HAND SANITIZER- alcohol gel
CA-Botana International

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 Dr Bump

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 65% 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

purified water USP, Aloe barbadensis extract, Hamamelis virginiana (Whitchhazel) extract, Polyquaternium-37, Cucumis sativus (Cucumber) extract, Centella asiatica (Gotu kola) extract, Equisetum fluviatitile (Horsetail) extract, Rosmarinus officinalis (Rosemary) extract, Aesculus hippocastanum (Horse chestnut) extract, Camellia sinensis (Green tea) extract, Olea europaea (Olive) extract, Eucalyptus globulus extract, Propanediol, Phenoxyethanol, Potassium Sorbate, Ethylhexylglycerin, Sodium Phytate, Benzalkoniun Chloride.

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:35192-035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CAMELLIA SINENSIS WHOLE (UNII: C5M4585ZBZ)	
PROPANEDIOL (UNII: 5965N8W85T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
EQUISETUM FLUVIATILE WHOLE (UNII: D6MQP77VTM)	
ROSEMARY (UNII: IJ67X351P9)	
HEXASODIUM PHYTATE (UNII: ZBX50UG81V)	
CENTELLA ASIATICA LEAF (UNII: 6810070TYD)	
HORSE CHESTNUT (UNII: 3C18L6RJAZ)	
WITCH HAZEL (UNII: 101I4J0U34)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
POLYQUATERNIUM 37 (200 MPA.S) (UNII: 67C1D6YV24)	
GREEN OLIVE (UNII: 6HD2W46UEG)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CUCUMBER (UNII: YY7C30VXJT)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:35192-035-13	118.3 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2020	

Marketing Information

Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph not final	part333A	05/18/2020	

Labeler - CA-Botana International (106276728)

Registrant - Rodolfo Ugelstad (106276728)

Establishment

Name	Address	ID/FEI	Business Operations
CA Botana International		106276728	manufacture(35192-035)