

MEDICAL SANITIZER- alcohol gel
Cedar Medical 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.

Medical[®] SANITIZER

Drug Facts

Active Ingredient(s)

Alcohol 70% v/v

Purpose

Antiseptic

Use(s)

- 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

- On children less than 2 months of age
- On open skin wound

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation occurs. These may be signs of a serious disease.

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 the product to avoid swallowing

Other information

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

Inactive ingredients fragrance, glycerin, *hydroxypropyl cellulose, *hydroxypropyl methylcellulose, water

***May or may not include**

UNSCENTED HAND GEL
KILLS 99.9% OF GERMS
WHO FORMULA
MADE IN THE USA
EPA APPROVED FACILITY
Distributed By:
Cedar Medical LLC
Tampa, FL, USA

Packaging



Medical
SANITIZER

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Medica?
SANITIZER

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70% ALCOHOL
KILLS 99.9% OF GERMS





4oz(118 ml)



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8oz(236 ml)



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16oz(473 ml)



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32oz(946 ml)

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MEDICAL SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80101-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80101-102-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
2	NDC:80101-102-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	

3	NDC:80101-102-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
4	NDC:80101-102-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
5	NDC:80101-102-32	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	
6	NDC:80101-102-10	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/18/2020	

Labeler - Cedar Medical Llc (117581534)

Revised: 8/2020

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