

SANITY HAND SANITIZER- alcohol liquid

Damon Industries

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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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

1893 mL NDC: 74682-001-07



FIGHT THE GOOD FIGHT

Renown[®]
HEALTH

HAND SANITIZER
ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
64 OZ (1893 ML)

Drug Facts

Active Ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

Use[s]

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For use when soap and water are not available.

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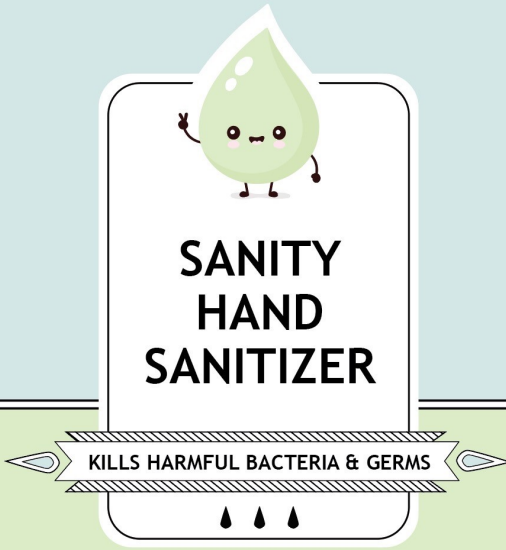
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Manufactured by Damon Industries
822 Packer Way, Sparks, NV 89431

1893 ml NDC: 74682-001-07




**SANITY
HAND
SANITIZER**

KILLS HARMFUL BACTERIA & GERMS

ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
NON-STERILE SOLUTION 64 OZ (1893 ML)

Drug Facts	
Active Ingredient[s] Alcohol 80% v/v.....	Purpose Antiseptic
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Manufactured by Damon Industries
822 Packer Way, Sparks, NV 89431



473 ml NDC: 74682-001-39



**FIGHT
THE
GOOD
FIGHT**

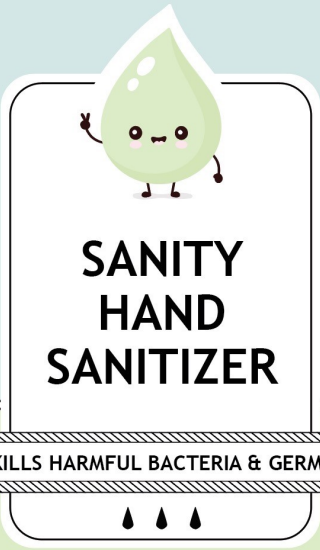
Renown[®]
HEALTH

HAND SANITIZER
ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
16 OZ (473 ML)

Drug Facts	
Active Ingredient[s] Alcohol 80% v/v.....	Purpose Antiseptic
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Manufactured by Damon Industries
822 Packer Way, Sparks, NV 89431

473 ml NDC: 74682-001-39



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HAND
SANITIZER**


KILLS HARMFUL BACTERIA & GERMS

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TOPICAL SOLUTION
NON-STERILE SOLUTION 16 OZ (473 ML)

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SANITY HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74682-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74682-001-39	473 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74682-001-07	1873 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 - Damon Industries (034954339)

Establishment

Name	Address	ID/FEI	Business Operations
Damon Industries		034954339	manufacture(74682-001)

Revised: 4/2020

Damon Industries