GAVISCON- aluminum hydroxide and magnesium carbonate liquid
GlaxoSmithKline Consumer Healthcare Holdings (US) LLC

Drug Facts

Active ingredient (in each 15mL tablespoonful) Regular Strength

Aluminum hydroxide 95mg

Magnesium carbonate 358mg

Active ingredient (in each 5mL teaspoonful) Extra Strength

Aluminum hydroxide 254mg

Magnesium carbonate 237.5mg

Purpose

Antacid

Uses

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

Warnings

Do not use if you have kidney disease

Ask a doctor or pharmacist before use if you are
- have kidney disease.
- are on a sodium-restricted diet or a magnesium-restricted diet.
- are taking a prescription drug. Antacids may interact with certain prescription drugs.

When using this product (Regular Strength)
- do not take more than 8 tablespoonfuls in 24 hours
- do not use the maximum dosage for more than 2 weeks
- laxative effect may occur

When using this product (Extra Strength)
- do not take more than 16 teaspoonfuls in 24 hours

- do not use the maximum dosage for more than 2 weeks except under the advice and supervision of a doctor
- laxative effect may occur

Keep out of reach of children.

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

Directions (Regular Strength)

- shake well
- take 1-2 tablespoonfuls four times a day or as directed by a doctor
- take after meals and at bedtime
- dispense product only by spoon or other measuring device

Directions (Extra Strength)

- shake well
- take 2-4 teaspoonfuls four times a day or as directed by a doctor
- take after meals and at bedtime
- dispense product only by spoon or other measuring device

Other information (Regular Strength)

- **each tablespoon (15mL) contains:**magnesium 115mg, sodium 52mg
- store at up to 25°C (77°F); avoid freezing
- keep tightly closed

Other information (Extra Strength)

- **each teaspoon (5mL) contains:**magnesium 80mg, sodium 14mg
- store at up to 25°C (77°F); avoid freezing
- keep tightly closed

Inactive ingredients (Regular Strength)

benzyl alcohol, D&C yellow #10, edetate disodium, FD&C blue #1, flavor, glycerin, saccharin sodium, sodium alginate, sorbitol solution, water, xanthan gum

Inactive ingredients (Extra Strength Cool Mint)

benzyl alcohol, edetate disodium, flavor, glycerin, saccharin sodium, simethicone emulsion, sodium alginate, sorbitol solution, water, xanthan gum

Inactive Ingredients (Extra Strength Cherry)

Benzyl alcohol, edentate disodium, flavor, glycerin, saccharin sodium, simethicone emulsion, sodium alginate, sorbitol solution, water, xanthan gum

Questions or comments?

call toll-free (English/Spanish) **1-888-367-6471**weekdays

Distributed by:

GlaxoSmithKlineConsumer Healthcare, L.P.

Moon Twp, PA 15108, Made in the U.S.A

IMPORTANT:

Do not use if foil inner seal imprinted "SEALED FOR YOUR PROTECTION" is disturbed or missing.

Principal Display Panel

NDC 0135-0094-41

Gaviscon ®

REGULAR STRENGTH

LIQUID ANTACID

- *Fast-Acting Heartburn Relief*
- *Helps Keep Acid Down for Hours*

COOL MINT

FLAVOR

12*fl oz (355 ml)*

FRONT: 100631XB

BACK: 100632XA

Principal Display Panel

NDC 0135-0095-41

Gaviscon **®**

EXTRA STRENGTH

LIQUID ANTACID

- *Fast-Acting Heartburn Relief*
- *Helps Keep Acid Down for Hours*

COOL MINT

FLAVOR

12*fl oz (355 ml)*

FRONT: 100651XB

BACK: 100652XA

Principal Display Panel

NDC 0135-0574-01

Gaviscon **®**

EXTRA STRENGTH

LIQUID ANTACID

- *Fast-Acting Heartburn Relief*
- *Helps Keep Acid Down for Hours*

CHERRY

FLAVOR

12*fl oz (355 ml)*

FRONT: 103698XA

BACK: 103699XA

GAVISCON

aluminum hydroxide and magnesium carbonate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0135-0094
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	95 mg in 15 mL
MAGNESIUM CARBONATE (UNII: 0E53J927NA) (CARBONATE ION - UNII:7UJQ5OPE7D)	MAGNESIUM CARBONATE	358 mg in 15 mL

Inactive Ingredients	
Ingredient Name	**Strength**

BENZYL ALCOHOL (UNII: LKG8494WBH)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCERIN (UNII: PDC6A3C0OX)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	green	**Score**	
Shape		**Size**	
Flavor	MINT (cool mint)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0094-41	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2011	
2	NDC:0135-0094-42	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	01/14/2011	

GAVISCON

aluminum hydroxide and magnesium carbonate liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0135-0095
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	254 mg in 5 mL
MAGNESIUM CARBONATE (UNII: 0E53J927NA) (CARBONATE ION -	MAGNESIUM	237.5 mg

UNII:7UJQ5OPE7D)	CARBONATE	in 5 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	green	**Score**	
Shape		**Size**	
Flavor	MINT (cool mint)	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0135-0095-41	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	01/14/2011	

GAVISCON

aluminum hydroxide and magnesium carbonate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0135-0574
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of**	**Strength**

Ingredient Name	Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	254 mg in 5 mL
MAGNESIUM CARBONATE (UNII: 0E53J927NA) (CARBONATE ION - UNII:7UJQ5OPE7D)	MAGNESIUM CARBONATE	237.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor	CHERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0135-0574-01	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	08/01/2014	

Labeler - GlaxoSmithKline Consumer Healthcare Holdings (US) LLC (079944263)

Revised: 11/2023 GlaxoSmithKline Consumer Healthcare Holdings (US) LLC