

KERICURE HAND SANITIZER- alcohol spray
Kericure Inc.

KeriCure® HAND SANITIZER

Drug Facts

Active Ingredient

Alcohol 70% (v/v)

Purpose

Antimicrobial

Uses • to decrease bacteria on the skin that could cause disease • recommended for repeated use

Warnings

For external use only • Flammable, keep away from heat or flame • **Do not use** on children less than 2 month 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 • Do NOT ingest

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

Directions • Shake bottle before use • Spray enough product on hands to cover all surfaces • Rub hands together until dry • Adult supervision

Other Information • Do not store bottle above 105°F • May discolor some fabrics • harmful to some wood finishes

Inactive Ingredients • Water, isopropyl alcohol, aloe barbadensis, glycerin, tocophoryl acetate, cellulose

MEDICAL

Made in the USA

KILLS MORE THAN 99.9% OF GERMS

FORMULATED FOR PROFESSIONALS
GENTLE ENOUGH FOR EVERYDAY USE

WITH ALOE & VITAMIN E
SKIN MOISTURIZERS

KeriCureMedical.com

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Distributed by KeriCure Inc 3701A Southwestern Blvd. Baltimore MD 21229

Packaging



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Distributed by KeriCure, Inc • KeriCureMedical.com
3701A Southwestern Blvd. Baltimore MD 21229 2oz (55mL)

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8oz (230mL)

Drug Facts

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KERICURE HAND SANITIZER

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78280-303
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)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
HYDROXYETHYL CELLULOSE (2000 MPA.S AT 1%) (UNII: S38J6RZN16)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78280-303-02	55 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/10/2020	
2	NDC:78280-303-08	230 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/10/2020	01/01/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/10/2020	

Labeler - Kericure Inc. (966638681)

Revised: 1/2025

Kericure Inc.