HAND SANITIZER- alcohol liquid LD ACEPRO

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

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)

Alcohol 80% 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

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel



HAND SANITIZER

Alcohol Antiseptic 80% Topical Solution Non-Sterile Solution

Drug Facts

Use(s)

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

Warnings

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1 GAL (3.78 L)

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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 150-300C (59-86F)
- Avoid freezing and excessive heat above 400C (104F)

Inactive ingredients: glycerin, hydrogen peroxide, purified water USP



LD ACEPRO INC.
South Heckensack, NJ 07606.
Made in USA

Lot 2020G1

3785 mL NDC: 79888-378-01

235mL NDC 79888-378-02



HAND SANITIZER

Alcohol Antiseptic 80%
Topical Solution
Non-Sterile Solution

8 fl.oz. (230 mL)

Drug Facts

Use(s)

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

Warnings

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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.

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LD ACEPRO INC. South Hackensack, NJ 07606. Mede in USA

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HAND SANITIZER

alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79888-378

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M)

ALCOHOL

Basis of Strength

ALCOHOL

80 mL in 100 mL

Inactive Ingredients

Ingredient Name Strength

GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

Pā	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:79888-378- 01	3785 mL in 1 JUG; Type 0: Not a Combination Product	03/30/2020			
_	NDC:79888-378- 02	235 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/30/2020		

Labeler - LD ACEPRO (091161772)

Revised: 9/2022 LD ACEPRO