# HAND SANITIZER- is opropyl alcohol gel RD LINCOLN INTERNATIONAL, 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.

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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. Isopropyl Alcohol (75%, 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)

Isopropyl Alcohol 75% 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**

glycerin, hydrogen peroxide, purified water USP

# Package Label - Principal Display Panel

236ml NDC 79689-002-08



50ml NDC 79689-002-02

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		Skin Care Facts		
	1.7 Fluid Ounces	Action Ingredient Isopropyl Alcohol 75% v/v	Purpose Antiseptic	
	7 Fluid	USE For external use- Hand Saniti the reduction of bacteria on the skin.	zer for assisiting Antiseptic.	
ANTI-MICROB HAND GEL For Advanced Hand Sanitizing Kills 99.9% of Illness Causing Germs		<b>Warnings</b> Flammable. Keep away from open flame or fire.		
		For external use only		
		Stop use and ask doctor if irritation or rash occurs. These may be signs of serious condition. Avoid use in the eyes, ears, mouth. In case, of contact with eyes, rinse thoroughly with water.		
		<b>Directions</b> Place enough product on hands to cover hands together until dry. Supervise childre age when using this product to avoid swal	n under 6 years of	
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Made in USA		50ml NDC: 79689-002-02		

HAND SANITIZER
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isopropyl alcohol gel							
Product Information							
Product Type	bduct Type HUMAN OTC DRUG Item Code (Source		xe) NDC:79689-002		689-002		
Route of Administration	TOPICAL						
Active Ingredient/Active Moiety							
				_			
I	ngredient Name		Basis of Str	rength	Strength		
I: ISOPROPYL ALCOHOL (UNII: ND2 UNII:ND2M416302)	•	L -	Basis of Str ISOPROPYL ALCOHOL	rength	Strength 75 mL in 100 mL		
ISOPROPYL ALCOHOL (UNII: ND2	•	L -	ISOPROPYL	rength	75 mL		
ISOPROPYL ALCOHOL (UNII: ND2	•	L -	ISOPROPYL	rength	75 mL		
ISOPROPYL ALCOHOL (UNII: ND2	•	L -	ISOPROPYL	rength	75 mL		
ISOPROPYL ALCOHOL (UNII: ND2 UNII:ND2M416302) Inactive Ingredients	•	L -	ISOPROPYL	rength Streng	75 mL in 100 mL		
ISOPROPYL ALCOHOL (UNII: ND2 UNII:ND2M416302) Inactive Ingredients	M416302) (ISOPROPYL ALCOHO ngredient Name	L -	ISOPROPYL	Streng	75 mL in 100 mL		
ISOPROPYL ALCOHOL (UNII: ND2 UNII:ND2M416302) Inactive Ingredients	M416302) (ISOPROPYL ALCOHO ngredient Name	L -	ISOPROPYL ALCOHOL	Streng	75 mL in 100 mL		

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:79689-002- 08	160000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			
2	NDC:79689-002- 02	25000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			
N	Aarketing Inf	ormation				
	Aarketing Inf Marketing Catego		Marketing Start Date	Marketing End Date		

Labeler - RD LINCOLN INTERNATIONAL, INC. (080317128)

Establishment						
Name	Address	ID/FEI	<b>Business Operations</b>			
RD LINCOLN INTERNATIONAL, INC.		080317128	manufacture(79689-002)			

Revised: 7/2020

RD LINCOLN INTERNATIONAL, INC.