SANI LUXE FOAMING HAND SANITIZER ALCOHOL FREE- benzalkonium chloride liquid
Celeste Industries Corporation

SaniLuxe™ Foaming Hand Sanitizer Alcohol Free

Drug Facts

Active ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses

- Recommended for repeated use
- For hand sanitizing to decrease bacteria on the skin

Warnings

- **For external use only.**
- **Keep out of reach of children.**
- **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.
- 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 hands together briskly until dry
- Rub thoroughly over all surfaces of both hands

Inactive ingredients

Water (Aqua), Triethylene Glycol, Glycereth-26, Cocamidopropyl PG-Dimonium Cloride Phosphate,Hydroxyethylcellulose, DMDM Hydantoin, Triethanolamine, Fragrance (Aroma), Iodopropynyl Butyl Carbamate, 1, 3-Butanediol

SaniLuxe™

FOAMING HAND SANITIZER

Alcohol free

- Requires no water or towels.
- Apply small amount to hands

  and rub until dry.

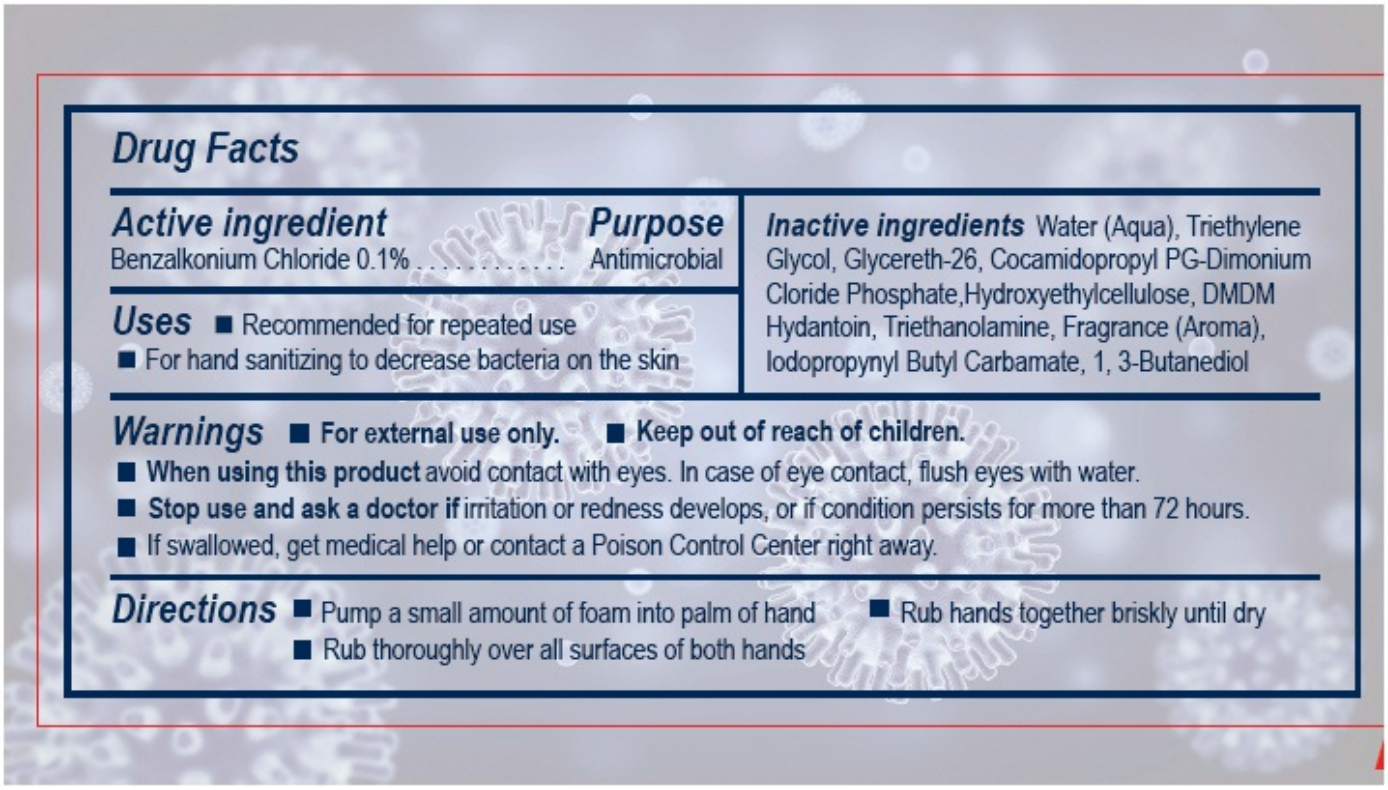
Kills 99%

of germs

on contact

Caution: Use on hands only.

8 fl oz • 236ml e



SANI LUXE FOAMING HAND SANITIZER ALCOHOL FREE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71489-006
Route of Administration	CUTANEOUS		

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)	
TRIETHYLENE GLYCOL (UNII: 3P5SU53360)	
GLYCERETH-26 (UNII: NNE56F2N14)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
HYDROXYETHYL CELLULOSE (5000 MPA.S AT 1%) (UNII: X70SE62ZAR)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
TROLAMINE (UNII: 9O3K93S3TK)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71489-006-01	237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/28/2019	
2	NDC:71489-006-02	56 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/25/2020	
3	NDC:71489-006-03	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/20/2020	
4	NDC:71489-006-04	284 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	
5	NDC:71489-006-05	355 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	
6	NDC:71489-006-06	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/20/2020	
7	NDC:71489-006-07	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
8	NDC:71489-006-08	177 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/25/2020	
9	NDC:71489-006-09	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
10	NDC:71489-006-10	284 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
11	NDC:71489-006-11	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
12	NDC:71489-006-12	3785 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/25/2020	
13	NDC:71489-006-13	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/30/2020	
14	NDC:71489-006-14	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/30/2020	
15	NDC:71489-006-15	530 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/14/2020	
16	NDC:71489-006-16	530 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/28/2019	

Labeler - Celeste Industries Corporation (047795034)

Revised: 7/2024 Celeste Industries Corporation