HAND SANITIZER- alcohol gel One Plastic 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) (70%, 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. Triethenolamone (0.125% v/v).
- d. Carbopol.

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 70% 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, Triethenolamone, Carbopol

Package Label - Principal Display Panel



Hand Sanitizer
Non-sterile Solution

33.81 oz (1000 mL)

Drug Facts

Active ingredient[s]

P urpose

Alcohol 70% v/v

.Antiseptic

Use[s]

Hand Sanitizento help reduce bacteria that potentially can cause disease. For use when so ap and water are not available.

Warnings

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

Do not use

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Directions

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Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 400 (104F)

Inactive ingredients Glycerin, Carbopol, Triethanolamine

1000 mL NDC: 74860-512-03 3750 mL NDC: 74860-512-05



Hand Sanitizer
Non-sterile Solution

126.80 oz (3750 mL)

Drug Facts

Active ingredient[s]

Alcohol 70% v/v

P urpose

.Antiseptic

Use[s]

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Warnings

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Directions

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Inactive ingredients Glycerin, Carbopol, Triethanolamine

500 mL NDC: 74860-512-02



Hand Sanitizer
Non-sterile Solution

16.9 oz (500 mL)

Drug Facts

Active ingredient[s]

Alcohol 70% v/v

Purpose

Antisentic

Use[s]

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Warnings

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Other information

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Inactive ingredients Glycerin, Carbopol, Triethanolamine

1500 ml NDC: 74860-512-04



Hand Sanitizer
Non-sterile Solution

50.72 oz (1500 mL)

Drug Facts

Active ingredient[s]

Purpose

Alcohol 70% v/v

Antiseptio

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Hand Sanitizento help reduce bacteria that potentially can cause disease. For use when so ap and water are not available.

Warnings

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Do not use

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Directions

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Other information

- Store between 15-30C (59-86F)
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Inactive ingredients Glycerin, Carbopol, Triethanolamine

150 ml NDC: 74860-512-01



Hand Sanitizer
Non-sterile Solution

5.07 oz (150 mL)

Drug Facts

Active ingredient[s]

P urpo se

Alcohol 70% v/v

.Antiseptic

Use[s]

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Warnings

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- Store between 15-30C (59-86F)
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Inactive ingredients Glycerin, Carbopol, Triethanolamine

alcohol gel

| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:74860-512 |
|--------------|----------------|--------------------|---------------|

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| ı | Ingredient Name | Basis of Strength | Strength |
|---|--|-------------------|-----------------|
| ı | 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 | | | |
| HYDRO GEN PERO XIDE (UNII: BBX060AN9V) | 0.125 mL in 100 mL | | | |
| WATER (UNII: 059QF0KO0R) | | | | |

| Packaging | | | | |
|-----------|------------------|--|-----------------------------|--------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:74860-512-01 | 150 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/10/2020 | |
| 2 | NDC:74860-512-02 | 500 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/10/2020 | |
| 3 | NDC:74860-512-03 | 1000 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/10/2020 | |
| 4 | NDC:74860-512-04 | 1500 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/10/2020 | |
| 5 | NDC:74860-512-05 | 3750 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/10/2020 | |

| Marketing Information | | | | |
|-------------------------|--|----------------------|--------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC monograph not final | part333A | 04/10/2020 | | |
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Labeler - One Plastic LLC (080529110)

| Establishment | | | | |
|-----------------|---------|-----------|------------------------|--|
| Name | Address | ID/FEI | Business Operations | |
| One Plastic LLC | | 080529110 | manufacture(74860-512) | |

Revised: 4/2020 One Plastic LLC