THRIVE HEALTH HAND SANITIZER- ethyl alcohol gel
Sanrace Biotechnology Co., 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.

Thrive Health Hand Sanitizer with Aloe

Drug Facts

Active Ingredient

Ethyl Alcohol 75% v/v

Purpose

Antiseptic

Use[s]

Hand sanitizer to help reduce bacteria that potentially can cause disease.

For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame.

Do not use

- in children less than 2 months of age
- on open wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

- Avoid freezing, store below 43°C (110°F)

Inactive Ingredients: Water, Glycerin, Propylene glycol, Carbomer, Aloe vera extract

Packaging

1013 THRIVE TAPE ALCOHOL HAND SANITIZER LABEL - 100% SCALE
LABEL SIZE: 1.1"W X 1.97"H

COLOR LIST:
PMS 108 YELLOW
BLACK
WHITE

FRONT -
PRINTED DIRECTLY ON BOTTLE

BACK

THRIVE HEALTH HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75448-322
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75448-322-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:75448-322-02	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
3	NDC:75448-322-03	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/29/2020	

Labeler - Sanrace Biotechnology Co., Ltd. (543000938)

Establishment

Name	Address	ID/FEI	Business Operations
Sanrace Biotechnology Co., Ltd.		543000938	manufacture(75448-322)

Revised: 5/2020 Sanrace Biotechnology Co., Ltd.