HAND SANITIZER- alcohol liquid Range & River Distilling, 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

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

3785 mL NDC: 74469-001-1



W.H.O. RECOMMENDED HAND RUB FORMULATION #1

FOR EXTERNAL USE ONLY NOT SAFE FOR HUMAN CONSUMPTION NON-STERILE SOLUTION

> DATE: BATCH: 1 gallon | 3.785 L

The hand sanitizer is manufactured using only to States Pharmacopole (USP) grade ingredients the product (percentage in final product formula World Health Organization (WHO) recommenda Achohol (ethanol) (USP or Food Chemical Co a (80%, volume/volume (w/v)) in an aqueous solution of according to Alcohol and Tobacco Tax and Trada according to Alcohol and Tobacco Tax and Trada (Sycerol (1.45% v/v)).

Alcohol (1.45% v/v)

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Drug Facts Active ingredient Ethyl alcohol 80% v/v.... 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 and rash occurs. These may be signs of a serious medical condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right Directions • Place enough 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 For questions/concerns, email info@singletrackspirits.com or call (307) 761-1296

HAND SANITIZER

Droduct Information

alcohol liquid

ı	r roduct milormation			
l	Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74469-0001

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	
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL	
WATER (UNII: 059QF0KO0R)		

l	Packaging				
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
l	1 NDC:74469-0001-1	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/02/2020		
	2 NDC:74469-0001-2	15140 mL in 1 PAIL; Type 0: Not a Combination Product	04/02/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	04/02/2020		

Labeler - Range & River Distilling, Inc. (117468361)

Establishment				
Name	Address	ID/FEI	Business Operations	
Range & River Distilling, Inc.		117468361	manufacture(74469-0001)	

Revised: 4/2020 Range & River Distilling, Inc.