HAND SANITIZER- antiseptic foam aerosol, foam Innovaciones Tecnologicas Concar 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

200 L NDC: 79279-521-05

PROPERTIES

- Alcohol, fragance and dye-free
- Contains emollients



7 503024 307283

52.8 gal (200 L)

LOT EXP DATE.___

www.sq2pro.com.mx

It is not medicine Only for external use

Drug Facts

Active ingredient

Use

on the skin Warnings For external use only. Do not use in the eyes Stop use and ask a doctor if

consult a doctor.

Directions

Benzalkonium chloride 0.13%Antiseptic

for hand washing to decrease bacteria

■ irritation and redness develop. If condition persists for more than 72 hours

Keep out of reach of children. If

swallowed, get medical help or contact a Poison Control Center right away.

wet hands thoroughly with product and

Citric Acid, Hydroxyethylcellulose, Lanolin Ethoxylated, Povidone, Propylene

Questions or Comments?

INNOVACIONES TECNOLOGICAS CONCAR, S.A. DE C.V.

Palenque No. 591, Col. Letrán Valle, Benito Juárez, C.P. 03650, CDMX, México.

allow to dry without wiping Other information avoid use in case of hypersensitivity to

benzalkonium chloride

Glycol, Water

818-619-2735

DISTRIBUTED BY:

Made in México.

Inactive ingredients

Purpose

HUMAN OTC DRUG	Item Code (Source)		NDC:79279-521	
TOPICAL				
ety				
Ingredient Name			Basis of Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)		BENZALKONIUM CHLORIDE		0.8 kg in 100 L
ľ	TOPICAL ety redient Name	ety redient Name	ety redient Name F5UM2KM3W7) (BENZALKONIUM - BENZALKONIUM	ety redient Name F5UM2KM3W7) (BENZALKONIUM - BENZALKONIUM

Inactive Ingredien	ts						
Ingredient Name			Strength				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			1.45 L in 100 L				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			0.125 L in 100 L				
WATER (UNII: 059QF0F	(O0R)						
Packaging # Item Code Package Description Marketing Start Date Marketing End Date							
	Package Description		Marketing Lift Date				
I NDC:/92/9-521-05	200 L in 1 DRUM; Type 0: Not a Combination Product	03/30/2020					
Marketing Information							
Marketing Info	rmation						
Marketing Info Marketing Category		Marketing Start	Date Marketing End Date				

Labeler - Innovaciones Tecnologicas Concar SA de CV (951578588)

Establishment

Name	Address	ID/FEI	Business Operations
Innovaciones Tecnologicas Concar SA de CV		951578588	manufacture(79279-521)

Revised: 6/2020

Innovaciones Tecnologicas Concar SA de CV