HAND SANITIZER- alcohol gel
Bit 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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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

PREVENTION STARTS WITH

SANE SANITIZER .COM

Hand Sanitizer Gel

with

Vitamin E and Aloe Vera

Lemongrass & Lavendar Scented

250 US gallons
NDC 76580-XXX-XX

62% Ethyl Alcohol

Made in America

Active Ingredients Ethyl Alcohol 62% 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 FDA NDC: 76580-XXX-XX

SaneSanitizer.com 8 The Green, Suite #5697, Dover, DE 19901 Made in the USA

000 mL NDC: 00000-000-00

HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76580-122
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF POLYSACCHARIDES (UNII: W21O437517)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:76580-122-01	946352 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/27/2020	

Marketing Information

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

Labeler - Bit Systems, LLC (117495902)

Establishment

Name	Address	ID/FEI	Business Operations
Bits Systems, LLC		117495902	manufacture(76580-122)

Revised: 7/2020 Bit Systems, LLC