

MELLOW INSTANT HAND SANITIZER 8 OZ- ethyl alcohol gel

Ashtel Studios, Inc

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.

MELLOW INSTANT Hand Sanitizer

Drug Facts

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&C Yellow No. 5(Tartrazine), FD&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 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.

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.
- ◆ Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact

a Poison Control Center immediately.

Other information

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

With Aloe and Vitamin E

Kills 99.99% of Germs

This product was not tested on animals.

Used under Lic. & Dist.By:

LANTERN BEAUTY AMERICA INC

Address: 1401 E, Cedar Street, Unit C, Ontario, CA 91761, USA

TEL: 909-799-9700

Web:LanternBeauty.com

Made in China

Packaging

MELLOW®

INSTANT Hand Sanitizer
With Aloe and Vitamin E

Kills 99.99% of Germs

8 fl.oz/236 mL

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Medicinal Ingredient	Purpose
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♻️ Item: **ML 305** Made in China

MELLOW INSTANT HAND SANITIZER 8 OZ

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70108-033
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)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108-033-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/06/2020	

Labeler - Ashtel Studios, Inc (148689180)

Revised: 4/2020

Ashtel Studios, Inc