HAND SANITIZER- alcohol liquid The American Bottling Company

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 1 gallon

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

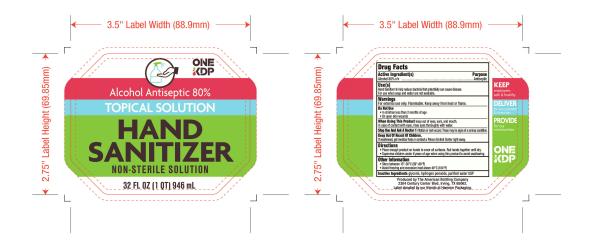
Package Label - Principle Display Panel

3.78 L NDC: 74362-001-01

7.0" Label Width (177.8mm)



946 mL NDC: 74362-001-02



alcohol liquid						
Product Informa	ation					
Product T ype		HUMAN OTC DRUG	Item Code (Source)		NDC:74362-001	
Route of Administr	ation	TOPICAL				
Active Ingredie	nt/Active Moie	ety				
Ingredient Name			Basis of Strength		Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL		80 mL in 100 mL	
				necone		
Inactive Ingredi	ents	redient Name				Strength
	ents Ing				1.45 mL in 10	Strength
GLYCERIN (UNII: PD	ents Ing C6A3C0OX)	redient Name				Strength 0 mL
Inactive Ingredi GLYCERIN (UNII: PD HYDROGEN PEROX WATER (UNII: 059Q)	ents Ing C6A3C0OX) IDE (UNII: BBX06)	redient Name			1.45 mL in 10	Strength 0 mL
GLYCERIN (UNII: PD HYDROGEN PEROX	ents Ing C6A3C0OX) IDE (UNII: BBX06)	redient Name			1.45 mL in 10	Strength 0 mL
GLYCERIN (UNII: PD HYDROGEN PEROX WATER (UNII: 059Q)	ents Ing C6A3C0OX) IDE (UNII: BBX06)	redient Name			1.45 mL in 10	Strength 0 mL
GLYCERIN (UNII: PD HYDROGEN PEROX	ents Ing C6A3C0OX) IDE (UNII: BBX064 F0KO0R)	redient Name		Marke	1.45 mL in 10	Strength 0 mL

Marketing Information							
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
part333A	03/30/2020						
	Application Number or Monograph Citation	Application Number or Monograph Citation Marketing Start Date					

Labeler - The American Bottling Company (095218129)

Revised: 4/2020

The American Bottling Company