

HAND SANITIZER- alcohol liquid
EJ Formula 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

100 mL NDC: 78955-001-33



DRUG FACTS	
Active Ingredient	Purpose:
Alcohol 80%	Antiseptic
Uses Health care personnel hand rub to help reduce bacteria that potentially can cause disease	
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 or mouth. In case of contact with eyes, rinse eyes thoroughly with water	
Stop use and ask 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	
Direction 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, Deionized Water, Fragrance	
Barcode	

Made in USA
Dist. by EJ Formula LLC.
8950 Fullbright Ave, Chatsworth, CA 91311



SPRAY SANITIZER
3.3 OZ (100 ML)

HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78955-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
		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	
		WATER (UNII: 059QF0KO0R)		
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78955-001-33	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/01/2020	
2	NDC:78955-001-06	177 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/01/2020	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part333A	05/01/2020	

Labeler - EJ Formula LLC (131410315)

Establishment

Name	Address	ID/FEI	Business Operations
EJ Formula LLC		131410315	manufacture(78955-001)

Revised: 7/2020

EJ Formula LLC