HAND SANITIZER- hand sanitizer gel LS Promotions 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.

Hand Sanitizer Gel

Drug Facts

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Active Ingredients

Ethyl Alcohol

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

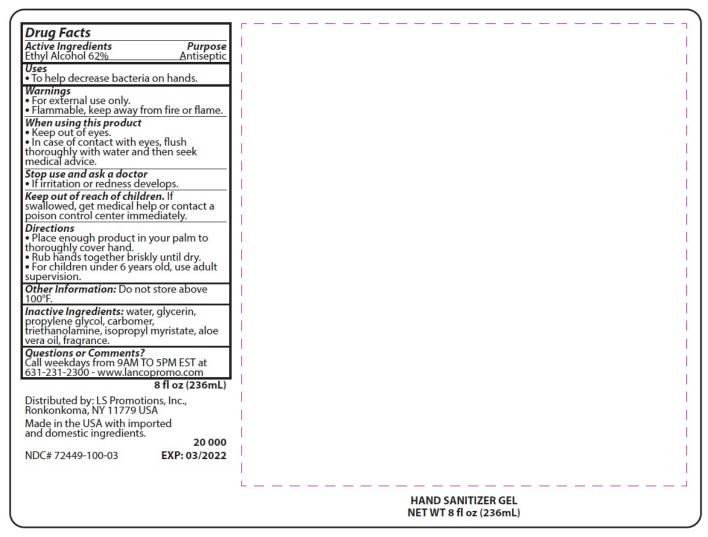
Inactive Ingredients

Inactive ingredients: water, glycerin, propylene glycol, carbomer, triethanolamine, isopropyl myristate, aloe vera oil, fragrance

Questions or Comments?

Call weekdays from 9 AM to 5 PM EST at 631-231-2300 - www.lancopromo.com

Principal display panel



ABL1900 Hand Sanitizer Gel NDC # 72449-100-03

HAND SANITIZER					
hand sanitizer gel					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:72449-100	
Route of Administration	TOPICAL				
Active Ingredient/Active Moiety					
Ingredient Name			Basis of Strengt	h Strength	1
ALCOHOL (UNII: 3K9958V90M) (ALC	OHOL - UNII:3K9958V90M)		ALCOHOL	0.62 mL in 1 m	L

Inactive Ingredients				
Ingredient Name	Strength			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
CARBOMER 1342 (UNII: 809 Y72KV36)				
TROLAMINE (UNII: 9O3K93S3TK)				
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)				
WATER (UNII: 059QF0KO0R)				
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)				
GLYCERIN (UNII: PDC6 A3C0 OX)				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72449-100- 03	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
2	NDC:72449-100- 04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
3	NDC:72449-100- 05	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
4	NDC:72449-100- 06	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
5	NDC:72449-100- 01	2.9 mL in 1 PACKET; Type 0: Not a Combination Product	03/23/2020	
6	NDC:72449-100- 02	5.9 mL in 1 PACKET; Type 0: Not a Combination Product	03/23/2020	
7	NDC:72449-100- 07	29 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
8	NDC:72449-100- 08	295 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
9	NDC:72449-100- 09	354 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/23/2020	
10	NDC:72449-100- 10	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/23/2020	
11	NDC:72449-100- 11	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/01/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	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(72449-100)	
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Revised: 12/2020 LS Promotions Inc.