

MAXIMUM STRENGTH ADVANCED ANTACID- aluminum hydroxide, magnesium hydroxide, simethicone liquid
DOLGENCORP, LLC

dg ant max cherry

Active ingredients (in each 10 mL dose)

Aluminum hydroxide (equivalent to dried gel, USP) 800 mg

Magnesium hydroxide 800 mg

Simethicone 80 mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- symptoms referred to as gas

Warnings

Ask a doctor before use if you have

- kidney disease
- a magnesium-restricted diet

Ask a doctor or pharmacist before use if you are presently 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.

Directions

- shake well before use
- adults and children 12 years of age and older: take 10ml to 20ml (1 to 2 doses) two

times a day or as directed by a doctor

- do not exceed 40ml (4 doses) in a 24 hour period or use the maximum dosage for more than 2 weeks
- children under 12 years of age: ask a doctor

Other information

- **each 10 mL dose contains:** magnesium 340 mg, sodium 10 mg
- do not freeze
- store at room temperature tightly closed

Inactive ingredients

benzyl alcohol, butylparaben, carboxymethylcellulose sodium, flavor, hypromellose, microcrystalline cellulose, propylparaben, purified water, saccharin sodium, sorbitol solution

Questions or Comments?

1-888-309-9030

package label

Since 1903
Rexall[®]

MAXIMUM STRENGTH

**Antacid
Advanced**

Antacid & Antigas

Fast relief of:

- Heartburn
- Acid Indigestion
- Pressure
- Bloating (Gas)

**Cherry
FLAVORED LIQUID**

**12 FL OZ
(355 mL)**

207-11322-0 REV0221

Drug Facts

TAMPER-EVIDENT: Do not use if the breakaway band on cap is broken or missing.

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907-11322-0 REV GC0221



MAXIMUM STRENGTH ADVANCED ANTACID

aluminum hydroxide, magnesium hydroxide, simethicone liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-638
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	800 mg in 10 mL

MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	800 mg in 10 mL
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	80 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0K00R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-638-12	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	02/01/2021	

Labeler - DOLGENCORP, LLC (068331990)

Registrant - GCP Laboratories (965480861)

Establishment

Name	Address	ID/FEI	Business Operations
GCP Laboratories		965480861	manufacture(55910-638)