

# **HAND SANITIZER GEL- hand sanitizer gel gel LS Promotions Inc.**

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## **Hand Sanitizer Gel**

### **Drug Facts**

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### **Active Ingredients**

Ethyl Alcohol 62%

### **Purpose**

Antiseptic

### **Uses**

- To help decrease bacteria on hands.

### **Warnings**

- For external use only
- Flammable, keep away from fire or flame

### **When using this product**

- Keep out of eyes
- In case of contact with eyes, flush thoroughly with water and then seek medical advise

### **Stop use and ask a doctor**

- If irritation or redness develops

### **Keep out of reach of children**

- If swallowed, get medical help or contact a poison control center immediately

### **Directions**

- Place enough product in your palm to thoroughly cover hand.
- Rub hands together briskly until dry.
- For children under 6 years old, use adult supervision.

### **Other information**

- Do not store above 100°F

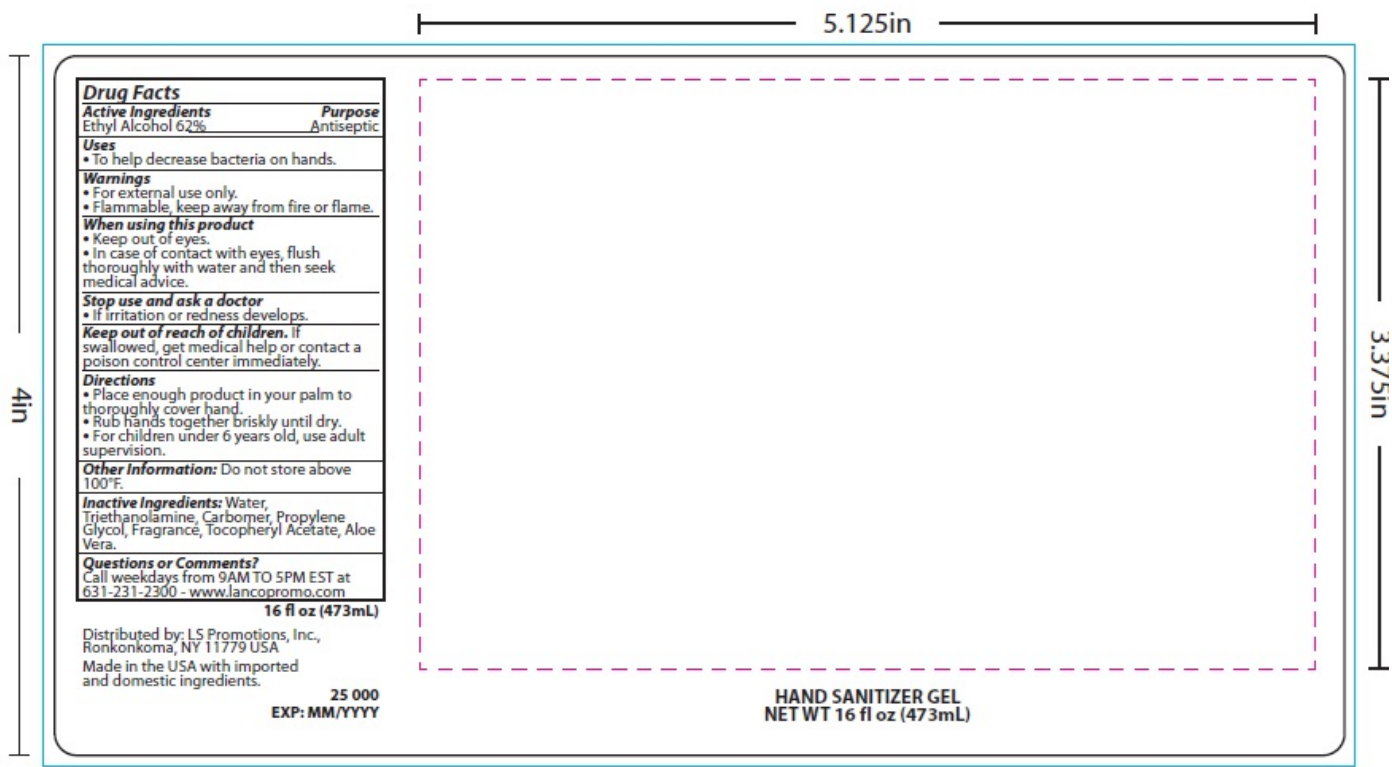
Inactive Ingredients

Inactive ingredients: water, triethanolamine, carbomer, propylene glycol, fragrance, tocopherol acetate, aloe vera oil.

Questions or Comments?

Call weekdays from 9 AM to 5 PM EST at 631-231-2300

Principal display panel



ABL1211 Hand Sanitizer Gel 16 NDC 72449-103-02

HAND SANITIZER GEL			
hand sanitizer gel gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72449-103
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	0.62 mL in 1 mL
Inactive Ingredients				
Ingredient Name				Strength
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)				
TRIETHANOLAMINE (UNII: 9O3K93S3TK)				
FRAGRANCE GREEN APPLE ORC1902709 (UNII: X2H4PXN0HA)				
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
CARBOMER 1342 (UNII: 809Y72KV36)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72449-103-06	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
2	NDC:72449-103-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
3	NDC:72449-103-02	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/04/2025	
4	NDC:72449-103-05	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
5	NDC:72449-103-01	2.9 mL in 1 PACKET; Type 0: Not a Combination Product	04/04/2025	
6	NDC:72449-103-10	29 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
7	NDC:72449-103-11	295 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
8	NDC:72449-103-09	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
9	NDC:72449-103-08	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/04/2025	
10	NDC:72449-103-07	295 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/04/2025	
11	NDC:72449-103-03	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/04/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		505G(a)(3)	03/23/2020	

**Labeler** - LS Promotions Inc. (024818391)

**Registrant** - LS Promotions Inc. (024818391)**Establishment**

Name	Address	ID/FEI	Business Operations
LS Promotions Inc.		024818391	manufacture(72449-103)

Revised: 4/2025

LS Promotions Inc.