HAND SANITIZER- alcohol gel JLA Distillery 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.

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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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.

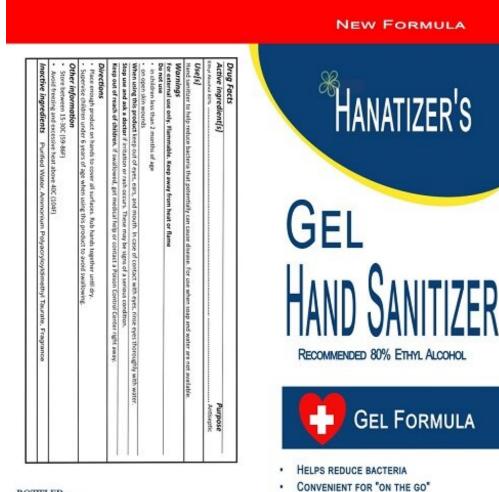
Other information

- 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



"FRAGRANCE FREE"

BOTTLED: AT JLA DISTILLERY ORLANDO, FL. 32822 WWW.SHOPJLA.NET



NDC 79005-101-05

HAND SANITIZER								
alcohol gel								
Product Information								
Product T ype		HUMAN OTC DRUG	Item Code (Source)		NDC:79005-101			
Route of Administra	ition	TOPICAL						
Active Ingredient/Active Moiety								
Ingredient Name				Basis of Strength	Strength			
ALCOHOL (UNII: 3K9	958V90M) (ALC	OHOL - UNII:3K9958V90M)		ALCOHOL	80 mL in 473 mL			
Inactive Ingredients								
	Strength							
GLYCERIN (UNII: PDC								
HYDRO GEN PERO XI								
WATER (UNII: 059QF								
Packaging								
# Item Code		Package Description	N	Marketing Start Date	Marketing End Date			
1 NDC:79005-101-05	473 mL in 1 BOTTLE; Type 0: Not a Combination		Product 0	7/06/2020				
Marketing Information								
Marketing Catego	ry Applicat	ion Number or Monograph Cit	ation 1	Marketing Start Date	Marketing End Date			
OTC monograph not fi	nal part333A	part333A		7/06/2020				

Labeler - JLA Distillery LLC (116280457)

Establishment							
Name	Address	ID/FEI	Business Operations				
JLA Distillery LLC		116280457	manufacture(79005-101)				

Revised: 12/2020

JLA Distillery LLC