

HAND SANITIZER ANTIBACTERIAL WITH MOISTURIZERS AND VITAMIN E

1.8OZ- alcohol gel

Ningbo Liyuan Daily Chemical Products Co.,Ltd.

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

Ethyl Alcohol 62.0%

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product

avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if

irritation or redness develops.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center.

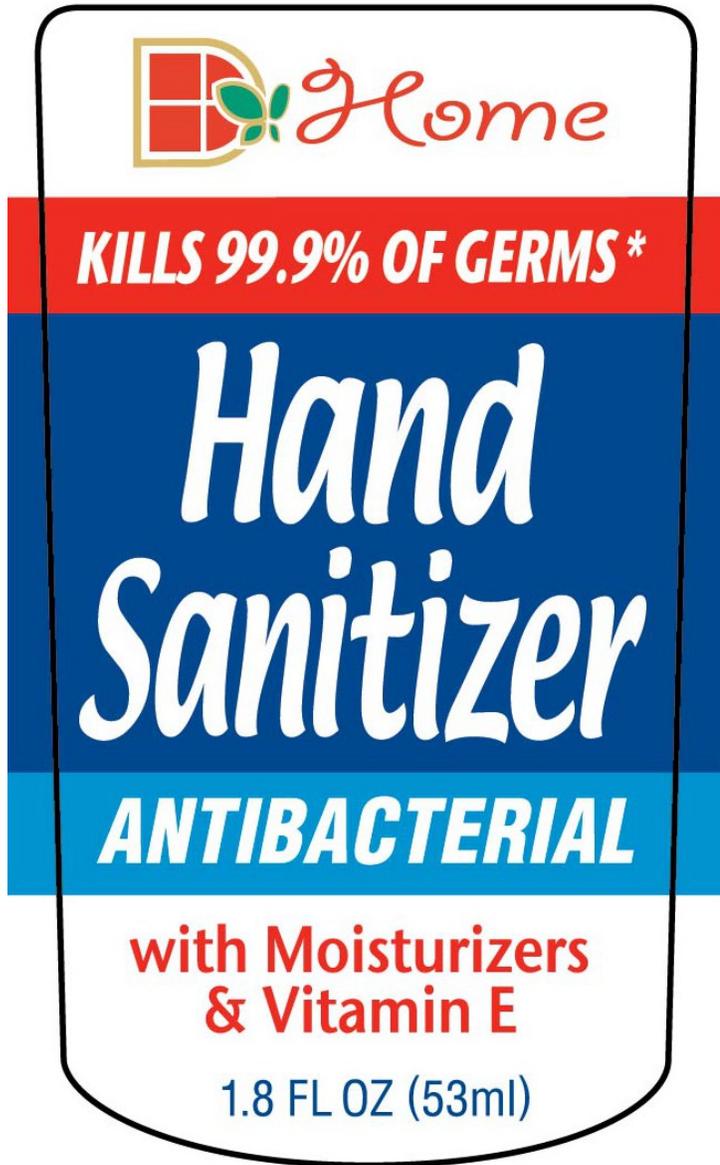
Directions

- wet hands thoroughly with product and rub into skin until dry.
- Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients

aloe barbadensis (aloe vera) leaf juice, carbomer, fragrance, glycerin, propylene glycol, tocopheryl acetate (vitamin E), triethanolamine, water (aqua)

Drug Facts



Home

KILLS 99.9% OF GERMS*

Hand Sanitizer

ANTIBACTERIAL

with Moisturizers & Vitamin E

1.8 FL OZ (53ml)



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Distributed by:
Trams Innovative Group
Miami USA
Made in China

HAND SANITIZER ANTIBACTERIAL WITH MOISTURIZERS AND VITAMIN E 1.8OZ

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76176-060
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)				
CARBOMER 934 (UNII: Z135WT9208)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
CUPRIC BIS(TRIETHANOLAMINE) (UNII: YBM44X0B6H)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-060-01	53 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	04/01/2018		

Labeler - Ningbo Liyuan Daily Chemical Products Co.,Ltd. (530766098)

Registrant - Ningbo Liyuan Daily Chemical Products Co.,Ltd. (530766098)

Establishment

Name	Address	ID/FEI	Business Operations
Ningbo Liyuan Daily Chemical Products Co.,Ltd.		530766098	manufacture(76176-060)

Revised: 10/2022

Ningbo Liyuan Daily Chemical Products Co.,Ltd.