HAND SANITIZER 59ML 01- alcohol liquid
Shenzhen Lantern Science 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.

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hand sanitizer 59ml

Drug Facts

Drug Facts Antibacterial Hand Sanitizer Gel

Active Ingredient **Purpose**

Ethyl Alcohol 75%(V/V) ..... Antibacterial

Use For handwashing to decrease bacteria on the skin.

Warnings Flammable.Keep away from fire or flame.

For external use only

When using this product avoid contact with eyes. if contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hours.

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

Directions

- Take a palmful (5grams) of product in one hand
- Spread on both hands and rub into skin

Other Information

- Do not store above 105°F
- May discolor some fabrics
- Harmful to wood finishes and plastics

Inactive Ingredients

Aqua (Water), Aminomethyl Propanol, Aloe Barbadensis Leaf Juice, Carbomer, FD&C Blue No.1, FD&C Yellow No.5, Glycerin, Isopropyl Alcohol, Isopropyl Myristate, Parfum (Fragrance), Propylene Glycol, Tocopheryl Acetate (Vitamin E).

2.0 fl oz (59ml) NDC 54860-175-01

VL2013 - #53138 - EXP: 043022
Logomark, Inc. ASI 67866
1201 Bell Avenue, Tustin, CA 92780, USA
Made In China

Active Ingredient

Active Ingredients Purpose

Ethyl Alcoh75% Antiseptic

USE

For handwashing to decrease bacteria on the skin.

Recommended for repeated use.

use anywhere without water.

Warning

flammable,keep away from fire or flame.

For external use only

When using this product,avoid contact with eyes.if contact occurs,rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash develops and continues for more than 72hours.

keep out of reach of children

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

For external use only.

Flammable, keep away from heat and flame.

Discontinue if skin becomes irritated and ask a doctor .

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a poson control center immediately.

Inactive ingredients

Alcohol□Aqua(water)□Isopropyl Alcohol□Glycerin□Carbomer□Aminomethyl Propanol□Parfum(Fragrance)□Propylene Glycol□ISOPROPYL MYRISTATE□ALOE BARBADENSIS LEAF JUICE□Tocopheryl Acetate(Vitamin E)□FD&C Yellow No.5,FD&C Blue No.1.

Directions

Take a palmful(5grams) of product in one hand

spread on both hands and rub into skin

When using this product

keep out of eyes. In case of contact with eyes, flush thoroughly with water.

Do not inhale or ingest.

Avoid contact with broken skin.

Other information

Do not store above 105F.

May discolor some fabrics.

Harmful to wood finishes and plastics.

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HAND SANITIZER 59ML 01

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-175
Route of Administration	TOPICAL		

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
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.1 mL in 100 mL
ALOE (UNII: V5VD430YW9)	0.2 mL in 100 mL
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	0.001 mL in 100 mL
CARBOMER 940 (UNII: 4Q93RCW27E)	0.4 mL in 100 mL
WATER (UNII: 059QF0KO0R)	23.057972 mL in 100 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302)	0.5 mL in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.00002 mL in 100 mL
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.14 mL in 100 mL
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	0.000008 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	0.5 mL in 100 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.1 mL in 100 mL
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.001 mL in 100 mL

Product Characteristics

Color	green	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-175-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/17/2020	

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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/17/2020	

Labeler - Shenzhen Lantern Science Co.,Ltd (421222423)

Registrant - Lantern Beauty America,INC. (117371139)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co., Ltd.		421222423	manufacture(54860-175)

Revised: 4/2020 Shenzhen Lantern Science Co.,Ltd