HAND SANITIZER- alcohol gel Veralmex, S.P.R. de R.L.

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.

Gel Hand Sanitazer

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, Trolamine, Carbomer 940

Package Label - Principal Display Panel

Alcohol Antiseptic 80% Topical Solution

Hand Sanitizer



Antiseptic Hand Rub Non-sterile Solution

33.81 fl Oz (1000 mL)

Drug Facts	
Active Ingredient(s)	Purpose
Ethyl Alcohol 80% v/y	Ansiteptic
Use(s)	
Health care personnel hand rub to help reduce bacteria that potencil	aly can cause diseases.
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
thorughly with water.	
Stop use and ask a doctor if irritation on rash occurs. These may be sig	gns of a serious condition
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Other information	
-Store between 15-30C (59-86F)	
-Avoid freezing and excessive heat above 40C(104F)	
Inactive Ingredients	•
Distilled Water Glycerin Trolamine, Carbomer 940, Hydrogen Peroxide	

Manufactured by: Veralmex, S.P.R. DE R.L. Camino Panteon S/N Montemotelos (Los Tamez) Nuevo Leon, Mex. 67500 521-818-335-2761 Imported by: Veralmex US LLC

Tiki Conduminium 6608 Padre Blvd, suite 120 South Padre Island Tx. 78597 Ph. 956-761-2694

PRODUCT OF MEXICO

HAND SANITIZER alcohol gel Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:75268-001 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		
CARBOMER 940 (UNII: 4Q93RCW27E)	0.5 mL in 100 mL		
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			
TROLAMINE (UNII: 9O3K93S3TK)	85 mL in 100 mL		

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:75268-001- 01	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Veralmex, S.P.R. de R.L. (812784188)

Registrant - Mario Hidalgo (080666336)

Establishment			
Name	Address	ID/FEI	Business Operations
Veralmex, S.P.R. de R.L.		812784188	manufacture(75268-001)

Revised: 3/2020 Veralmex, S.P.R. de R.L.