

## **HAND SANITIZER- 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.*

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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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away.

**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)

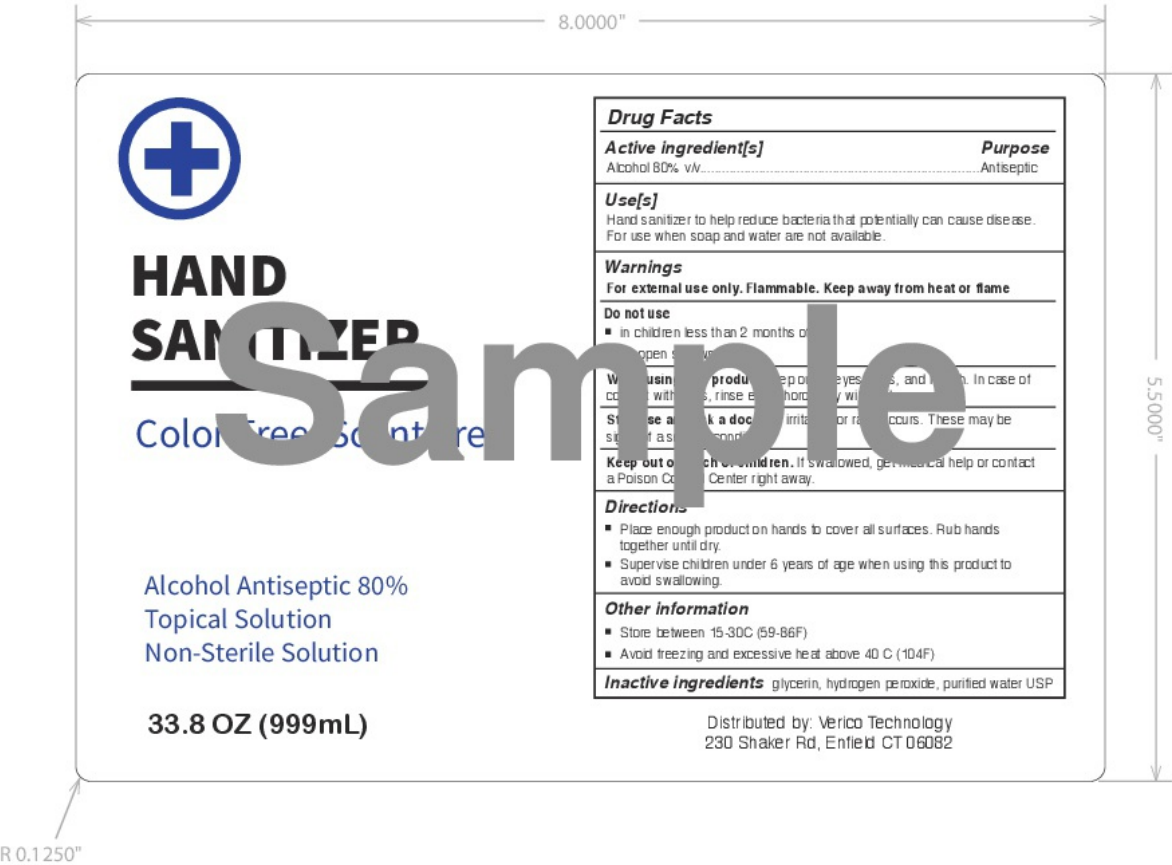
**Inactive ingredients**





glycerin, hydrogen peroxide, purified water USP

**1000 mL NDC: 75647-003-03**

33.8oz Sanitizer Label  
Labels per roll: 1,000  
Rolls per case: 2  
Core ID: 3"

WIND DIRECTION →



 <b>SPECIALTY PRINTING</b> CONVERTING BETTER IDEAS 4 Thompson Rd. • East Windsor, CT 06028 • M: 860.623.8870 • F: 860.623.8861	<b>- THIS PROOF MAY NOT DISPLAY OR PRODUCE ACCURATE COLOR MATCH -</b> Please review the design, copy and ink breakdown carefully. If any corrections are required, please make them on this sheet. Orders will not proceed without written approval. Your signature gives us permission to proceed with this design, tooling, and order.	<b>Wind 3</b> 
Customer: <b>Verico Technology</b> Date: <b>04/29/20</b> Job ID: <b>VTECH010</b> Rev: _____ Label Size: <b>5.5</b> W x <b>8.0</b> H    CR: <b>.125</b> Approval: _____      Date: _____	Front:  <b>Black</b> <b>Blue</b> Back:  <b>Black</b> <b>Blue</b>	

FORM Q0007 ISSUE A8      \*\*\* THIS IS A CONTROLLED DOCUMENT \*\*\*      REV DATE: 10/17/17

HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75647-002
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)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75647-002-03	1000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/06/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/06/2020	

**Labeler** - Verico Technology (183615426)

## Establishment

Name	Address	ID/FEI	Business Operations
Verico Technology		183615426	manufacture(75647-002)

Revised: 8/2020

Verico Technology