HAND SANITIZER- alcohol gel Bits Systems, LLC

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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Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right

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



Active Ingredients Ethyl Alcohol 70% v/v Antiseptic - Lab Tested

Use Hand sanitizer kills 99.99% of germs that can cause disease. For use when soap and water are not available.

Warnings For external use only. Flammable. Keep away from heat or flame. Allergen warning, contains essential oils for fragrance. See Inactive Ingredients.



Do Not Use on children less than 2 months of age, or 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, this may be a sign of a serious condition.

Keep Out of Reach of Children

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

Directions

•Place enough product on hands to cover all surfaces.

•Rub hands together until dry. Thoroughly rub into crevices under and around nails.

•Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information Store between 15-30C (59-86F), Avoid freezing or excessive heat above 40C (104F)

Inactive Ingredients hydrogen peroxide, aloe vera, glycerin, sterile water, carbomer and/or acrylate copolymer, cymbopogon citratus, lavandula, rosmarinus officinalis, triethanolamine

Questions? (844) 530-8788 support@sanesanitizer.com

SaneSanitizer.com 8 The Green, Suite #5697, Dover, DE 19901

Made in the USA

FDA NDC: 76580-840-XX





Lemongrass, Lavender, & Rosemary

- Antiseptic - 70% Alcohol -

4 fl oz

Made in America

000 mL NDC: 00000-000-00

HAND SANITIZER alcohol gel						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) ND		NDC:76580-119		
Route of Administration	TOPICAL					
Active Ingredient/Active Moi	ots					
Ingredient Name			Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	80 mL in 100 mL		
Inactive Ingredients						
Ingredient Name				Strength		
ALOE VERA LEAF POLYSACCHARII						
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)						
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL					

HYDRO GEN PERO X	0.125 mL in 100 mL						
WATER (UNII: 059QF							
Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:76580-119- 01	3900 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/22/2020					
Marketing Information							
Marketing Catego	ry Application Number or Monograph Citatio	n Marketing Start Date	Marketing End Date				
OTC monograph not f	inal part333A	03/30/2020					

Labeler - Bits Systems, LLC (117495902)

Establishment						
Name	Address	ID/FEI	Business Operations			
Bits Systems, LLC		117495902	manufacture(76580-119)			

Revised: 7/2020

Bits Systems, LLC