HAND SANITIZER- alcohol gel
Rx SAN 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.

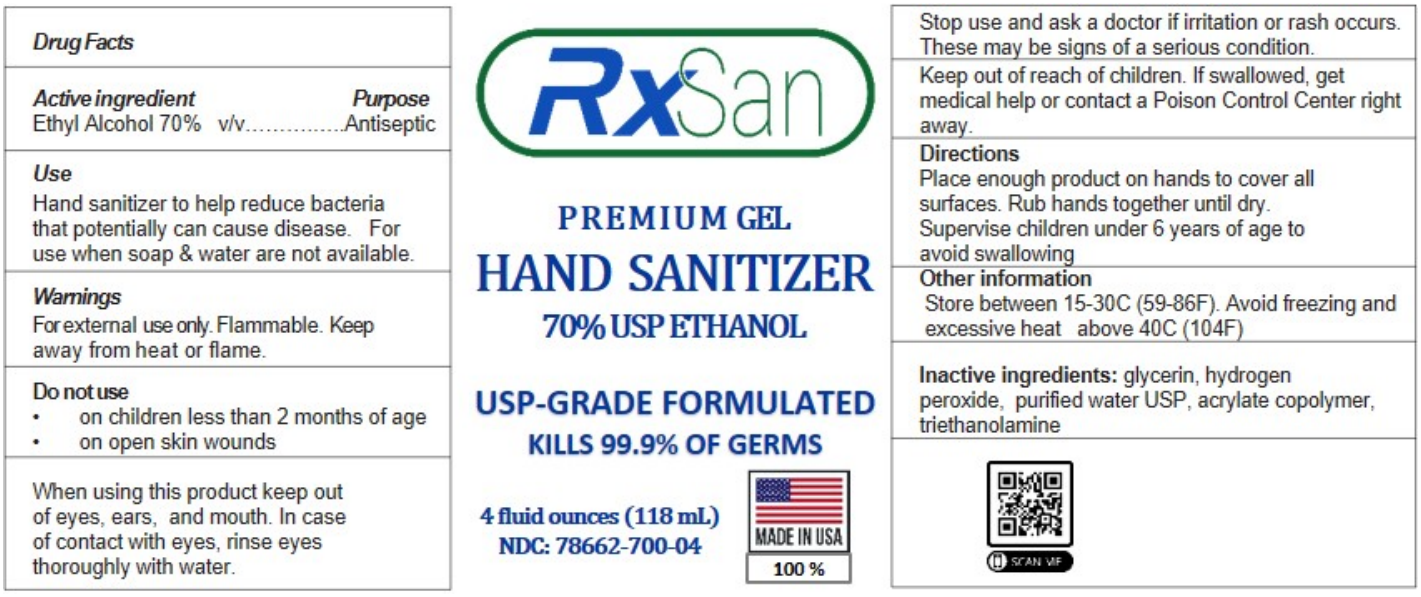
Other information

- 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% v/v.....Antiseptic

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Inactive ingredients: glycerin, hydrogen peroxide, purified water USP, acrylate copolymer, triethanolamine

SCAN ME

RxSan

PREMIUM GEL

HAND SANITIZER

70% USP ETHANOL

USP-GRADE FORMULATED

KILLS 99.9% OF GERMS

8 fluid ounces (237 mL)
NDC: 78662-700-08

MADE IN USA
100 %

Drug Facts

Active ingredient — **Purpose**
Ethyl Alcohol 70% v/v.....Antiseptic

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SCAN ME

RxSan

PREMIUM GEL

HAND SANITIZER

70% USP ETHANOL

USP-GRADE FORMULATED

KILLS 99.9% OF GERMS

12 fluid ounces (355 mL)
NDC: 78662-700-12

MADE IN USA
100 %

Drug Facts

Active ingredient — **Purpose**
Ethyl Alcohol 70% v/v.....Antiseptic

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Inactive ingredients: glycerin, hydrogen peroxide, purified water USP, acrylate copolymer, triethanolamine

SCAN ME

RxSan

PREMIUM GEL

HAND SANITIZER

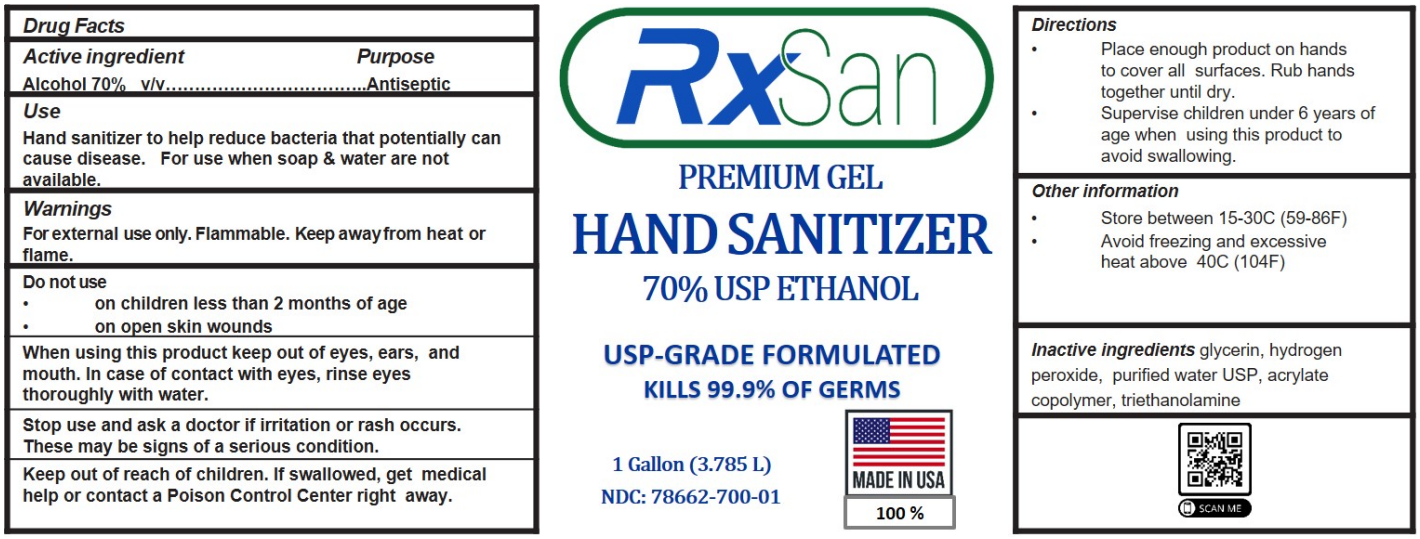
70% USP ETHANOL

USP-GRADE FORMULATED

KILLS 99.9% OF GERMS

16 fluid ounces (473 mL)
NDC: 78662-700-16

MADE IN USA
100 %



HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78662-700(NDC:74171-300)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 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)	
TROLAMINE (UNII: 9O3K93S3TK)	0.1287 mL in 100 mL
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	1.8 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78662-700-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/03/2020	
2	NDC:78662-700-16	473 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/03/2020	
3	NDC:78662-700-12	355 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/03/2020	
4	NDC:78662-700-04	118 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	06/03/2020	
5	NDC:78662-	1041000 mL in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0:	06/03/2020	

5	700-75	Not a Combination Product	06/03/2020	
6	NDC:78662-700-50	18927 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/03/2020	

Labeler - Rx SAN LLC (117536764)

Registrant - Rx SAN LLC (117536764)

Establishment

Name	Address	ID/FEI	Business Operations
Rx SAN		117536764	repack(78662-700)

Revised: 1/2021 Rx SAN LLC