

SANI EZE- isopropyl alcohol spray
MENPER DISTRIBUTORS, 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.

Active Ingredient	Purpose
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Isopropyl alcohol 72% v/v.....	Antiseptic
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Purpose

Antiseptic

Use

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 fire or flame.

Do not use

in the eyes. In case of contact, rinse eyes thoroughly with water.

in children less than two months of age

on open skin wounds

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

wet hands thoroughly with product

briskly rub hands together until dry

supervise children under 6 years of age when using this product to avoid swallowing.

Other information

store at 20° to 25°C (68° to 77°F) Avoid excessive heat and protect from freezing.

may discolor certain fabrics

Inactive ingredients

Aloe barbadensis leaf juice, aminomethyl propanol, carbomer, dl-alpha tocopherol acetate, fragrance, glycerin, isopropyl myristate, propylene glycol, water.

Questions or Comments?

Call 1-800-560-5223 M-F 9 AM TO 4 PM Eastern.

Drug Facts

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**INSTANT HAND
SANITIZER**

**Kills 99.99% of
Germs**

2 Fl Oz (60mL)



Distributed by/Distribuido por:
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 11421 N.W. 107 Street Suite # 24
 Miami, FL 33178
 www.menperdistributors.com



0 42279 59349 4

SANI EZE

isopropyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53145-415
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	72 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE (UNII: V5VD430YW9)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53145-415-02	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/29/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/29/2011	

Labeler - MENPER DISTRIBUTORS, INC. (101947166)

Revised: 4/2021

MENPER DISTRIBUTORS, INC.