LE VITAL DREAMY GLOW- ethyl alcohol gel Lantern Enterprises 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.

Mecicical Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Features

To decrease bacteria on the skin that could cause disease.

- Recommended for repeated use.
- Use anywhere without water.

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Le Vital ® Signature

dreamy glow

hand sanitizer

Directions

- Wet hands thoroughly with product and rub until dry without wiping
- For children under 6, use only under adult supervision.
- Not recommended for infants

Warning

- For external use only hands
- Flammable, keep away from heat and flame
- Discontinue if skin becomes irritated and ask a doctor

CAUTION: For external use only. This product was not tested on animals.

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.

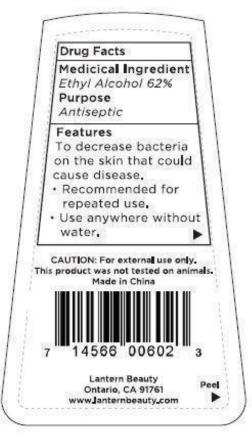
Other information

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

Non-Medicinal

Iwater, isopropyl alcohol, glycerin, carbomer, aminomethyl propanol, parfum, propylene glycol, isopropyl myristate, aloe barbadensis leaf juice, tacopheryl acetate





Drug Facts (continued)

Warning

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- 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 Poison Control Center immediately.

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Drug Facts (continued)

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Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:50154-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ı	Ingredient Name	Basis of Strength	Strength
ı	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 g

Inactive Ingredients

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
GLYCERIN (UNII: PDC6A3C0OX)		
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		

ı	Packaging				
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:50154-002-01	29 g in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Inform			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/23/2015	

Labeler - Lantern Enterprises Ltd. (004005196)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shanghai Lantern Cosmetic Co., Ltd.		421252043	manufacture(50154-002)	

Revised: 8/2015 Lantern Enterprises Ltd.