

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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

gerilanta max

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

NuCare Pharmaceuticals, Inc.

NDC: 68071-3423-1
Geri-Lanta II 400mg/400mg/40mg/5mL

12oz Oral Susp.
(in each 5mL teaspoonful) Aluminum Hydroxide (equivalent to dried gel, USP) 400mg
Magnesium Hydroxide 400mg
Simethicone 40mg
See manufacturer's label for full list of ingredients.

Product #: R0297012

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

Geriatric Information:
Geri-Lanta II 400mg/400mg/40mg/5mL
Lot: 00000 NDC: 68071-3423-01
MFR NDC: 57896-619-12 Exp.: 00-00
Serial# 0000000002

Geriatric Information:
Geri-Lanta II 400mg/400mg/40mg/5mL
Lot: 00000 NDC: 68071-3423-01
MFR NDC: 57896-619-12 Exp.: 00-00
Serial# 0000000002

GTIN 00368071342314
Serial# 0000000002
Exp. Date 00-00
LOT#: 00000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Patent Instructions:
Take _____ teaspoonful(s) every _____ hours _____ times a day.

Rev 01/01/19

Distributed by: 3 68071 34231 4
Geriatric Pharmaceuticals
Brooklyn, NY 11204
Packaged By:
NuCare Pharmaceuticals, Inc.
Orange, CA 92867

6807134230112-000000-00000

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 | 06/01/2023 | |

| Marketing Information | | | |
|-----------------------|------------------------------------------|----------------------|--------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph final | part331 | 01/01/2000 | |

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

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

Revised: 6/2023

NuCare Pharmaceuticals, Inc.