

KERICURE HAND SANITIZER- alcohol spray

Kericure 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.

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

• Keep out of reach of children • Do NOT ingest • 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, natural fragrances, tocophoryl acetate, cellulose

■ LAVENDER ■ PEPPERMINT

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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Packaging



HAND SANITIZER

Made in the USA

KILLS OVER 99.9% OF GERMS IN 20 SECONDS

WITH ALOE & VITAMIN E SKIN MOISTURIZERS

*FORMULATED FOR PROFESSIONALS
GENTLE ENOUGH FOR EVERYDAY USE*

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

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8oz (230mL)

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KERICURE HAND SANITIZER

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78280-304
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 MPAS AT 1%) (UNII: S38J6RZN16)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78280-304-02	55 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/10/2020	
2	NDC:78280-304-08	230 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/10/2020	

Labeler - Kericure Inc. (966638681)

Revised: 6/2020

Kericure Inc.