HAND SANITIZER- alcohol liquid KIMJEM 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.

HAND SANITIZER

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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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



29.57 mL NDC: 77904-001-01

alcohol liquid							
Product Information	n						
Product Type		HUMAN OTC DRUG	Item Code (Source)		e)	NDC:77904-001	
Route of Administration	n	TOPICAL					
Active Ingredient/A	ctive Moi	ety					
Ingredient Name				Basis of Strength		Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)				ALCOHOL		80 mL in 100 mL	
Inactive Ingredients Ingredient Name					Strength		
GLYCERIN (UNII: PDC6A3C0OX)				1.45 mL in 100 mL			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				0.125 mL in 100 mL			
HYDROGEN PEROXIDE (011100 ()			0.125 IIIL III 10	UIIIL	
	·				0.125 IIIL III IC	0 IIIL	
	·				0.125 IIIE III 10	U IIL	
WATER (UNII: 059QF0KO	·				0.125 mL m to	U III.	
water (UNII: 059QF0KO Packaging	·	Package Description		Marketin		Marketing End Dat	
WATER (UNII: 059QF0KO Packaging # Item Code	00R)				ng Start Date		
WATER (UNII: 059QF0KO Packaging # Item Code	00R)	Package Description			ng Start Date		
WATER (UNII: 059QF0KO Packaging # Item Code 1 NDC:77904-001-01 29.	57 mL in 1 B0	Package Description			ng Start Date		
WATER (UNII: 059QF0KO Packaging # Item Code 1 NDC:77904-001-01 29.	57 mL in 1 B0	Package Description			ng Start Date		
WATER (UNII: 059QF0KO Packaging # Item Code	57 mL in 1 BC mation	Package Description	tion Product	05/01/2020	ng Start Date		

Labeler - KIMJEM LLC (102474907)

Registrant - KIMJEM LLC (102474907)

Revised: 7/2020

KIMJEM LLC