

SANI-CARDZ- alcohol solution
TVG Products, 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.

Sani-Cardz Hand Sanitizer

Drug Facts

Active ingredient

Alcohol 80% v/v

Purpose

Antiseptic

Uses

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

Inactive ingredients glycerin, hydrogen peroxide, purified water USP

MADE IN USA OF US & IMPORTED INGREDIENTS

Distributed by Sani-Cardz

2045 Biscayne Blvd #341

Miami, FL 33137

To report a serious adverse event contact 1-877-214-3305

ALCOHOL ANTISEPTIC 80%

Sani-Cardz
ON-THE-GO HAND SANITIZER
TOPICAL SOLUTION
NON-STERILE SOLUTION
0.7 FL. OZ. (20ML)



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SANI-CARDZ

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80805-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80805-001-01	20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/07/2020	
2	NDC:80805-001-12	12 in 1 PACKAGE	10/07/2020	
2		20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/07/2020	

Labeler - TVG Products, LLC (117689793)