HAND SANITIZER- is opropyl alcohol liquid Verico Technology

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. 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

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

Core ID: 3"



SPECIALTY PRINTING CONVERTING BETTER IDEAS	- THIS PROOF MAY NOT DISPLAY OR PRODUCE ACCURATE COLOR MATCH - Please review the design, copy and ink breakdown carefully, if any corrections are required, please make them on this sheet. Orders	Wind 3
4 Thompson Rd. * East Windsor, CT 06086 * Mt 860.623.8870 * F; 860.623.8861	will not proceed without written approval. Your signature gives us permission to proceed with this design, tooling, and order.	
Customer: Verico Technology Date: 04/07/20 Job ID: VTECH006 Rev:	Front: Black Blue 2388 6	
Label Size: 4.3125 W x 6.25 H CR: .125	Back:	
Approval: Date:		

Core ID: 3"



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Customer: Verico Technology Date: 04/07/20 Job ID: VTECH005 Rev:	Frunt: Black Blue	
Label Size: 5.125 W x 6.75 H CR: .125	Back:	
Approval: Date:		

FORM# QC007 ISSUE A8

*** THIS IS A CONTROLLED DOCUMENT ***

REV. DATE: 10/17/17

354 ml NDC: 75647-001-02 236 ml NDC: 75647-001-04 1000 ml NDC: 75647-001-03

HAND SANITIZER

isopropyl alcohol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75647-001	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
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		
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			

Product Characteristics			
Color	Score		
Shape	Size		
Flavor	Imprint Code		
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:75647-001-04	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020		
2	NDC:75647-001-02	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020		
3	NDC:75647-001-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020		
4	NDC:75647-001-05	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/11/2020		

32oz Sanitizer Label Labels per roll: 1,000 Rolls per case: 2 Core ID: 3"

WIND DIRECTION —



SPECIALTY PRINTING - THIS PROOF MAY NOT DISPLAY OR PRODUCE ACCURATE COLOR MATCH -Wind 3 CONVERTING BETTER IDEAS Please review the design, copy and ink breakdown carefully. If any corrections are required, please make them on this sheet. Ord will not proceed without written approval. Your signature gives us permission to proceed with this design, tooling, and order. 4 Thompson Rd. • East Windsor, CT 06088 • M: 860.623.8870 • F: 860.623.8861 Customer: Verico Technology Date: 04/29/20 Job ID: VTECH009 Rev: Label Size: 5.5 W x 8.0 H CR: Back: Approval: Date: FORM# QC007 ISSUE A8 *** THIS IS A CONTROLLED DOCUMENT ***

Marketing InformationMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateOTC monograph not finalpart333A04/20/2020

Labeler - Verico Technology (183615426)

Registrant - Verico Technology (183615426)

Establishment			
Name	Address	ID/FEI	Business Operations

Verico Technology	183615426	manufacture(75647-001)
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Revised: 5/2020 Verico Technology