HAND SANITIZER- alcohol solution Cape Spirits Inc

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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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.

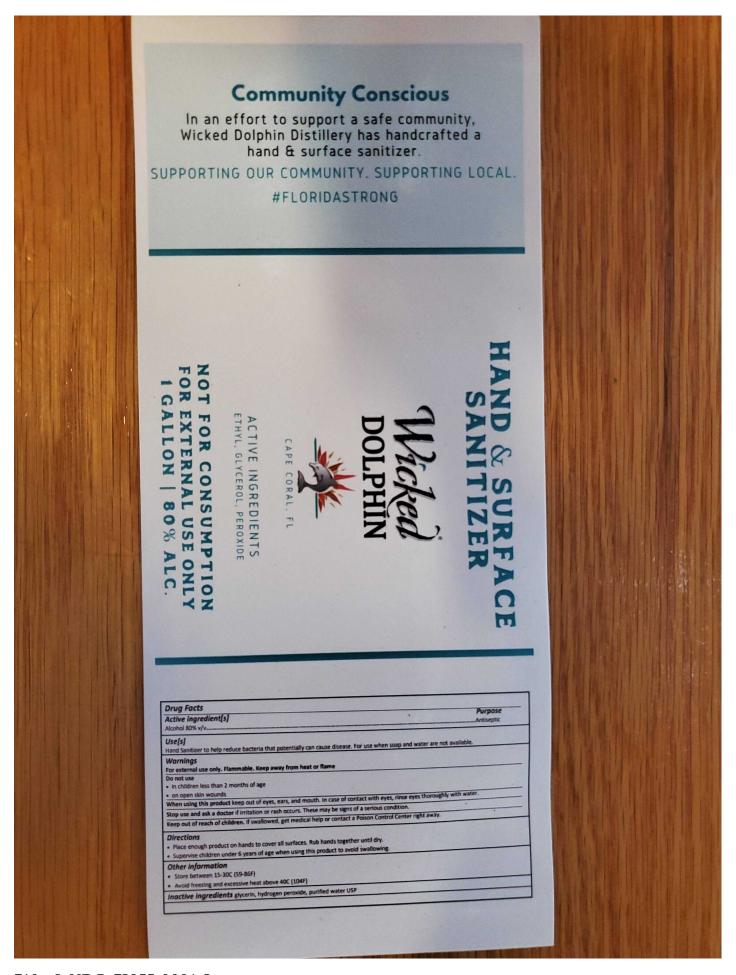
Other information

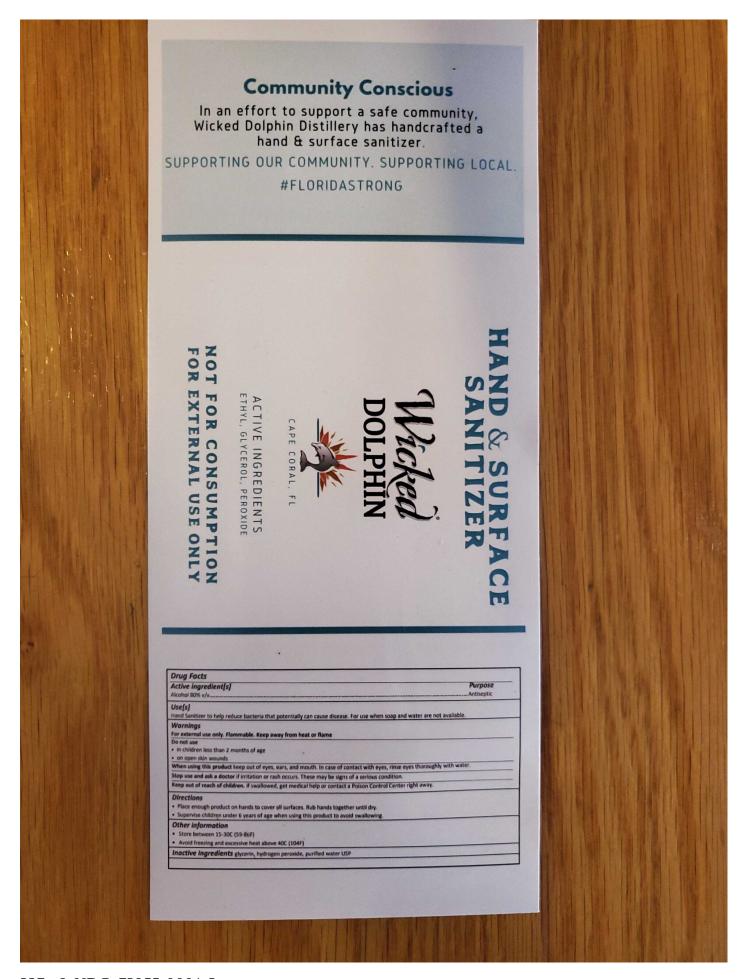
- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel







FOR EXTERNAL USE ONLY 00 NOT FOR CONSUMPTION 02 80% ALC

ETHYL, CLYCEROL, PEROXIDE ACTIVE INCREDIENTS WORLD HEALTH ORGANIZATION FORMULA

CAPE CORAL, FLORIDA

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40C (104F)

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■ Store between 15-30C (59-86F) Other information

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reduce bacteria that potentially can cause disease Use[s] Health care personnel hand rub to help

Active ingredient[s] Alcohol 80% v/v..... . Antiseptic ...Purpose

Drug Facts

Florida's Cruft Run

WORLD HEALTH ORGANIZATION FORMULA

MICKEDDOLPHIN COM

ETHYL, CLYCEROL, PEROXIDE

ACTIVE INCREDIENTS

FOR EXTERNAL USE ONLY NOT FOR CONSUMPTION

80% ALC

CAPE CORAL, FLORIDA

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Florida's Craft Rum

WORLD HEALTH ORGANIZATION FORMULA

ACTIVE INGREDIENTS
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WICKEDDOLPHIN COM

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FOR EXTERNAL USE ONLY NOT FOR CONSUMPTION

CAPE CORAL, FLORIDA

HANDCRAFTED IN

Drug Facts

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HAND SANITIZER

alcohol solution

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Product	Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:73955-0001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ı	Ingredient Name	Basis of Strength	Strength
ı	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)	

Packaging

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#	Item Code Package Description		Marketing Start Date	Marketing End Date
1	NDC:73955-0001- 1	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:73955-0001- 2	710 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/30/2020	
3	NDC:73955-0001-3	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:73955-0001-	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/30/2020	
5	NDC:73955-0001-	59 mL in 1 BOTTLE, SPRAY; 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 - Cape Spirits Inc (042574182)

Establishment

Establishment				
Name	Address	ID/FEI	Business Operations	
Cape Spirits Inc		042574182	manufacture(73955-0001)	

Revised: 6/2020 Cape Spirits Inc