

DERMACEN LOTIONIZED HAND SANITIZER- alcohol liquid
Central Solutions Inc

Drug Facts 62654-151

Active Ingredient

Ethanol 70%

Purpose

Antiseptic

Uses

- An effective antimicrobial hand cleanser for topical application
- Recommended for repeated use
- Reduces transient microorganisms on intact skin
- Keeps hands from drying out due to repeated uses

Warnings

- **For External Use Only.**
- **Flammable, keep away from fire or open flame**

When using this product:

- Avoid contact with eyes
- In case of eye contact, flush with water for 15 minutes
- Discontinue use and see a doctor if irritation occurs
- Avoid contact with broken skin

Keep out of reach of children.

Directions

- Wet hands thoroughly with product. Allow to dry without wiping. Use no water or towels

Other information

- Not tested on animals

Inactive Ingredients

Aloe Barbadensis Leaf Juice, Ammonium Acryloyldimethyltaurate/VP Copolymer, Butylene Glycol, Calcium Pantothenate (Vitamin B5), Cetyl Alcohol, Fragrance, Glycerin, Glyceryl Laurate, Hydroxypropyl Cellulose, Isopropyl Myristate, Maltodextrin, Niacinamide

(Vitamin B3), Pyridoxine HCl (Vitamin B6), Silica, Sodium Ascorbyl Phosphate (Vitamin C), Sodium Starch Octenylsuccinate, Titanium Dioxide, Tocopherol (Vitamin E), Tocopheryl Acetate (Vitamin E), Trehalose, Water

Questions?

Call 800-255-0262

NDC: 62654-151-64

DermaCen

Lotionized

Hand

Sanitizer

For hand washing

to decrease bacteria

on the skin

Net Contents:

8 oz (236mL)

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MADE IN USA

Re-Order No:
14014
SDA-KS-664 14014REV0324

Central Solutions, Inc.
Kansas City, KS 66115
www.centralsolutions.com
(800) 255-0262

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DERMACEN LOTIONIZED HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62654-151
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
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TOCOPHEROL (UNII: R0ZB2556P8)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
CALCIUM PANTOTHENATE (UNII: 568ET80C3D)	
OCTENYLSUCCINIC ACID (UNII: 12UZE4X73L)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
TREHALOSE (UNII: B8WCK70T7I)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL LAURATE (UNII: Y98611C087)	
NIACINAMIDE (UNII: 25X51I8RD4)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62654-151-64	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/07/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/07/2024	

Labeler - Central Solutions Inc (007118524)

Registrant - Central Solutions (007118524)

Establishment

Name	Address	ID/FEI	Business Operations
Central Solutions Inc		007118524	manufacture(62654-151)

Revised: 8/2024

Central Solutions Inc