KEEP N IT CLEAN SANALL HAND SANITIZER FRESH SCENT- alcohol gel Sanall Enterprises, 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.

Keep n It Clean Sanall Hand Sanitizer Fresh Scent

Drug Facts

Active Ingredient[s]

Alcohol 70% v/v

Purpose

Antiseptic

Use:

- Hand sanitizer to help reduce bacteria on the skin that potentally can cause disease.
- For use when soap and water are not available.

Warnings:

- For exeternal use only.
- Flammable.
- Keep away from heat or flame.

Do not use

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

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 when using this product to avoid swallowing.

Other information

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

Inactive ingredients

acrylates copolymer, alpha-tocopherol acetate, denatonium benzoate, glycerin, isopropyl myristate, propylene glycol, tert-butyl alcohol, triisopropanolamine, fragrance.

Package Labeling:59ml



Package Labeling: 3785ml



KEEP N IT CLEAN SANALL HAND SANITIZER FRESH SCENT

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79537-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

retive ingredient/retive violety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL	

Inactive Ingredients				
Ingredient Name	Strength			
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				
DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2)				
GLYCERIN (UNII: PDC6A3C0OX)				
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)				
TRIISOPROPANOLAMINE (UNII: W9 EN9 DLM9 8)				
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)				

	Packaging					
:	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:79537-001-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2020			
	NDC:79537-001-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/15/2020			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/15/2020	

Labeler - Sanall Enterprises, LLC (117566912)

Revised: 8/2020 Sanall Enterprises, LLC