HANDY SOLUTIONS INSTANT HAND SANITIZER WITH ALOE AND VITAMIN E- alcohol liquid
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 Aloe & Vitamin E

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

- A hand sanitizer to help reduce bacteria on the skin
- A hand and lower arm sanitizer to help reduce bacteria on the skin

Warnings

Warnings For external use only

Flammable. Keep away from fire or flame or sparks

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 rash appears on the skin.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough to cover both hands in the palm, and rub hands together until dry.
- Children under 6 years of age should be supervised by adult when applying this product.

- Open cap. Place a small amount (enough to cover both hands) in the palm of your hand.
- Rub hands together, spreading the sanitizer and rubbing vigorously until dry

Other Information

- Store below 110 degrees F (43 degrees C)
- May discolor certain fabrics or surfaces.

Inactive Ingredient

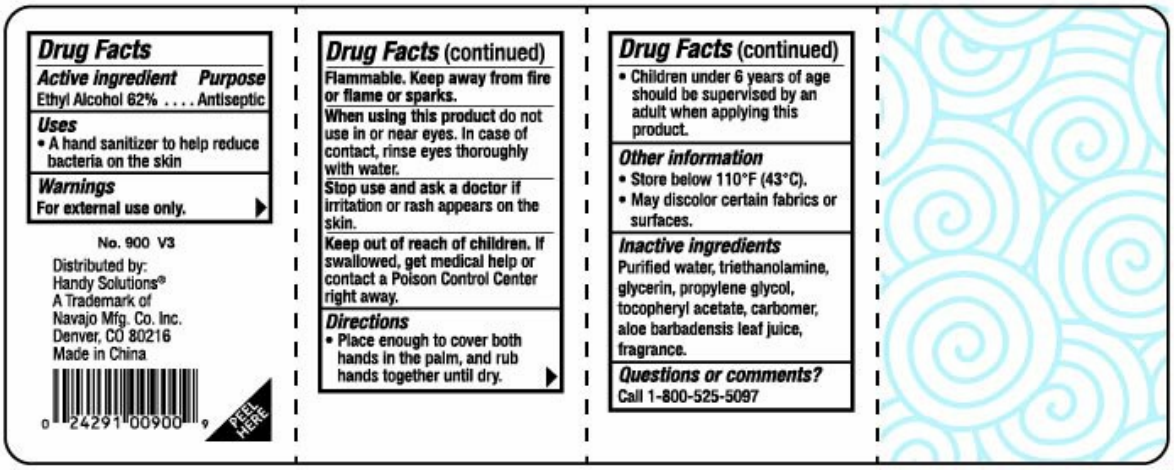
Purified water, triethanolamine, glycerin, isopropyl myristate, propylene glycol, tocopheryl acetate, carbomer, aloe barbadensis leaf juice, fragrance.

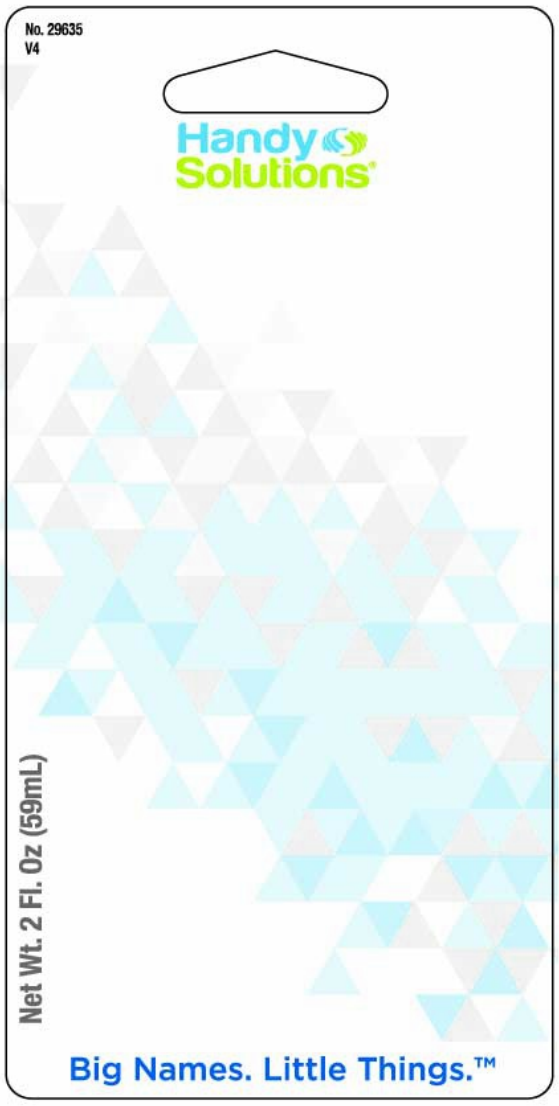
Questions or comments?

