

SANI LUXE HAND SANITIZER WITH ALCOHOL- alcohol gel
Celeste Industries Corporation

Sani Luxe[®] Hand Sanitizer Gel with Alcohol

Drug Facts

Active ingredient

Ethyl Alcohol 62.5%

Purpose

Antimicrobial

Uses

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

Warnings

- **For external use only.**
- **Flammable. Keep away from open flame and sources of heat.**
- **Stop use and ask a doctor if** irritation or redness develops, or if condition persists for more than 72 hours.
- **When using this product,** avoid contact with eyes. In case of eye contact, flush eyes with water.
- **If swallowed,** get medical help or contact a Poison Control Center right away.
- **Keep out of reach of children.**

Directions

- Children under 6 years of age should be supervised when using this product
- Place enough product in your palm to thoroughly cover your hands
- Rub hands together briskly until dry

Other information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces.

Inactive ingredients

Aqua, PEG-4 Diheptanoate, Acrylates/C10-30 Alkyl Acrylate crosspolymer, Fragrance,

Aminomethyl Propanol

May contain hydroxypropylcellulose.

284 ml Bottle Label

Sani Luxe[®]

HAND SANITIZER GEL

with Alcohol

- 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.

284ml e

SaniLuxe[®]

HAND SANITIZER GEL

with Alcohol

- 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.

284mL

Celeste Industries Corporation
Easton, MD 21601 USA 36M
++1.410.822.5775
www.celestecorp.com

Drug Facts

Active ingredient Ethyl Alcohol 62.5%	Purpose Antimicrobial	Inactive ingredients Aqua, PEG-4 Diheptanoate, Acrylates/C10-30 Alkyl Acrylate crosspolymer, Fragrance, Aminomethyl Propanol May contain hydroxypropylcellulose.
Uses <ul style="list-style-type: none"> Recommended for repeated use For hand sanitizing to decrease bacteria on the skin 		

Warnings

- For external use only.
- Flammable. Keep away from open flame and sources of heat.
- Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.
- When using this product, avoid contact with eyes. In case of eye contact, flush eyes with water.
- If swallowed, get medical help or contact a Poison Control Center right away.
- Keep out of reach of children.

Directions

- Children under 6 years of age should be supervised when using this product
- Place enough product in your palm to thoroughly cover your hands
- Rub hands together briskly until dry

Other information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces.

SANI LUXE HAND SANITIZER WITH ALCOHOL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71489-007
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.625 mL in 1 mL

Inactive Ingredients

Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
PEG-4 DIHEPTANOATE (UNII: 2DQ7O61VHJ)				
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71489-007-01	284 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/24/2019	
2	NDC:71489-007-02	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2020	11/14/2022
3	NDC:71489-007-03	59 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/25/2020	
4	NDC:71489-007-04	237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020	11/14/2022
5	NDC:71489-007-05	355 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/25/2020	11/14/2022
6	NDC:71489-007-06	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2020	11/14/2022
7	NDC:71489-007-07	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
8	NDC:71489-007-08	177 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/25/2020	11/14/2022
9	NDC:71489-007-09	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	11/14/2022
10	NDC:71489-007-10	284 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	
11	NDC:71489-007-11	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/25/2020	11/14/2022
12	NDC:71489-007-12	3785 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/25/2020	11/14/2022
13	NDC:71489-007-13	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/30/2020	
14	NDC:71489-007-14	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/30/2020	
15	NDC:71489-007-15	530 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/14/2020	
16	NDC:71489-007-16	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)	01/24/2019	

Labeler - Celeste Industries Corporation (047795034)

