HAND SANITIZER- alcohol liquid Leidos Biomedical Researh, Inc.

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

Hand Sanitizer Labels (front label and drug facts label)

Alcohol 80 % v/v

Antiseptic

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

For external use only. Flammable. Keep away from heat or flame

- In children less than 2 months of age
- On open skin wounds

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.

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 40 C (104F)

Inactive ingredients glycerin, hydrogen peroxide, purified water USP

450 mL NDC 74167-8001-1

Alcohol Antiseptic 80% Topical Solution

Antiseptic Hand Rub Non-sterile Solution

450 mL L/N: 20-XXX

Mfg for VRC by Leidos Biomedical Research Frederick National Lab

Not for Sale

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

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)
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Inactive ingredients glycerin, hydrogen peroxide, purified water USP

450 mL NDC: 74167-8001-1

alcohol liquid

Product Information

Product Type HUMAN OTC DRUG NDC:74167-8001 Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (LINII) 2K0059X00M (ALCOHOL LINII) 2K0059X00M	AT COHOT	90 mI in 100 mI

ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII: 3K9958 V90M) 80 mL in 100 mL

Inactive Ingredients

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Ingredient Name	Strength			
GLYCERIN (UNII: PDC6 A3C0 OX)				
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)				

WATER (UNII: 059QF0KO0R)

Packaging

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	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:74167- 8001-2	1 in 1 BOX	03/30/2020	
	1 NDC:74167- 8001-1	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

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

Labeler - Leidos Biomedical Researh, Inc. (079940150)

Revised: 3/2020 Leidos Biomedical Researh, Inc.