HAND SANITIZER BY SOURCERY- ethyl alcohol liquid Sourcery Ltd

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.

Hand Sanitizer 300ml imported by Sourcery Ltd asi/55000

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)

Ethyl Alcohol 75% 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 years 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

300 mL NDC: 76693-300-12



185mm

Drug Facts

Active ingredient

Purpose

- Hand sanitizer to help reduce bacteria on the skin that
- Recommended for repeated use

Warnings

- Flammable. Keep away from fire or flame.
- For external use only.
- •When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
- $\,$ ^Stop use and ask a doctor if irritation or rash appears and lasts.
- •Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away.



Drug Facts(cont.) Directions

- Place enough product in your palm to thoroughly cover your hands
- Rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product

Other information

- Do not store above 110°F (43 °C)
 May discolor certain fabrics or surfaces

Inactive ingredients Purified Water, Glycerin, Carbomer, Triethanolamine, DMDM Hydantoin, Vitamin E, Aloe Barbadensis Leaf Extract.

NOT FOR RETAIL SALE

FOR OVER THE COUNTER USE ONLY

Distributed by idegy, inc.

70mm

ethyl alcohol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76693-301	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	75 mL in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL		
HYDROGEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)	50 mL in 100 mL		

Product Characteristics			
Color	white (clear)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76693-301- 10	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/14/2020	
2	NDC:76693-301- 02	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/14/2020	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/14/2020	

Labeler - Sourcery Ltd (800787645)

Registrant - Sourcery Ltd (800787645)

Establishment				
Name	Address	ID/FEI	Business Operations	
Sourcery Ltd		800787645	manufacture (76693-301)	

Revised: 5/2020 Sourcery Ltd