HAND SANITIZER- ethyl alcohol gel ZHEJIANG JINGHUI COSMETICS SHARE 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. 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 70% 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

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

236mL NDC: 73905-016-04

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HAND SANITIZER	HAND SANITIZER	KEEP OUT OF REACH OF CHILDREN.
Kills 99.9% of Germs	Drug Facts Active impredient[s] Purpose Active impredient[s] Purpose Active impredient[s] Antiseptic Use[s] Hadd sanitare to help reduce bacteria that potentially can cause disease. For use when soo and water and validable. Wornings For external use only, Flammable. Keep away from heat or flame De not use Hand Sanitare to help reduce bacteria that potentially can cause disease. For use when soo and water and use only, Flammable. Keep away from heat or flame When using this horaclassic on organ. San away from heat or flame to not use Hand Sanitare to a subscription of the sanitare of the sanitare to reach or cause a subscription of the sanitare to a sanitare to a subscription of the sanitare to a sanitare to	Distributed by: Mosaic Bath & Spa LLC New York NY 10016 PRO:05/10/2020 EXP:05/09/2022 MADE IN CHINA Not tested on animals NDC: 73905-016-04 MULTION AND AND AND AND AND AND AND AND AND AN
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HAND SANITIZER					
ethyl alcohol gel					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:73905-016	
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	ety				
Ingred	lient Name		Basis of Strengtl	h	Strength
ALCOHOL (UNII: 3K9958V90M) (ALC	COHOL - UNII:3K9958V90M)		ALCOHOL	5	70 mL in 100 mL

Iı	nactive Ingredie	ents			
			Ingredient Name		Strength
A	MINO METHYL PRO	PAN	OL (UNII: LU49E6626Q)		
Pl	ROPYLENE GLYCO	L (U	JNII: 6DC9Q167V3)		
B	UTYLENE GLYCOL	L (UN	VII: 3XUS85K0RA)		
B	UTYL ACRYLATE (UNII	: 705NM8U35V)		
W	ATER (UNII: 059QF	0KO	0 R)		
D	ENATONIUM (UNII:	4IK2	2DF4OU)		
P	ackaging				
#	Item Code		Package Description	Marketing Start Date	Marketing End Date
1	NDC:73905-016-01	2 m	L in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:73905-016-02	60 1	mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:73905-016-03	150	mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:73905-016-04	236	mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:73905-016-05	500	mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:73905-016- 06	980) mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
7	NDC:73905-016-07	150	0 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
8	NDC:73905-016- 08	180	0 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
9	NDC:73905-016- 09	380	0 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
N	Iarketing Inf	orı	mation		
	Marketing Catego	ry	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
0	TC monograph not fi	nal	part333A	03/30/2020	

Labeler - Zhejiang Jinghui Cosmetics Share Co., LTD (529558167)

Establishment

Name	Address	ID/FEI	Business Operations
ZHEJIANG JINGHUI COSMETICS SHARE CO.,LTD		529558167	manufacture(73905-016)

Revised: 5/2020

ZHEJIANG JINGHUI COSMETICS SHARE CO.,LTD