

**DERMACEN INSTANT HAND SANITIZER WITH ALOE- alcohol liquid**  
**Central Solutions Inc**

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**Drug Facts 62654-143**

Active Ingredients

Ethanol 64.5% (w/w)

Purpose

Hand Sanitizer

Uses

An antiseptic hand sanitizer for topical application

Prevents infection and cross contamination

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 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.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Wet hands thoroughly with product.

Allow to dry without wiping.

Use no water or towels.

Other Information

DermaCen products are not tested on animals

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice, EDTA, Fragrance, Glycerin, Tetrahydroxypropyl Ethylenediamine, Tocopherol (Vitamin E), Water, Yellow 5, Blue 1

DermaCen

Instant

Hand Sanitizer

with Aloe

Contains

Vitamin E

For hand washing

to decrease bacteria

on the skin

Net Contents

2.5 ounces (74 ml)

**DERMACEN INSTANT HAND SANITIZER WITH ALOE**

alcohol liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:62654-143
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	64.5 mL in 100 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)</b> (UNII: 59TL3WG5CO)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>EDETIC ACID</b> (UNII: 9G34HU7RV0)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>EDETOL</b> (UNII: Q4R969U9FR)	
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	

**Packaging**

#	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:62654-143-19	74 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
2	NDC:62654-143-12	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
3	NDC:62654-143-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
4	NDC:62654-143-02	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
5	NDC:62654-143-03	443 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
6	NDC:62654-143-92	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
7	NDC:62654-143-37	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2001	
8	NDC:62654-143-04	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2020	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	505G(a)(3)	01/01/2001	

**Labeler** - Central Solutions Inc (007118524)

**Establishment**

Name	Address	ID/FEI	Business Operations
Central Solutions Inc		007118524	manufacture(62654-143)

Revised: 12/2024

Central Solutions Inc