HAND SANITIZER- alcohol gel Franslux, S.A. de C.V.

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 70% Alcohol

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



ANTI-BACTERIAL GEL GEL ANTIBACTERIAL

INGREDIENTS: Alcohol, Aqua (Water), Glycerin, Triethanolamine, Carbomer. DIRECTIONS: Apply and rub product onto hands and allow to dry. Apply sparingly as part of your daily cleansing routine. CAUTION: Avoid contact with eyes, in case of contact rinse thoroughly with water. Keep away from flame. Keep closed. Keep out of reach of children. For external use only.

INGREDIENTES: Alcohol, Agua, Glicerina, Trietanolamina, Carbomero. MODO DE EMPLEO: Aplicar una cantidad considerable del Gel Antibacterial en la palma de las manos dar un ligero masaje hasta la total absorción del gel. PRECAUCIONES: Evite el contacto con los ojos, si esto ocurre enjuagar con abundante agua. No ingerirse. Uso externo solamente. Manténgase alejado del fuego o llamas. Manténgase tapado. No se deje al alcance de los niños.

Made in Mexico for: / Fabricado en México para: AIR-VAL INTERNATIONAL S.A., Ayala 82, 28001 MADRID, SPAIN / ESPAÑA.

Made in Mexico by: / Fabricado en México por: FRANSLUX, S.A. DE C.V., Cuautipark II, Nave 2-B, Carr. Pte. Grande Las Ánimas s/n, Col. La Victoria, Cuautitlán, Estado de México C.P. 54834.



240 mL NDC: 55260-003-00

HAND SANITIZER

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55260-003	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	73.06 mL in 100 mL		

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6 A3C0 OX)	0.99 mL in 100 mL		
WATER (UNII: 059QF0KO0R)	24.67 mL in 100 mL		
TROLAMINE (UNII: 9O3K93S3TK)	0.68 mL in 100 mL		
CARBOMER 940 (UNII: 4Q93RCW27E)	0.6 mL in 100 mL		

l	Packaging				
	# It	tem Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC	:55260-003-	240 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/29/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	05/29/2020		

Labeler - Franslux, S.A. de C.V. (810312348)

Establishment				
Name	Address	ID/FEI	Business Operations	
Franslux, S.A. de C.V.		8 10 312348	manufacture(55260-003)	

Revised: 5/2020 Franslux, S.A. de C.V.