HAND SANITIZER- alcohol aerosol, spray Cosmetic Specialty Labs, 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) (75%, 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. Aloe Barbadenis Leaf Juice.
- c. Caprylyl Glycol.
- d. Glycerin.
- e. Phenoxyethanol.
- f. Polysorate-20.
- g. Purified Water.
- h. Tocopheryl Acetate.

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

SD Alcohol 40 75%

Antimicrobial

Use

Hand Sanitizer to help reduce bacteria on the skin.

Warnings

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 thoroughtly with water.

Stop use 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 Center right away.

Directions

- Use enough product in your palm to cover hands and rub hands together biskly until dry.
- Children under 6 years of age should be supervised when using this product.

Other information

- Store below 110F (43C)
- May discolor fabrics or surfaces

Inactive ingredients

Aloe Barbadensis Leaf Juice, Caprylyl Glycol, Fragrance, Glycerin, Phenoxyethanol, Polysorbate-20, Purified Water, Tocopheryl Acetate.

Package Label - Principal Display Panel

HAND SANITI	ZER				
alcohol aerosol, sp	ау				
Product Inform	ation				
Product Type		HUMAN OTC DRUG Item Code (Source)			NDC:58133-959
Route of Administr	ation	TOPICAL			
Active Ingredie	nt/Active Moiet	у			
	Ingredie	ent Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) AI				ALCOHOL	75 mL in 100 mL
Inactive Ingred	ents				
	Strength				
GLYCERIN (UNII: PDC6	(3C0OX)				
ALOE VERA LEAF (UN	: ZY81Z83H0X)				
POLYSORBATE 20 (UN	II: 7T1F30V5YH)				
PHENOXYETHANOL (U	NII: HIE492ZZ3T)				
WATER (UNII: 059QF0k	DOR)				
ALPHA-TOCOPHEROL	ACETATE (UNII: 9E8X	80D2L0)			
CAPRYLYL GLYCOL (U	NII: 00YIU5438U)				
Packaging					
# Item Code		Package Description		Marketing Start Date	Marketing End Date
1 NDC:58133-959-65	178 mL in 1 BOTTLE,	PUMP; Type 1: Convenience Kit of Co	-Package	08/30/2021	
Marketing In	ormation				
Marketing In Marketing Categ		tion Number or Monograph Cit	ation	Marketing Start Date	Marketing End Date

Labeler - Cosmetic Specialty Labs, Inc. (032973000)

Establishment						
Name	Address	ID/FEI	Business Operations			
Cosmetic Specialty Labs, Inc.		032973000	manufacture(58133-959) , label(58133-959) , pack(58133-959) , analysis(58133-959)			

Revised: 8/2021

Cosmetic Specialty Labs, Inc.