

SANI-CHOICE INSTANT HAND SANITIZER- ethyl alcohol gel
SANI-CHOICE CORPORATION

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-CHOICE INSTANT HAND SANITIZER GEL

Drug Facts

Active ingredient

Ethyl Alcohol- 70%

Purpose

Antiseptic

Use- To Sanitize hands without water or rinse

Warnings

For external use only: hands

Flammable, keep away from fire or flame.

■ Keep out of eyes. ■ In case of contact with eyes, flush thoroughly with water ■ Do not inhale or ingest.

If ingested, seek medical help or contact a Poison Control Center right away.

Discontinue if skin irritation or redness occurs

■ Keep away from children

Directions

Apply Gel to hands; Rub thoroughly; Allow to dry without wiping or rinsing. Store at room temperature

Inactive Ingredients

Water, Acrylic Copolymer, Glycerin, Fragrance, Sodium Hydroxide

FOR ANTISEPTIC USE

Distributed By

Sani- Choice Corporation

4000 West 6th St. Suite B-208

Lawrence, Kansas 66049

MADE IN USA

Packaging

SANI-CHOICE!

INSTANT HAND SANITIZER GEL

FOR ANTISEPTIC USE

16 fl oz (473 mL)

Drug Facts

Active ingredient	Purpose
Ethyl Alcohol- 70%.....	Antiseptic

Use- To Sanitize hands without water or rinse

Warnings

For external use only: hands

Flammable, keep away from fire or flame.

■ Keep away from children. ■ Keep out of eyes. ■ In case of contact with eyes, flush thoroughly with water. ■ Do not inhale or inject. If injected, seek medical help or contact a Poison Control Center right away.

Discontinue if skin irritation or redness occurs

Directions

Apply Gel to hands; Rub thoroughly; Allow to dry without wiping or rinsing. Store at room temperature

Inactive Ingredients

Water, Acrylic Copolymer, Glycerin, Fragrance, Sodium Hydroxide

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Expires 09/2023

SANI-CHOICE INSTANT HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80412-070
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
WATER (UNII: 059QF0KO0R)	
ACRYLIC ACID/2-ETHYLHEXYL ACRYLATE/STYRENE COPOLYMER (500000 MW) (UNII: 3K8D4YRM4R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80412-070-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/04/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/04/2020	

Labeler - SANI-CHOICE CORPORATION (117632085)

Establishment

Name	Address	ID/FEI	Business Operations
SANI-CHOICE CORPORATION		117632085	manufacture(80412-070)

Revised: 9/2020

SANI-CHOICE CORPORATION