HAND SANITIZER- hand sanitizer gel Zhejiang Enchant Cosmetic Co., 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.

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. Isopropyl Alcohol (75%, 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)

Isopropyl 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 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

59 ml in 1 bottle NDC: 52728-016-01



Drug Facts

Active Ingredient

Purpose

Ethyl Alcohol 75.0% v/v.....Antimicrobial

Uses · for handwashing to decrease bacteria on the skin · recommended for repeated use.

Warnings

For external use only.

Flammable, keep away from heat and flame.

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation and redness develop and persistfor more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • wet hands thoroughly with product briskly rub hands together until dry • supervise children under 6 years in the use of this product.

Other Information

- store at 20°C to 25°C (68°to 77°F)
 may discolor certain fabrics.

Inactive Ingredients: Water, Glycerin, Carbomer, Aloe Verg, Aminomethyl propanol, Vitamin E.

* Effective at eliminating 99.99% of many common harmful germs & bacteria. DISTRIBUTED BY HEALING SOLUTIONS, LLC - 4703 W BRILL ST, PHOENIX, AZ 85043 800-986-9240 WWW.HEALINGSOLUTIONS.COM MADE IN CHINA



Front

Back

尺寸: 30*50MM,

正面标:透明PE材质,彩色印刷,版面后面加白底,覆亮膜(左出标)

背面标: 白色PE材质, 彩色印刷, 覆亮膜 (右出标)

HAND SANITIZER

hand sanitizer gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:52728-016

Route of Administration EXTRACORPOREAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	44.25 mL in 59 mL
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Inactive Ingredients			
Ingredient Name	Strength		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.059 mL in 59 mL		
GLYCERIN (UNII: PDC6A3C0OX)	0.295 mL in 59 mL		
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)	0.177 mL in 59 mL		
WATER (UNII: 059QF0KO0R)	14.0951 mL in 59 mL		
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	0.118 mL in 59 mL		
TOCOPHEROL (UNII: R0ZB2556P8)	0.0059 mL in 59 mL		

ı	Packaging			
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:52728-016-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/22/2020	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	05/22/2020			

Labeler - Zhejiang Enchant Cosmetic Co., Ltd. (527284802)

Establishment					
Name	Address	ID/FEI	Business Operations		
Zhejiang Enchant Cosmetic Co., Ltd.		527284802	manufacture(52728-016)		

Revised: 5/2020 Zhejiang Enchant Cosmetic Co., Ltd.