MELLOW INSTANT HAND SANITIZER WITH ALOE AND VITAMIN E - 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.

Instant Hand Sanitizer With Aloe and Vitamin E

Medicinal Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Non-Medicinal

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

Features

- To decrease bacteria on the skin that could cause disease.
- Recommended for repeated use.
- Use anywhere without water.

Directions

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

Warning

Warning
- For external use only - hands.
- 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 Poison Control Center immediately.

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 degrees F.
- May disscolor some fabrics.
- Harmful to wood finishes and plastics.

This product was not tested on animals.

Made in China

Used under Lic. and Dist. By:
LANTERN ENTERPRISES LTD.
Bellingham, WA 98225
1-888-273-7955
Web: LanternBeauty.com

MELLOW INSTANT HAND SANITIZER WITH ALOE AND VITAMIN E
ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50154-0030
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)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50154-0030-4	59 mL in 1 BOTTLE		
2	NDC:50154-0030-5	236 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/30/2009	

Labeler - Lantern Enterprises Ltd. (240084249)

Revised: 10/2011 Lantern Enterprises Ltd.