GERI-LANTA MAXIMUM STRENGTH- aluminum hydroxide, magnesium hydroxide, dimethicone suspension
NuCare Pharmaceuticals, Inc.

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gerilanta max 619

Active ingredients (in each 5 mL teaspoonful)

Aluminum hydroxide 400mg (equivalent to dried gel, USP)
Magnesium hydroxide 400 mg
Simethicone 40mg

Purposes

Antacid

Antigas

Uses

relieves

- heartburn
- sour stomach
- acid indigestion
- the 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

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 and older: take 2 to 4 teaspoonfuls between meals, at bedtime, or as directed by a doctor
- do not exceed 12 teaspoonfuls in a 24 hour period or use the maximum dosage for more than 2 weeks
- children under 12 years: ask a doctor

Other information

- **each 5 mL teaspoonful contains:**magnesium 165 mg, sodium 5 mg
- do not freeze
- store at room temperature tightly closed

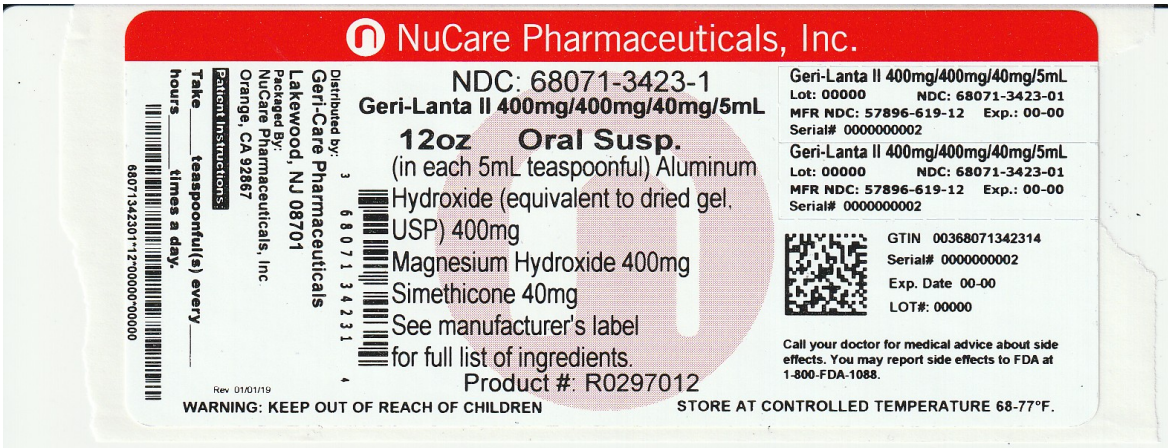
Inactive ingredients

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

Questions or comments?

1-800-540-3765

package Label



GERI-LANTA MAXIMUM STRENGTH

aluminum hydroxide, magnesium hydroxide, dimethicone suspension

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68071-3423(NDC:57896-619)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	ALUMINUM HYDROXIDE	400 mg in 5 mL
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)	MAGNESIUM HYDROXIDE	400 mg in 5 mL
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	40 mg in 5 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: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	LEMON (Lemon)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-3423-1	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/11/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	01/01/2000	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-3423)

Revised: 12/2024 NuCare Pharmaceuticals, Inc.