SANI-KURE EXTRA STRENGTH HAND SANITIZER- alcohol solution KOLORCURE CORPORATION

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.

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antimicrobial

Use

Use:

Antimicrobial

Warnings

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

Do not use

in or around the eys.

In case of contact immediately flush eyes with water.

If swallowed, contact physician.

Keep out of reach of children.

Directions

- Place enough product in your palm to cover hands. Rub hands together until dry.
- Children under 6 years of age should be supervised when using Sani-Kure.

Other information

Do not store above 110°F.

Inactive ingredients

Pure Aloe Vera, vegetable glycerin, isopropyl alcohol, hydrogen peroxide, distilled water.

Warnings

Keep out of reach of children.

Package Label - Principal Display Panel

2 FL OZ (59 mL)



Drug Facts

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Purpose Active Ingredient Ethyl alcohol 70% v/v. Antomicrobial Use: Help reduce bacteria on skn

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Directions: Spray enough product in your pains to cover hands and ne hands together briskly until day "Children under 6 years of age sholl be supervised when using Sani-Kun hand.

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SANI-KURE EXTRA STRENGTH HAND SANITIZER

alcohol solution

Product Information								
Product Type	HUMAN OTC DRUG	ltem C	ode (Source)	NDC:78923-001				
Route of Administration	TOPICAL							
Active Ingredient/Active Moiety								
Ingredient Name			Basis of Strength	Strength				
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)							

In	active Ingre	edients						
Ingredient Name					Strength			
GLYCERIN (UNII: PDC6A3C0OX)				2.1 mL in 100 mL				
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				1.8 mL in 100 mL				
WATER (UNII: 059QF0KO0R)			7.1 mL in 100 mL					
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)				15 mL in 100 mL				
Packaging								
#	Item Code	Package Description		ing Start ate	Marketing End Date			
1	NDC:78923- 001-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/30/202	D				
Marketing Information								
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date		Marketing End Date			
OT fin	C monograph no al	part333A	03/30/2020					

Labeler - KOLORCURE CORPORATION (113087894)

Revised: 1/2022

KOLORCURE CORPORATION