HAND SANITIZER- alcohol liquid Givaudan Fragrances Corporation

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

230 mL NDC: 74672-139-01

Alcohol Antiseptic 80% Topical Solution

Antiseptic Hand Rub Non-Sterile Solution

230 mL

Givaudan Flavors Corporation

Production Date: Batch Number:

Drug Facts	
Active ingredient[s] Alcohol 80% v/v	Purpose Antiseptic
Use[s] Health care personnel hand rub to help reduce bacteria that potentially can cause dis	sease.
Warnings	STITLE OF
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Other information	IRI

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Inactive ingredients glycerin, hydrogen peroxide, purified water USP

Avoid freezing and excessive heat above 40C (104F)

250 mL NDC: 74672-139-02

Store between 15-30C (59-86F)

Alcohol Antiseptic 80% Topical Solution

Antiseptic Hand Rub Non-Sterile Solution

250 mL

Givaudan Flavors Corporation

Production Date: Batch Number:

Drug Facts	
Active ingredient[s]	Purpose
Alcohol 20% v/v	Anticontic

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

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- · 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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Inactive ingredients glycerin, hydrogen peroxide, purified water USP



HAND SANITIZER

alcohol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74672-139	
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: PDC6 A3C0 OX)	1.45 mL in 100 mL			
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL			
WATER (UNII: 059QF0KO0R)				

Pa	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:74672-139-	230 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	

2 NDC:74672-139- 02	250 mL in 1 BOTTLE, GLASS; Type 0: Not a Co Product	mbination 03/30/2020		
Marketing Information				
Marketing Categ	ory Application Number or Monograph	Citation Marketing Start Date	Marketing End Date	
OTC monograph not	final part333A	03/30/2020		

Labeler - Givaudan Fragrances Corporation (008645475)

Registrant - Givaudan Fragrances Corporation (008645475)

Establishment				
Name	Address	ID/FEI	Business Operations	
Givaudan Fragrances Corporation		008645475	manufacture(74672-139)	

Revised: 4/2020 Givaudan Fragrances Corporation