HAND SANITIZER- benzalkonium chloride gel
NATURATLALI S DE RL 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.

77143-002-01
Naturatlali 946mL

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)

Benzalkonium Chloride 0.13% w/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.

Do not use

- on open 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

water, hydroxyethyl cellulose, glycerin, methylparabene, propylparabene, propylene glycol, DMDM Hydantoin, FD&C Blue No 1

Package Label - Principal Display Panel

946 mL NDC: 77143-002-01

HAND SANITIZER
benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77143-002
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	0.25 mL in 100 mL
HYDROXYETHYL CELLULOSE (100 MPA.S AT 2%) (UNII: R33S7TK2EP)	0.01 g in 100 mL
WATER (UNII: 059QF0KO0R)	
METHYLPARABEN (UNII: A2I8C7HI9T)	0.025 g in 100 mL
PROPYLPARABEN (UNII: Z8IX2SC1OH)	0.025 g in 100 mL
DMDM HYDANTOIN (UNII: BYR0546TOW)	0.025 g in 100 mL
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	0.001 g in 100 mL
DIPROPYLENE GLYCOL (UNII: E107L85C40)	0.025 mL in 100 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77143-002-01	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:77143-002-18	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020	

Marketing Information

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

Labeler - NATURATLALIS DE RL DE CV (951577202)

Registrant - NATURATLALIS DE RL DE CV (951577202)

Establishment

Name	Address	ID/FEI	Business Operations
NATURATLALIS DE RL DE CV		951577202	manufacture(77143-002) , pack(77143-002) , label(77143-002)

Revised: 9/2020 NATURATLALIS DE RL DE CV