

QUALITY CHOICE ORIGINAL FLAVOR- aluminum hydroxide and magnesium carbonate tablet, chewable
CHAIN DRUG MARKETING ASSOCIATION INC.

Quality Choice Antacid Tablets Extra Strength Original Flavor

Active ingredients (in each tablet)

Aluminum hydroxide 160 mg

Magnesium carbonate 105 mg

Purpose

Antacid

Antacid

Uses

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

Warnings

Ask a doctor or pharmacist before use if you are

- taking a prescription drug. Antacids may interact with certain prescription drugs
- If you are on a sodium-restricted diet

When using this product

- do not take more than 16 tablets in 24 hours
- do not use the maximum dosage for more than 2 weeks

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center immediately at 1-800-222-1222

Directions

- chew 2 to 4 tablets four times a day or as directed by a doctor
- take after meals and at bedtime or as needed
- for best results follow by a half glass of water or other liquid
- DO NOT SWALLOW TABLETS WHOLE

Other information

- **each tablet contains:** magnesium 31 mg, sodium 19 mg
- Store at room temperature

Inactive ingredients

alginic acid, calcium stearate, compressible sugar, flavor, sodium bicarbonate, starch (corn), sucrose.

Questions or comments

1-866-467-2748

Principal Display Panel

QUALITY CHOICE®

NDC 83324-219-01

Compare to the active ingredients in Gaviscon® Extra Strength Antacid*

Antacid Tablets

Extra Strength

Fast Acting Heartburn Relief

Helps Keep Acid Down For Hours


Original Flavor

100 Chewable Tablets

100% QC SATISFACTION GUARANTEED

- **DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS BROKEN OR MISSING.**
- Distributed by CDMA., Inc.
- Novi, MI 48375
- www.qualitychoice.com
- Questions: 800-935-2362

*This product is not manufactured or distributed by GlaxoSmithKline, the distributor of Gaviscon® Extra Strength Antacid.



NDC 63868-996-01


Compare to **Gaviscon**
active ingredients*

Extra Strength Antacid Tablets

Foaming

Long Lasting Antacid Barrier
Fast Acting Heartburn Relief

**Original
Flavor**



100 Chewable Tablets

Do not use if printed under cap is broken or missing

Drug Facts

Active ingredients (in each tablet) Purpose
Aluminum hydroxide 160 mg.....Antacid
Magnesium carbonate 105 mg.....Antacid

Uses relieves ■ acid indigestion ■ heartburn ■ sour stomach ■ upset stomach associated with these symptoms

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Directions
■ chew 2 to 4 tablets four times a day or as directed by a doctor
■ take after meals and at bedtime or as needed
■ for best results follow by a half glass of water or other liquid
■ DO NOT SWALLOW WHOLE


Other information
■ each tablet contains: sodium 19 mg, magnesium 31 mg
■ store at up to 25°C (77°F) in a dry place

Inactive ingredients alginic acid, calcium stearate, compressible sugar, flavor, sodium bicarbonate, starch (corn), sucrose.

Questions or comments? 1-866-467-2748

*This product is not manufactured or distributed by GlaxoSmithKline, the distributor of Gaviscon® Extra Strength.

Distributed by CD.M.A., Inc.®
43157 W. Nine Mile
Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300



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UNVARNISHED AREA

EXP:

QUALITY CHOICE ORIGINAL FLAVOR

aluminum hydroxide and magnesium carbonate tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-219
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	160 mg
MAGNESIUM CARBONATE (UNII: 0E53J927NA) (CARBONATE ION - UNII:7UJQ5OPE7D)	MAGNESIUM CARBONATE	105 mg

Inactive Ingredients

Ingredient Name	Strength
ALGINIC ACID (UNII: 8C3Z4148WZ)	
CALCIUM STEARATE (UNII: 776XM7047L)	
CORN SYRUP (UNII: 9G5L16BK6N)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
STARCH, CORN (UNII: O8232NY3SJ)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	17mm
Flavor		Imprint Code	RP105
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-219-01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/29/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	07/29/2024	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC. (011920774)

Revised: 12/2025

CHAIN DRUG MARKETING ASSOCIATION INC.