# HAND SANITIZER- alcohol liquid ningbo fareast industry co.,ltd

Reference Label Set Id: a68674ba-455a-4d63-e053-2a95a90aed00

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.

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to help reduce bacteria that potentially can cause disease.for use when soap and water are not available for external use only,flammable,keep away from heat or flame

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 thorough 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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place enough product on hands to cover all surface, rub hands together until dry supervise children under 6 years when using this product to avoid swallowing



Active ingredients

Purpose

Ethyl Alcohol 75%......Antiseptic

Uses ■ Hand sanitizer to help decrease bacteria on the skin When water, soap & towel are not available. Recommended for repeated use.

#### Warnings

- For external use only
  Flammable
- Keep away from fire or flame
- Do not apply around eyes. Do not use in ears & mouth.
- When using this product. avoid contact with eyes.
- In case of contact flush eyes with water
- Stop use and ask a doctor it redness or irritation develop and persist for more than 72 hours
- Keep out of reach of children. Children must be supervised in use of this product

#### Directions

■ Pump as needed into your palms and thoroughly spread on both hands. Rub into skin until dry

Other Information Store at 20C(88" to 77"F) may discolor fabrices

#### Inactive ingredients

Water, Glycerin, Carbomer, Triethanolamine, propylene





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made in china

20ML NDC 78623-016-20 30ML NDC 78623-016-30 60ML NDC 78623-016-60 75ML NDC 78623-016-75

#### HAND SANITIZER

alcohol liquid

Prod	luct	Info	uma	tion
Prou		11110	riiia	111111

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78623-016

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

**Ingredient Name Basis of Strength** Strength

ALCOHOL

75 mL in 100 mL

### **Inactive Ingredients**

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:78623-016- 20	20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/11/2021				
2	NDC:78623-016- 30	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/11/2021				
3	NDC:78623-016- 60	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/11/2021				
4	NDC:78623-016- 75	75 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/11/2021				

Marketing Information						
Marketing Category Application Number or Monograph Citation		Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	12/10/2020				

Labeler - ningbo fareast industry co.,ltd (554535808)

Revised: 1/2021 ningbo fareast industry co.,ltd