HAND SANITIZER- is opropyl alcohol gel Greensys of North America 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 ingredient

Isopropyl alcohol 70% v/v

Purpose

Antiseptic

Uses

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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.

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

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Labeling:

Alcohol Antiseptic 70% Topical Solution Hand Sanitizer Non-sterile Solution 250 MI (8.5 FI Oz.)

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Produced by: Greensys of North America LLC 5001 W. Military Hwy suite 100 McAllen, Tx 78503 1(956) 609-2888

MADE IN USA

HAND SANITIZER

isopropyl alcohol gel

P	roc	luct	Inf	orma	tion

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:76692-000

Route of Administration TOPICAL

Activo	Ingro	diant/	Activo	Moiety
ACTIVE	111914		ACTIVE	VIII

Ingredient Name	Basis of Strength	Strength
	ISOPROPYL ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	

l	Packaging					
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:76692-000-85	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	04/24/2020		

Labeler - Greensys of North America LLC (075577393)

Revised: 4/2020 Greensys of North America LLC