

HANDY SOLUTIONS INSTANT HAND SANITIZER WITH VITAMIN E 70% ALCOHOL-ethyl alcohol gel

Navajo Manufacturing Company 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.

Handy Solutions Instant Hand Sanitizer with Vitamin E 70% Alcohol

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

- for hand sanitizing to decrease bacteria on the skin.
- recommended for repeated use

Warnings

For external use only.

Flammable. Keep away from heat or flame.

When using this product, do not use in or near eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or redness develop and persist 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
- briskly rub hands together until dry
- supervise children under 6 years in the use of this product

Other information

- store at room temperature
- may discolor certain fabrics

Inactive ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, tocopheryl acetate.

Questions or comments?

Call 1-800-525-5097

Principal Display Panel



Drug Facts
Active ingredient Purpose
Ethyl Alcohol
70% v/v Antimicrobial

Uses
• for hand sanitizing to decrease bacteria on the skin.
• recommended for repeated use

Distributed by: Handy Solutions®
A Trademark of Navajo Mfg. Co. Inc.
Denver, CO 80216
Made in China

No. 15942 V1

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Drug Facts (continued)
Warnings
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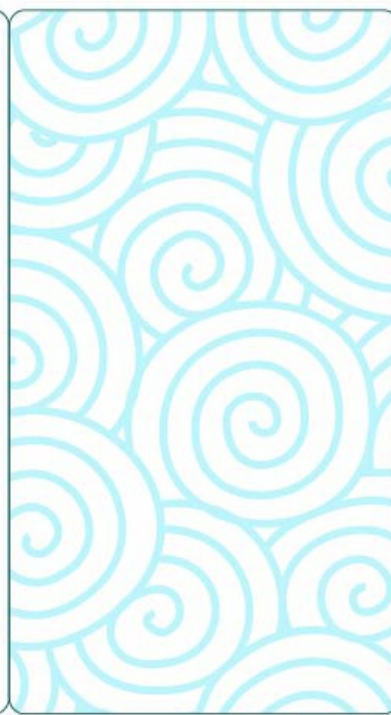
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| | |
|--|---------------------------------|
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| Active ingredient Ethyl Alcohol 70% v/v | Purpose Antimicrobial |
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| | |



HANDY SOLUTIONS INSTANT HAND SANITIZER WITH VITAMIN E 70% ALCOHOL

ethyl alcohol gel

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:67751-197 |
| 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 |
|--|----------|
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| .ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| TROLAMINE (UNII: 9O3K93S3TK) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| WATER (UNII: 059QF0KO0R) | |
| CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|-----------|---------------------|----------------------|--------------------|
|---|-----------|---------------------|----------------------|--------------------|

| | | | | |
|---|------------------|---|------------|--|
| 1 | NDC:67751-197-01 | 59 mL in 1 BOTTLE; Type 0: Not a Combination Product | 05/12/2020 | |
| 2 | NDC:67751-197-03 | 236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product | 05/12/2020 | |
| 3 | NDC:67751-197-02 | 2 in 1 BOX | 05/12/2020 | |
| 3 | | 59 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333E | 05/12/2020 | |

Labeler - Navajo Manufacturing Company Inc. (091917799)

Registrant - Navajo Manufacturing Company Inc. (136941411)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--|---------|-----------|------------------------|
| China Ningbo Shangge Cosmetic Technology Corp. | | 529287434 | manufacture(67751-197) |

Revised: 5/2020

Navajo Manufacturing Company Inc.