

HAND SANITIZER- ethyl alcohol gel

Allure Labs 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.

DRUG FACTS

Active Ingredients:

Ethyl Alcohol - 70%

Purpose: Antimicrobial

Uses: Helps to reduce bacteria on skin. Recommended for repeated use.

Warning: For External use only - Hands

Flammable: Keep away from heat and Flame.

When using this products:

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin.

Stop use and ask a doctor if skin irritation develops.

Keep out of reach of children:

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

Directions:

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

Inactive Ingredients: Water (Aqua), Acrylates Copolymer, Glycerin, Sodium Hydroxide.

MANUFACTURED FOR

ECZEMA HONEY CO BY,

ALLURE LABS,

HAYWARD, CA, 94545



**eczema
honey®**

HAND SANITIZER

EFFECTIVE & UNSCENTED

ECZEMAHONEYCO.COM
@ECZEMAHONEYCO

**8 FL OZ.
(237 ML)**

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Directions:

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Other Information:

- Do not store at temperature above 105 °F

Inactive Ingredients:

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ECZEMA HONEY CO BY
ALLURE LABS:

HAYWARD, CA, 94545
NDC 62742-4187-2

HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4187
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4187-1	59 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/23/2020	
2	NDC:62742-4187-2	237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/23/2020	

Labeler - Allure Labs Inc (926831603)

Revised: 4/2020

Allure Labs Inc