

SANI-SURE AF- benzalkonium chloride solution
Whisk Products, Inc.

Sani-Sure AF

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

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 eyes with water.

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

Keep out of reach of children. If swallowed, get medical help or contact a 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

Water, Glycerin, Propylene Glycol, Phenoxyethanol, Lauramine Oxide, Panthenol, Fragrance, Citric Acid

Sani-Sure AF

**Foaming Instant Hand Sanitizer
Alcohol Free**

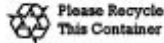
Contact To Reorder:



CHATTACHEM
423.322.3921

Drug Facts	
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LOT #: EXP:
 Contains: Four Bags
 Net Contents Each: 33.8 fl. oz. (1 qt. 1.8 fl. oz.) 1000 mL
 Total Net Volume: 135.2 fl. oz. (1 gal. 7.2 fl. oz.) 4 L



Manufactured by:
 Whisk Products, Inc.
 Wentzville, MO 63385

1004-1.0

SANI-SURE AF

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65585-360
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PANTHENOL (UNII: WW9CM0O67Z)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Product Characteristics

Color	white (water white - colorless, dispensed as a white foam)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65585-360-01	4 in 1 BAG	09/04/2024	
1		1000 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/04/2024	

Labeler - Whisk Products, Inc. (834270639)

Establishment

Name	Address	ID/FEI	Business Operations
Whisk Products, Inc.		834270639	manufacture(65585-360)

Revised: 9/2024

Whisk Products, Inc.