HAND SANITIZER- alcohol gel 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.

alcohol 75%v/v purpose : antibacterial

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

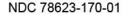
in children less than 2 months of age .on open skin wounds

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





Drug Facts

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
- 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, Glyerin, Vitamin E, Carbomer, Trolamine.



1000ml NDC 78623-275-01

HAND SANITIZER

alcohol gel

Product Information

HUMAN OTC DRUG NDC:78623-275 Item Code (Source) Product Type

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

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# It	em Code	Package Description	Marketing Start Date	Marketing End Date

1 NDC:78623-275-01 1000 mL in 1 BOTTLE; 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	12/31/2020	

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

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