

**GERI-LANTA ANTACID ANTIGAS- aluminum hydroxide, magnesium hydroxide, dimethicone suspension**  
**Preferred 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.*

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**Geri-lanta original 629**

**Active ingredients (in each 5 mL teaspoonful)**

Aluminum hydroxide 200 mg (equivalent to dried gel, USP)  
Magnesium hydroxide 200 mg  
Simethicone 20mg

**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** 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

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68788-8169(NDC:57896-629)
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Route of Administration		ORAL		
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)		ALUMINUM HYDROXIDE	200 mg in 5 mL	
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S) (MAGNESIUM CATION - UNII:T6V3LHY838, HYDROXIDE ION - UNII:9159UV381P)		MAGNESIUM HYDROXIDE	200 mg in 5 mL	
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)		DIMETHICONE	20 mg in 5 mL	
Inactive Ingredients				
Ingredient Name			Strength	
BENZYL ALCOHOL (UNII: LKG8494WBH)				
BUTYLPARABEN (UNII: 3QPI1U3FV8)				
HYDROXYETHYL CELLULOSE (4000 MPA.S AT 1%) (UNII: ZYD53NBL45)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
WATER (UNII: 059QF0KO0R)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SORBITOL SOLUTION (UNII: 8KW3E207O2)				
Product Characteristics				
Color		Score		
Shape		Size		
Flavor	LEMON (citrus mint)	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8169-3	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/11/2022	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph final	part331		07/11/2022	

**Labeler** - Preferred Pharmaceuticals Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals Inc. (791119022)

Establishment			
Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8169)

Revised: 7/2023

Preferred Pharmaceuticals Inc.