HAND SANITIZER- alcohol liquid Dancing Goat Distillery, 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)

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

18,927.1 mL NDC: 74544-001-01

ALCOHOL ANTISEPTIC 80% TOPICAL SOLUTION



ANTISEPTIC HAND RUB NON-STERILE SOLUTION

VOLUME: 18,927.10 ML

DO NOT DRINK

BOTTLED 03/ 29 / 2020 BATCH # 001

Drug Facts

Active Ingredient(s)

Alcohol 80% v/v

Purpose Antiseptic

Use[s]

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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HAND SANITIZER

alcohol liquid

Product Information				
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:74544-111
Route of Administration	TOPICAL			
Active Ingredient/Active Mo	iety			
Ingre	dient Name		Basis of Strength	Strength
8			Dasis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (AL			LCOHOL	80 mL in 100 mL
			0	-
ALCOHOL (UNII: 3K9958V90M) (AL Inactive Ingredients			LCOHOL	
ALCOHOL (UNII: 3K9958V90M) (AL Inactive Ingredients	COHOL - UNII:3K9958V90M)		LCOHOL	80 mL in 100 mL
ALCOHOL (UNII: 3K9958V90M) (AL Inactive Ingredients Ir	COHOL - UNII:3K9958V90M) agredient Name		LCOHOL	80 mL in 100 mL trength mL
ALCOHOL (UNII: 3K9958V90M) (AL Inactive Ingredients If GLYCERIN (UNII: PDC6A3C0OX)	COHOL - UNII:3K9958V90M) agredient Name		LCOHOL S 1.45 mL in 100	80 mL in 100 mL trength mL
ALCOHOL (UNII: 3K9958V90M) (AL Inactive Ingredients In GLYCERIN (UNII: PDC6A3C0OX) HYDROGEN PEROXIDE (UNII: BBX0	COHOL - UNII:3K9958V90M) agredient Name		LCOHOL S 1.45 mL in 100	80 mL in 100 mL trength mL
ALCOHOL (UNII: 3K9958V90M) (AL Inactive Ingredients In GLYCERIN (UNII: PDC6A3C0OX) HYDROGEN PEROXIDE (UNII: BBX0	COHOL - UNII:3K9958V90M) agredient Name		LCOHOL S 1.45 mL in 100	80 mL in 100 mL trength mL

Packaging							
# Item Code	Package Description	Marketing Start	Date Marketing End Date				
1 NDC:74544-111-11	18927.1 mL in 1 PAIL; Type 0: Not a Combi	nation Product 04/03/2020					
Marketing Information							
Marketing Categ	ory Application Number or Monog	aph Citation Marketing Start	Date Marketing End Date				
OTC monograph not	inal part333A	03/30/2020					

Labeler - Dancing Goat Distillery, LLC (117469685)

Establishment

Name	Address	ID/FEI	Business Operations
Dancing Goat Distillery, LLC		117469685	manufacture(74544-111)

Revised: 4/2020

Dancing Goat Distillery, LLC