HAND SANITIZER- alcohol gel TV Direct LLC

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.

Hand Sanitizer

Drug Facts

Active Ingredients

Ethyl Alcohol 75% v/v

Purpose

Antiseptic

Uses:

Hand Sanitizer to help reduce bacteria on the skin

Warnings:

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

Do not use

- in children less than 2 months of age
- on open skin 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. Rubs hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

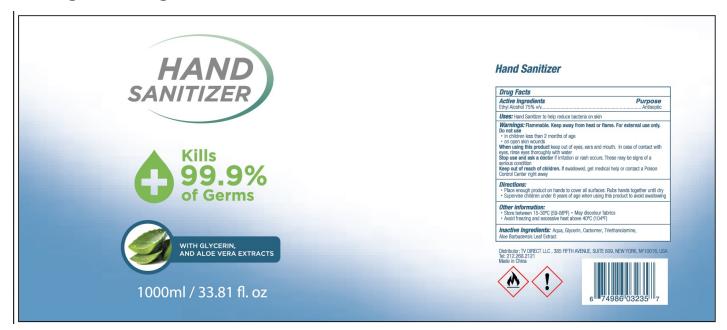
Other information:

- Store between 15-30°C (59-86°F)
- May discolor fabrics
- Avoid freezing and excessive heat above 40°C (104°F)

Inactive Ingredients:

Aqua, Glycerin, Carbomer, Triethanolamine, Aloe Barbadensis Leaf Extract

Package Labeling:1000ml



Package Labeling:60ml



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Distributor: TV DIRECT LLC , 385 FIFTH AVENUE, SUITE 809, NEW YORK, NY10016, USA Made in China I Tel: 212.268.2121



HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78915-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (LINII: 3K9958\/90M) (ALCOHOL - LINII: 3K9958\/90M)	ALCOHOL	0.75 ml in 1 ml

Inactive Ingredients

l	Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)

GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 903K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:78915-001-33	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020		
NDC:78915- 001-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/18/2020	

Labeler - TV Direct LLC (078829674)

Revised: 2/2023 TV Direct LLC