

**ANTACID- calcium carbonate tablet, chewable**  
**Advance Pharmaceutical Inc.**

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**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

### Active ingredient (in each tablet)

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



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LA1112

Lot No.:

**NON-VARNISH AREA**

Exp. Date:

# Calcium Carbonate 500 mg

## Antacid tablets

**NDC: 17714-041-01 - 100 COUNT**

### ANTACID

calcium carbonate tablet, chewable

#### Product Information

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

#### Active Ingredient/Active Moiety

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

#### Inactive Ingredients

Ingredient Name	Strength
<b>DEXTROSE</b> (UNII: IY9XDZ35W2)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>CHERRY</b> (UNII: BUC5I9595W)	

#### Product Characteristics

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

#### 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 Drug	M001	11/30/2007	

**Labeler** - Advance Pharmaceutical Inc. (078301063)

**Registrant** - Advance Pharmaceutical Inc. (078301063)

**Establishment**

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

Revised: 12/2024

Advance Pharmaceutical Inc.