

HAND SANITIZER WITH MOISTURIZERS- ethyl alcohol gel
China Ningbo Shangge Cosmetic Technology Corp.

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.

Active Ingredient

Purpose

Ethyl Alcohol 62.0%.....Antimicrobial

Uses

- for handwashing to decrease the bacteria on the skin
- recommended for repeated use

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

Other information

- store at 20oC to 25oC (68o to 77oF)
- may discolor certain fabrics

Warnings

For external use only.

Flammable, keep away from heat and flame.

Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.

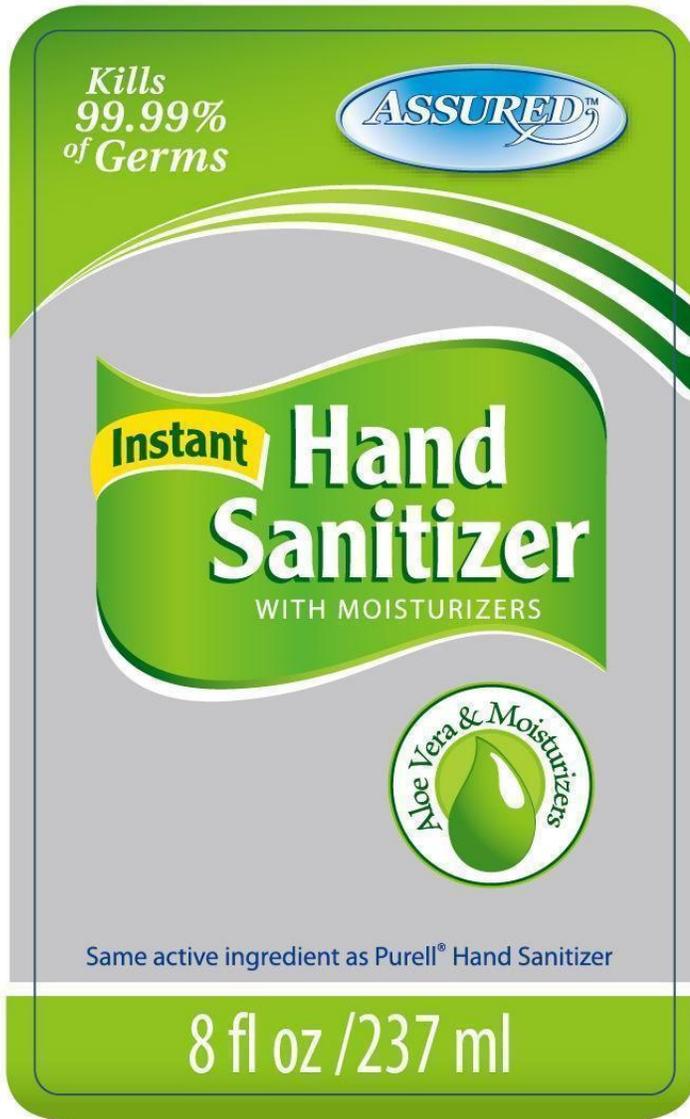
Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Directions

- wet hands thoroughly with product
- briskly rub hands together until dry
- supervise children under 6 years in the use of this product

Inactive Ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, fd&c blue no.1, fd&c red no.33.



Drug Facts

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*This product is not manufactured or distributed by Gojo Industries Inc, distributor of the registered trademark Purell®.

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DISTRIBUTED BY:
GREENBRIER INTERNATIONAL, INC.
500 VOLVO PARKWAY, CHESAPEAKE, VA 23320
MADE IN CHINA

HAND SANITIZER WITH MOISTURIZERS

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58503-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	147 mL in 237 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	

TROLAMINE (UNII: 903K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58503-005-01	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/24/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/24/2013	

Labeler - China Ningbo Shangge Cosmetic Technology Corp. (529287434)

Establishment

Name	Address	ID/FEI	Business Operations
China Ningbo Shangge Cosmetic Technology Corp.		529287434	manufacture(58503-005)

Revised: 11/2022

China Ningbo Shangge Cosmetic Technology Corp.