HAND SANITIZER WITH ALOE AND VITAMIN E- ethyl alcohol liquid Top's Healthcare Product 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.

TANSOR INSTANT Hand Sanitizer With Aloe and Vitamin E

Antibacterial Hand Sanitizer

Active Ingredient

Ethyl Alcohol 71% (V/V)

Purpose

Antiseptic

Features:

To decrease bacteria on the skin. Recommended for repeated use. Delicate and fast dry. Use anywhere without water.

Inactive Ingredients:

Water (Aqua), Isopropyl Alcohol, Glycerin, Aloe Yohyju Matsu Ekisu, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine, Fragrance (Parfum).

Directions:

- Place enough product in your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.
- Children under 8 years of age should be supervised when using this product.

Warning:

- Flammable. Keep away from fire or flame.
- For external use only.
- When using this product, do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.
- Stop using and ask a doctor, if irritation or rash appears and lasts.
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Other information:

- Do not store above 110°F (43°C).
- May discolor certain fabrics or surfaces.

Kills 99.99% of Germs

Distributor: Tops Healthcare Product Inc.

2233 E 49th Street, Vernon 90058 CA, USA

E-mail: customerservice@topshealthcare.com Production Date: Please find on the package

Shelf life: 2 years

Executive Standard: GB 26373

Made in China

Packaging



TANSOR

INSTANT

Hand Sanitizer

With Aloe and Vitamin E

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Kills 99.99% of Germs 3.3 fl(oz)/300ml

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TANSOR

INSTANT Hand Sanitizer

With Aloe and Vitamin E

Kills 99.99% of Germs

3.3 fl(oz)/500ml

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HAND SANITIZER WITH ALOE AND VITAMIN E

ethyl alcohol liquid

Product Information	oduct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75227-151	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ISOPROPYL ALCOHOL (UNII: ND2M416302)		
GLYCERIN (UNII: PDC6A3C0OX)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)		
TROLAMINE (UNII: 9O3K93S3TK)		

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75227-151-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	
2	NDC:75227-151-03	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	
3	NDC:75227-151-05	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	
4	NDC:75227-151-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/26/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/26/2020		

Labeler - Top's Healthcare Product Inc (117482389)

Revised: 4/2020 Top's Healthcare Product Inc