

NDC: 17714-041-01 – 100 COUNT

ANTACID

calcium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17714-041
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CARBONATE	500 mg

Inactive Ingredients

Ingredient Name	Strength
DEXTROSE (UNII: IY9XDZ35W2)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
CHERRY (UNII: BUC5I9595W)	

Product Characteristics

Color	white	Score	score with uneven pieces
Shape	ROUND	Size	16 mm
Flavor	CHERRY	Imprint Code	AP;041
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17714-041-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/30/2007	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part331	11/30/2007	

Labeler - Advance Pharmaceutical Inc. (078301063)

Registrant - Advance Pharmaceutical Inc. (078301063)

Establishment

Name	Address	ID/FEI	Business Operations
Advance Pharmaceutical Inc.		078301063	manufacture(17714-041)

Revised: 12/2019

Advance Pharmaceutical Inc.