SNTZR- alcohol gel Applichem, 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.

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Medicinal ingredient: Ethyl alcohol 70% v/v

Purpose

Antiseptic

Use

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 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.

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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 heat above 40C (104F)

Inactive ingredients

Water (Aqua), Glycerin, Hydrogen Peroxide, Alkyl Acrylate Crosspolymer, Triethanolamine, Aloe, Lauryl Alcohol, Myristyl Alcohol, Fragrance (Parfum), PEG Dimethicone

Package Label - Principal Display Panel



3800 mL NDC: 74751-006-01

SNT	ZR
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alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74751-006
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety		
Ingredient Name		Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII: 3K9958 V90M)	ALCOHOL	70.6586 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	0.2665 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.3344 mL in 100 mL

HYDRO GEN PERO XIDE (UNII: BBX060 AN9 V)	0.1258 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	0.0216 mL in 100 mL
WEST INDIAN LEMONGRASS OIL (UNII: 5BIA40E9ED)	0.0187 mL in 100 mL
MYRISTYL LACTATE (UNII: 1D822OC34X)	0.2656 mL in 100 mL
TROLAMINE (UNII: 903K93S3TK)	0.6023 mL in 100 mL
METHACRYLIC ACID - ETHYL ACRYLATE COPOLYMER (4500 MPA.S) (UNII: T967IEU43C)	7.6152 mL in 100 mL

Packaging				
# It	em Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC 01	:74751-006-	3800 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/12/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/12/2020	

Labeler - Applichem, Inc (039829528)

Revised: 11/2020 Applichem, Inc