HAND SANITIZER- is opropyl alcohol liquid Josh Shamoilia Enterprises LLC

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 (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Isopropyl Alcohol (75%, 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)

Isopropyl Alcohol 75% 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

carbomer, lavender essential oil, purified water USP, vitamin e oil, aloe vera leaf, dimethicone

Package Label - Principal Display Panel

DRIPPIES MOISTURIZING HAND SANITIZER Fast Absorbing, Hydrating, Lightly Scented, Hand Sanitizer Mist Sprays

LAVENDE

0.5 FL, OZ, / 15ML







Drug Facts

Active Ingredients

Purpose

Ethyl Alcohol 70%

Antiseptic

- · For hand washing to decrease bacteria on skin
- · Protect and restores moisture
- Recommended for repeated use

Warnings

- Flammable. Keep away from fire or flame.
- · For external use only. Keep out of eyes. In case of contact eyes, rinse with water.
- Stop use and consult a doctor if irritation and redness develope and persist.
- Keep out of reach of children.
- · If swallowed, get medical help promptly.

Directions

Spray 2-3 times on hands and rub together until dry.

Other Information

Store Under 104 F.

Inactive Ingredients

Aloe Barbodensis Leaf, Carbomer, Dimethicone, Water, Vitamin E Oil, Lavender Essential Oil

15 ml NDC: 76826-1216-1

HAND SANITIZER

isopropyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76826-1216

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	75 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
DIMETHICO NE 100 (UNII: RO266O364U)	1 mL in 100 mL		
.ALPHATO COPHERO L ACETATE, D- (UNII: A7E6112E4N)	1 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			
LAVENDER O IL (UNII: ZBP1YXW0 H8)	1 mL in 100 mL		
ALOE VERA LEAF (UNII: ZY81Z83H0X)	1 mL in 100 mL		
CARBOMER 934 (UNII: Z135WT9208)	0.125 mL in 100 mL		

ı	Packag	ing			
	# Iten	n Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:76	6826-1216-	15 mL in 1 BOTTLE, SPRAY; 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 - Josh Shamoilia Enterprises LLC (121195511)

Establishment				
Name	Address	ID/FEI	Business Operations	
Josh Shamoilia Enterprises LLCN		121195511	manufacture(76826-1216)	

Revised: 5/2020 Josh Shamoilia Enterprises LLC