

EDDDIES HAND SANITIZER GEL- alcohol 75% gel
Nutralife Biosciences, 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.

Eddie's Gel Hand Sanitizer 8oz Nutralife

Active Ingredient

Alcohol 75%

Purpose

Antiseptic

Uses

To help reduce bacteria on the skin. For use when soap and water are not available

Warnings

For External Use only

Flammable. Keep away from heat or flame

Do not Use

On children less than 2 months of age- On open skin wounds -around eyes -in ears and mouth

When using this product

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water. - avoid contact with broken skin. - do not inhale or ingest

Stop use and ask a doctor if

Irritation or rash occurs

Keep out of reach of children

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

Directions

-Place enough product on hands to cover all surfaces. Rub hands together until dry. -Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

Store between 15-30C (59-86F). -Avoid freezing and excessive above 40C (104F)

Inactive Ingredients

Aloe Barbadensis (leaf) Extract, Carbomer, Fragrance, Glycerin, purified water

Eddie's Clean Hand Sanitizer

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Drug Facts

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Manufactured By: NUTRALIFE BIOSCIENCES Inc.
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Stock Symbol: NLBS | nutralifbiosciences.com

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EDDIES HAND SANITIZER GEL

alcohol 75% gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73761-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	177.75 mL in 237 mL

Inactive Ingredients

Ingredient Name	Strength
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ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73761-102-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/02/2020	

Labeler - Nutralife Biosciences, Inc (078834338)

Registrant - Nutralife Biosciences, Inc (078834338)

Revised: 3/2020

Nutralife Biosciences, Inc