HAND SANITIZER- hand sanitizer liquid Avantek Capital 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. 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)

Denatured Ethyl Alcohol 80% v/v

Purpose

Antiseptic

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

• Do not use on 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.

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.

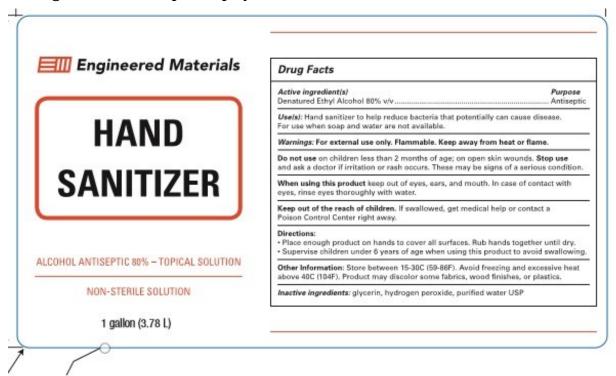
Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)
- Product may discolor some fabrics, wood finishes, or plastics.

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel



Emgineered Materials

HAND SANITIZER

ALCOHOL ANTISEPTIC 80% - TOPICAL SOLUTION

NON-STERILE SOLUTION

10 FL OZ (296 mL)

Drug Facts

Active ingredient(s)
Denatured Ethyl Alcohol 80% v/v............

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HAND SANITIZER

ALCOHOL ANTISEPTIC 80% - TOPICAL SOLUTION

NON-STERILE SOLUTION

8 FL OZ (237 mL)

Drug Facts

Active ingredient(s)
Denatured Ethyl Alcohol 80% v/v.

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Warnings: For external use only. Flammable. Keep away from heat or flame

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Other Information: Store between 15-30C (59-86F). Avoid freezing and excessive heat above 40C (104F). Product may discolor some fabrics, wood finishes, or plastics.

Inactive ingredients: glycerin, hydrogen peroxide, purified water USP

HAND SANITIZER

hand sanitizer liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:74261-0001

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 189.6 mL in 237 mL

Inactive Ingredients

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Ingredient Name	Strength		
GLYCEROL FORMAL (UNII: 3L7GR2604E)	3.44 mL in 237 mL		
WATER (UNII: 059QF0KO0R)	43.66 mL in 237 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.3 mL in 237 mL		

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74261- 0001-1	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74261- 0001-2	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	
3	NDC:74261- 0001-3	3780 mL in 1 BOTTLE, PLASTIC; 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 - Avantek Capital LLC (087020850)

Establishment					
Name	Address	ID/FEI	Business Operations		
Avantek Capital LLC		087020850	manufacture(74261-0001)		

Revised: 4/2020 Avantek Capital LLC