EESOME HAND SANITIZER- alcohol gel Draga Laboratories

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.

E Hands anitizer

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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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)

Inactive ingredients

Aloe Barbadensis Leaf Juice, Aminomethyl Propanol, Aqua (Water), Carbomer

Package Label - Principal Display Panel

237 mL NDC: 75748-4520-01



473 mL NDC: 75748-4520-02 front



473 mL NDC: 75748-4520-02 back

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Other Information
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In active Ingredients Ane Barbadenis Leaf Juke, Aminomethyl Propanol, Aqua (Water), Carborner

3785 mL NDC: 75748-4520-03

ees me Distributed by Eesome, LLC 8046 Roswell Road, Suite 202 Atlanta, GA 30350 770-671-0657

E E S TO M E HAND SANITIZER FRAGRANCE FREE 80% ETHYL ALCOHOL

WITH 💥 ALOE

1 GALLON 128 FL.OZ (3,785 ML)

Drug Facts	
Active ingredient Bhil Akolol 80% vir	Parpose
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EESOME HAND SANIT	IZER				
alcohol gel					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75748-4520		
Route of Administration	TOPICAL				
Active Ingredient/Active Mo	liety				
Ingredient Name		Basis of Stre	ength Strength		
ALCOHOL (UNII: 3K9958V90M) (AI	COHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL		
T (1 T) 11 (
Inactive Ingredients					
	Ingredient Name		Strength		
CARBOMER HOMOPOLYMER, UNS	SPECIFIED TYPE (UNII: 0 A5MM	307FC)	0.5 mL in 100 mL		

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ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	0.36 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.1 mL in 100 mL

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
T	1	10000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/19/20/20			
	2	10000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product				
3	NDC:75748-4520- 3	10000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/19/2020			
Marketing Information						
	Marketing Categ	ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
0	TC monograph not	final part333A	10/19/2020			

Labeler - Draga Laboratories (610096534)

Establishment

Name	Address	ID/FEI	Business Operations
Draga Laboratories		610096534	manufacture(75748-4520)

Revised: 10/2020

Draga Laboratories