

HAND SANITIZER- handwash gel

Mexiquim SA de CV

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

480 mL NDC: 80137-420-02

Antibacterial Handwash ABSCLEAN Natural Mint protects, softens and hydrates your skin every day leaving a refreshing cleansing sensation as it contains Aloe Vera. Its formula acts against germs, viruses and bacteria and helps prevent diseases by direct and indirect contact.

Instructions for use

1. Apply a small amount to the palm of the hand.
2. Gently rub into a rich lather.
3. Rinse hands with plenty of water.

Warnings

Keep the product out of the reach of children. Do not apply to cut or irritated skin. External use. Avoid contact with the eyes. In case of irritation, discontinue use and consult a doctor.

Ingredients

Water, Sodium laureth sulfate, Pentaerythrityl Tetrastearate, Betaine, Coconut Amide, Phenoxyethanol, Triethanolamine, Sodium Chloride.

Made in Mexico by: Grupo Quimae SA de CV, Jose Marti 203 B, Col. Tlacopa Toluca Edo de Mex C.P. 50010

Distributed and commercialized by: Mexiquim, S.A. de C.V. Calle de la Barranca 3-A Col, Tlacopa Toluca, Edo. de México, C.P. 50010

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EXTRA SOFT

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QR CODE

HAND SANITIZER

handwash gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80137-420
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
COCAMIDE (UNII: 3YXD33R71G)	1 g in 100 g
SODIUM CHLORATE (UNII: T95DR77GMR)	1.5 g in 100 g
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	12 g in 100 g
PEG-150 PENTAERYTHRITYL TETRASTEARATE (UNII: 8L40OQ76AM)	0.25 g in 100 g
WATER (UNII: 059QF0K00R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.2 g in 100 g

CO CAMIDO PROPYL BETAINE (UNII: 5OCF3O11KX)

3 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80137-420-02	480 g in 1 BOTTLE; 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 - Mexiquim SA de CV (589907237)

Establishment

Name	Address	ID/FEI	Business Operations
Mexiquim SA de CV		589907237	manufacture(80137-420)

Revised: 11/2020

Mexiquim SA de CV