HAND SANITIZER- 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 70%

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only.

Flammable. Keep away from fire or flame

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest.

Stop use and ask a doctor if

- irritation or redness develops
- 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

- 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

Do not store above 105°F

May discolor some fabrics

Harmful to wood finishes and plastics

Inactive ingredients

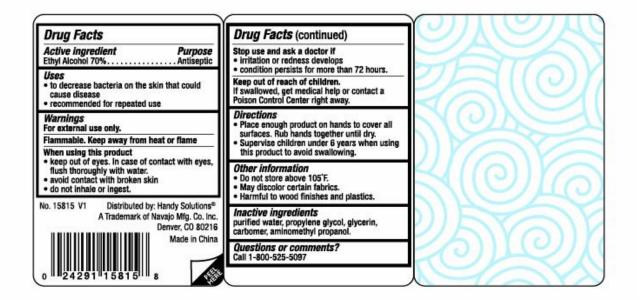
purified water, propylene glycol, glycerin, carbomer, aminomethyl propanol.

Questions or Comments?

Call 1-800-525-5097

Principal Display Panel





HAND SANITIZER			
ethyl alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67751-196

Route	of Administration	
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Active Ingredient/A	ctive Moiety			
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958	3V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL	
Inactive Ingredients				
macuve ingreutente			C two w of th	
	Ingredient Name		Strength	
WATER (UNII: 059QF0KC	,			
CARBOMER HOMOPOL	YMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
AMINOMETHYLPROPAN	NOL (UNII: LU49E6626Q)			
GLYCERIN (UNII: PDC6A)	3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:67751-196-01 59	mL in 1 BOTTLE; Type 0: Not a Combination Product	05/12/2020		
Marketing Infor	mation			
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
ZHEJIANG JINGHUI COSMETICS SHARE CO.,LTD		529558167	manufacture(67751-196)

Revised: 5/2020

Navajo Manufacturing Company Inc.