

SANIX ALCOHOL HAND RUB- alcohol liquid
Topiderm, Inc

Sanix Alcohol Hand Rub

Drug Facts

Active ingredients

Ethyl Alcohol 62% (v/v)

Purpose

Antimicrobial

Uses

Hand sanitizer for reduction of bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from fire and flame.

Keep out of and away from the eyes. In case of eye contact flush thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. Not for use on infants.

If product is swallowed, get medical help or contact a Poison Control Center immediately.

Directions

Apply enough product into the palm and rub hands together to thoroughly wet them

- Continue until the product dries without wiping off
- Supervise children when they are using the product
- Store below 110°F (43°C)
- May discolor some types of fabrics & surfaces.

Inactive ingredients

Water, Aloe Barbadosensis Leaf Juice, Glycerin, Tocopheryl Acetate, Polysorbate 20, Isopropyl Alcohol, Isopropyl Palmitate, Tetrahydroxypropyl Ethylenediamine, Carbomer, Fragrance.


PRINCIPAL DISPLAY PANEL - 237 ml Bottle Label

ANTIBACTERIAL
Hand Sanitizer Gel
by REPLENIX®

Kills 99.9% of Most Illness Causing Germs*
Triple Action Moisturizers
Softens Hands with Aloe, Vitamin E & Glycerin

Made in USA
Trusted By
MEDICAL
PROFESSIONALS
Manufactured by Topix Pharmaceuticals


Net 8 fl.oz. (237ml)



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SANIX ALCOHOL HAND RUB

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51326-900
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	
EDETOL (UNII: Q4R969U9FR)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51326-900-08	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M003	03/17/2020	

Labeler - Topiderm, Inc (049121643)

Registrant - Topiderm, Inc. (049121643)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc		049121643	MANUFACTURE(51326-900)

Revised: 3/2020

Topiderm, Inc