HAND SANITIZER- alcohol liquid University of Alaska System

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)

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

236 mL NDC:76817-001-01

Drug facts	
0 11	Purpose Antiseptic
Use[s] 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 • on children less than 2 months of age • on open skin wounds	
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Glycerin, hydrogen peroxide, purified water USP

Produced by UAF Environmental Health, Safety, and Risk Management. For more information, contact the EHSRM office at 907-474-5413 or uaf-ehsrm@alaska.edu. www.uaf.edu/safety/



Alcohol antiseptic 80% topical solution

Hand sanitizer Nonsterile solution

8 oz / 237 mL

472 mL NDC: 76817-001-02

Drug Facts	
Active ingredient[s] Purp Alcohol 80% v/v Antise	

Use[s] Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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

Do not use

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Directions

Drug Facts

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 15-30°C (59-86°F)

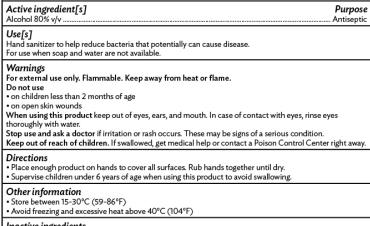
Avoid freezing and excessive heat above 40°C (104°F)

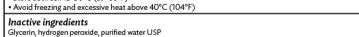
Inactive ingredients

Glycerin, hydrogen peroxide, purified water USP

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3785 mL NDC: 76817-001-03





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Alcohol antiseptic 80% topical solution

Hand sanitizer Nonsterile solution

16 oz / 473 mL



Alcohol antiseptic 80% topical solution

Hand sanitizer Nonsterile solution

1 gallon / 3785 mL

HAND SANITIZER

alcohol liquid

Product Information HUMAN OTC DRUG Product Type Item Code (Source) NDC:76817-001 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		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:76817-001- 01	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/30/2020		
2	NDC:76817-001- 02	472 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/30/2020		
3	NDC:76817-001- 03	3780 mL in 1 BOTTLE; 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 - University of Alaska System (048679567)

Registrant - University of Alaska System (048679567)

Establishment					
Name	Address	ID/FEI	Business Operations		
University of Alaska System		048679567	manufacture (76817-001)		

Revised: 4/2020 University of Alaska System