HAND SANITIZER- ethyl alcohol gel Jiangsu Ouvali Daily Cosmetic 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 75%

Purpose

Antiseptic

Uses

For hand washing to decrease bacteria on the skin.

Recommended for repeated use.

Warnings

Flammable. Keep away from fire or flame.

For external use only.

Avoid contact with broken skin.

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help.

Directions

Not recommended for infants.

Wet hands thoroughly with product and allow to dry without wiping.

Children under 6 years of age should be supervised when using this product.

Other Information

Do not store above 110°F. May discolor some fabrics.

Inactive Ingredients

Water, Aloe Barbadensis Leaf Juice, Maltodextrin, Glycerin, Propylene Glycol, Acrylates/C1O-30 Alkyl Acrylate crosspolymer, Triethanoamine, Fragrance, Tocopheryl Acetate.



Drug Facts

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Jiangsu Ouyali Daily Cosmetio Co., Ltd No. 9-9, Chuangye Road, Shatou Town, Guangling District, Yangshou oity, Jiangsu province, China Made in China



HAND SANITIZER

ethyl alcohol gel

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76719-002
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Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

Inactive Ingredients				
Ingredient Name	Strength			
MALTO DEXTRIN (UNII: 7CVR7L4A2D)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
GLYCERIN (UNII: PDC6A3C0OX)				
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)				
TROLAMINE (UNII: 9O3K93S3TK)				
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				

Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:76719-002-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/25/2020					
Marketing Information							
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not fin	nal part333E	04/25/2020					

Labeler - Jiangsu Ouyali Daily Cosmetic Co., Ltd. (542920070)

Revised: 4/2020 Jiangsu Ouyali Daily Cosmetic Co., Ltd.