

HAND SANITIZER - LAVENDER- alcohol liquid

Farmers & Distillers

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.

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.

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (73%, 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. Lavender Oil (8% v/v)
- e. Sterile distilled water or boiled cold water.

Active Ingredient(s)

Alcohol 73% 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, lavender oil, purified water USP

Package Label - Principal Display Panel

208198 L NDC: 74183-002-04

Drug Facts	
Active Ingredient(s) Alcohol 73% v/v	Purpose Antiseptic
Use(s) Health care personnel hand rub to help reduce bacteria that potentially can cause disease.	
Warnings For external use only. Flammable. Keep away from heat or flame.	
Do not use <ul style="list-style-type: none">• on children less than 2 months of age• on open skin wound	
When using this product keep out of eyes, ears and mouth. In case of contact with eyes rinse 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.	
Directions <ul style="list-style-type: none">• 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 <ul style="list-style-type: none">• Store between 15-30°C (59-86°)• Avoid freezing and excessive heat (above 40°)	
Inactive Ingredients Glycerin, hydrogen peroxide, purified water, lavender oil	

FOUNDING SPIRITS

HAND

SANITIZER

LAVENDER

Alcohol Antiseptic 73%, Topical Solution
Non-sterile Solution
55 gal

NOT FOR INTERNAL CONSUMPTION

HAND SANITIZER - LAVENDER

alcohol liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:74183-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	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: 059QF0K00R)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	8 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74183-002-04	208198 mL in 1 DRUM; Type 0: Not a Combination Product	09/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/16/2020	

Labeler - Farmers & Distillers (076425254)

Registrant - Farmers & Distillers (076425254)

Establishment

Name	Address	ID/FEI	Business Operations
Farmers & Distillers		076425254	manufacture(74183-002)

Revised: 10/2020

Farmers & Distillers