HAND SANITIZER PLUS- alcohol gel Clinical Resolution Laboratory, Inc.

Hand Sanitizer Plus

DRUG FACTS

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses:

- To decrease bacteria on the skin that could cause disease.
- Recommended for repeated use.

Warnings:

For external use only: Hands

Flammable, keep away from fire or flame.

When using this product

- Keep out of eyes. In case of contact with eyes, rinse thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.

Stop use and ask a doctor if

- Irritation and redness develop.
- Condition persists for more than 72 hours.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands thoroughly with product and allow to dry without wiping.
- For children under 6, use only under adult supervision.
- Not recommended for infants.

Other information

• Do not store above 105°F.

- May discolor some fabrics or surfaces.
- Harmful to wood finishes and plastics.

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Vera Leaf Juice, Aminomethyl Propanol, Fragrance, Glycerin, Hamamelis Virginiana (Witch Hazel) Leaf Extract, Isopropyl Alcohol, Isopropyl Myristate, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Tocopheryl Acetate, Water

Package Labeling:



Package Labeling: 60ml

Ebanel

HAND **SANITIZER**

PLUS

INSTANTLY KILLS 99.9% OF GERMS

With moisturizing Aloe and Vitamin E



Dermatologist-Tested Non-Irritating No Propylene Glycol Made in the USA

2 FL.OZ | 60 mL

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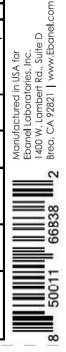
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HAND SANITIZER PLUS

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63742-040

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) **ALCOHOL** 0.7 mL in 1 mL

Inactive Ingredients

| Ingredient Name | Strength | | | |
|---|----------|--|--|--|
| CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO) | | | | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | | | | |
| AMINOMETHYLPROPANOL (UNII: LU49E6626Q) | | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | | | | |
| HAMAMELIS VIRGINIANA LEAF (LINII: TO7111161SV) | | | | |

ISOPROPYL ALCOHOL (UNII: ND2M416302) ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)

| TEA TREE OIL (UNII: VIF565UC2G) | |
|---|--|
| .ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| WATER (UNII: 059QF0KO0R) | |

| Packaging | | | | | | |
|---------------------------|---|-------------------------|-----------------------|--|--|--|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date | | | |
| 1 NDC:63742-040-00 | 118 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | 03/01/2020 | | | | |
| NDC:63742- 040-01 | 60 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product | 05/11/2020 | | | | |

| Marketing Information | | | | |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | 505G(a)(3) | 03/01/2020 | | |
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Labeler - Clinical Resolution Laboratory, Inc. (825047942)

Revised: 12/2023 Clinical Resolution Laboratory, Inc.