HAND SANITIZER- alcohol gel Jon Davler, 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.

LASplash So Fresh Lemon Gel Hand Sanitizer 74044-1008-4 100ml

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Ethyl Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water (aqua), Isopropyl Alcohol, Propylene Glycol, Glycerin, Peg-6 (and) AMP-Acrylates/Vinyl Isodecanoate Crosspolymer, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate, Citus Medica Limonum (Lemon) Peel Oil.

Package Label - Principal Display Panel

100 mL NDC:74044-1008-4



HAND SANITIZER

alcohol gel

Product Information

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:74044-1008 |
|-------------------------|----------------|--------------------|----------------|
| Route of Administration | TOPICAL | | |

| Active Ingredient/Active Moiety | | | |
|--|-------------------|-----------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 70 mL in 100 mL | |

| Inactive Ingredients | | | | |
|---|--------------------------|--|--|--|
| Ingredient Name | Strength | | | |
| ALOE (UNII: V5VD430 YW9) | $0.094\;mL$ in $100\;mL$ | | | |
| POLYETHYLENE GLYCOL 300 (UNII: 5655G9 Y8 AQ) | 2.821 mL in 100 mL | | | |
| CITRUS MEDICA FRUIT (UNII: ZE5Q6PN9ON) | 0.12 mL in 100 mL | | | |
| ISOPROPYL ALCOHOL (UNII: ND2M416302) | 6.996 mL in 100 mL | | | |
| .ALPHATO COPHEROL ACETATE, DL- (UNII: WR1WPI7EW8) | 0.094 mL in 100 mL | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | 0.451 mL in 100 mL | | | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | $0.677\;mL$ in $100\;mL$ | | | |
| WATER (UNII: 059QF0KO0R) | | | | |

| ı | Packaging | | | |
|---|------------------------|--|------------|-----------------------|
| | # Item Code | Item Code Package Description | | Marketing End Date |
| | 1 NDC:74044- 1008-4 | 100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 06/01/2020 | |

| Marketing Information | | | |
|-------------------------|--|----------------------|--------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | part333A | 06/01/2020 | |
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Labeler - Jon Davler, Inc. (097710185)

| Establishment | | | |
|------------------|---------|-----------|-------------------------|
| Name | Address | ID/FEI | Business Operations |
| Jon Davler, Inc. | | 097710185 | manufacture(74044-1008) |

Revised: 6/2020 Jon Davler, Inc.