EDDIES HAND SANITIZER- alcohol 75% spray
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 Spray 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, Glycerin, Fragrance, Purified water

Eddie's Clean Hands Hand Sanitizer

EDDIES HAND SANITIZER

alcohol 75% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73761-101
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**

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73761-101-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/02/2020	
2	NDC:73761-101-02	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/02/2020	04/01/2020
3	NDC:73761-101-60	60 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: 4/2020 Nutralife Biosciences, Inc.