

DR JS HAND SANITIZER GEL- alcohol gel

Froggy's Fog 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

Drug Facts
Active ingredient Purpose Ethyl Alcohol 70%.....Antiseptic
Uses: • Hand Sanitizer to help decrease bacteria on the skin that could cause disease. • Recommended for repeated use.
Warnings For external use only, use on hands only.
Flammable. Keep away from fire or flame.
When using this product: • Avoid contact with eyes. In case of contact with eyes, flush thoroughly with water. • Avoid contact with broken skin. • Do not inhale or ingest.
Stop use and ask a doctor if • Irritation and redness develops. • Condition persists for more than 72 hours.
Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
Directions • Pump as needed into your palms and thoroughly spread on both hands. Rub into skin until dry. • For children under 6, use only under adult supervision. • Not recommended for infants.
Other information • Store below 110°F (43°C) • May discolor some fabrics.
Inactive ingredients: acrylate polymer, aminoethyl propanol, fragrance, glycerin, hydrogen peroxide, polyethylene glycol, purified water USP


 Dr. J's[™]
 Natural
 ADVANCED
 FORMULA
**HAND
 SANITIZER**
 ALOE VERA
 REFRESHING GEL
 Kills More Than 99.99% of
 Bacteria
 16.9 FL OZ (500 mL)




 Dr. J's[™]
 Natural

Distributed By: QYK Brands, LLC.

Manufactured:
 302 Rutherford Ln
 Columbia, TN 38401
www.DrJsNatural.com

Questions or Comments
 Please Call: 1-888-308-7078
 NDC : 75682-997-16



500 mL NDC: 75682-997-16

DR JS HAND SANITIZER GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75682-997
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE (UNII: V5VD430 YW9)	

GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75682-997-16	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

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

Labeler - Froggy's Fog LLC (144348161)

Registrant - Froggy's Fog LLC (144348161)

Establishment

Name	Address	ID/FEI	Business Operations
Froggy's Fog LLC		144348161	manufacture(75682-997)

Revised: 12/2020

Froggy's Fog LLC