

GRANDWAY HAND SANITIZER 1.8OZ- ethyl 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.

Drug Facts

Active ingredient

Ethyl Alcohol 62 percent

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:

Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Fragrance, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E)



kills 99.9% of germs

Hand Sanitizer

— DISINFECTION —

- VITAMIN E ADDED
- MOISTURIZE HAND SKIN

1.8 fl.oz/53mL

Drug Facts

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Distributed by:
GRANDWAY
USA
Made in China

GRANDWAY HAND SANITIZER 1.8OZ

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76 176-194
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

CARBOMER 934 (UNII: Z135WT9208)	
TRIETHANOLAMINE BENZOATE (UNII: M3EN4GC19W)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-194-01	53 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/01/2020	

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(76176-194)

Revised: 3/2020

NINGBO LIYUAN DAILY CHEMICAL PRODUCTS CO., LTD.