DERMACEN NON-ALCOHOL FOAMING HAND SANITIZER- benzalkonium chloride solution
Central Solutions 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.

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Uses

For hand sanitizing to decrease bacteria on the skin

Recommended for repeated use

Warnings

For external use only

When using this product avoid contact with eyes.

In case of eye contact, flush with water.

Stop use and ask a doctor if irritation or redness develops, or if conditions persists for more than 72 hours.

Keep out of reach of children. If swallowed, call a physician or Poison Control Center right away.

Directions

Pump a small amount of foam into palm of hand

Rub thoroughly over all surfaces of both hands

Rub hands together briskly until dry

Inactive ingredients

Purified Water, Glycerin, Cocamidopropyl Betaine, Triethanolamine

Purpose

Hand Sanitizer

DermaCen

Non-Alcohol

Foaming Instant

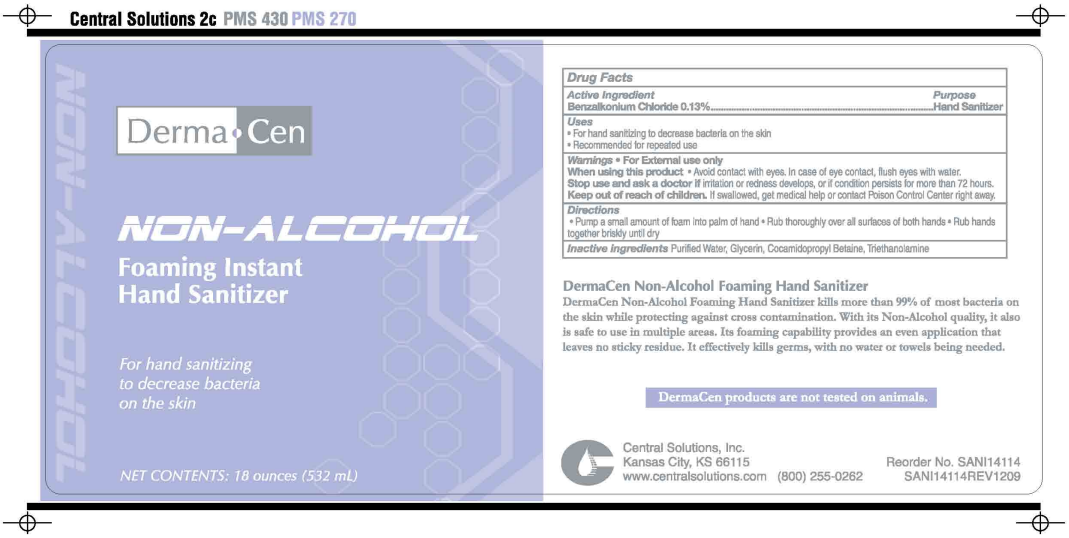
Hand Sanitizer

For hand sanitizing

to decrease bacteria

on the skin

NET CONTENTS: 18 ounces (532 mL)



DERMACEN NON-ALCOHOL FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62654-141
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62654-141-14	532 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/18/2010	
2	NDC:62654-141-16	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/18/2010	
3	NDC:62654-141-37	532 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/18/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/18/2010	

Labeler - Central Solutions Inc (007118524)

Establishment

Name	Address	ID/FEI	Business Operations
Central Solutions Inc		007118524	manufacture(62654-141)

Revised: 1/2023 Central Solutions Inc