ISOPROPYL ALCOHOL ANTISEPTIC- alcohol solution Hill Dermaceuticals, 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.

Isopropyl Alcohol Antiseptic

Drug Facts

Active Ingredient[s]

Isopropyl alcohol 75% v/v

Purpose

Antiseptic

Use[s]

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.

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

glycerin, hydrogen peroxide, purified water USP

Question?

PRINCIPAL DISPLAY PANEL - 100 mL Bottle Label

HILL DERMACEUTICALS, INC.® Established In 1943

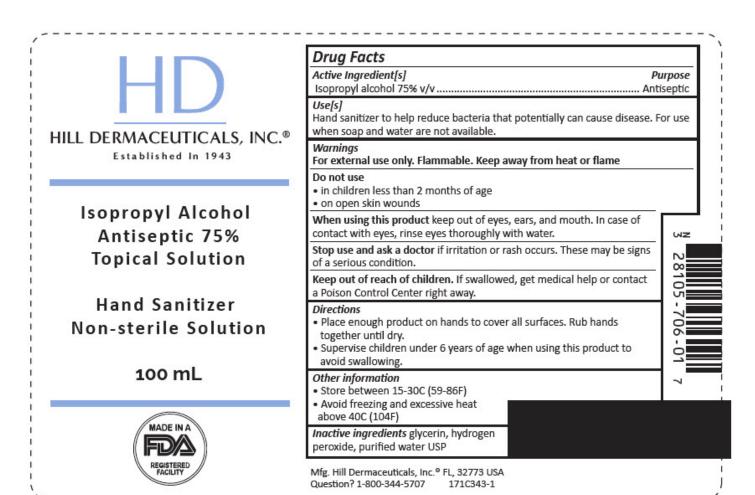
Isopropyl Alcohol Antiseptic 75% Topical Solution

Hand Sanitizer Non-sterile Solution

100 mL

MADE IN A FDA REGISTERED FACILITY

Product Type



ISOPROPYL ALCOHOL ANTISEPTIC alcohol solution Product Information

Item Code (Source)

NDC:28105-706

HUMAN OTC DRUG

Route of Administration	TOPICAL

Active	Ingred	dient/Active	Moiety
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ı	8				
	Ingredient Name		Basis of Strength	Strength	
	Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958	V90M)	Alcohol	75 mL in 100 mL	

Inactive Ingredients

mactive ingredients				
Ingredient Name	Strength			
glycerin (UNII: PDC6A3C0OX)				
hydrogen peroxide (UNII: BBX060AN9V)				
water (UNII: 059QF0KO0R)				

Packaging

1					
ı	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:28105-706-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	04/01/2020	

Labeler - Hill Dermaceuticals, Inc. (098366990)

Establishment			
Name	Address	ID/FEI	Business Operations
Hill Dermaceuticals, Inc.		098366990	MANUFACTURE(28105-706)

Hill Dermaceuticals, Inc. Revised: 4/2020