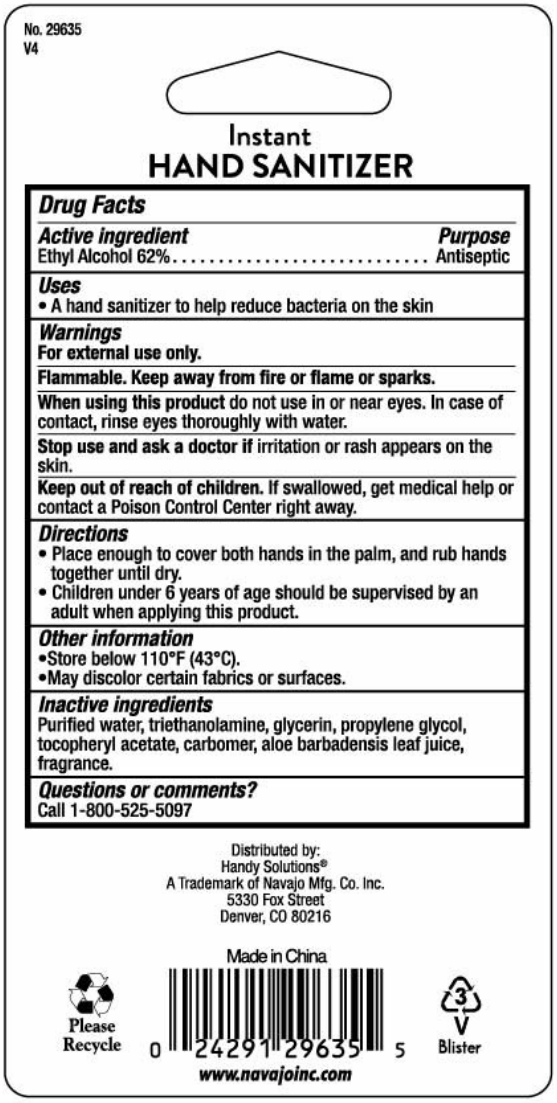
Call **toll-free 1-800-525-5097**

or visit is on the web at **www.navajomfg.com**

Package Labeling







HANDY SOLUTIONS INSTANT HAND SANITIZER WITH ALOE AND VITAMIN E

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67751-085
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67751-085-02	1 in 1 PACKAGE	10/28/2011	
1		59 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:67751-085-03	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/28/2011	
3	NDC:67751-085-04	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/28/2011	01/01/2016

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/2011	

Labeler - Navajo Manufacturing Company Inc. (091917799)

Establishment

Name	Address	ID/FEI	Business Operations
Navajo Manufacturing Company Inc.		136941411	repack(67751-085)

Establishment

Name	Address	ID/FEI	Business Operations
Nantong Health & Beyond Hygienic Products Inc.		421280161	manufacture(67751-085)

Revised: 5/2020 Navajo Manufacturing Company Inc.