MOISTURIZING FOAM HAND SANITIZER- alcohol liquid Volu-Sol

Active Ingredient(s)

Alcohol 74.3% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water, Glycerin, 2-(Perfluorohexyl)ethanol, PEG-10 Acrylate/Perfluorohexylethylacrylate Copolymer, Hydrogen Peroxide

Package Label - Principal Display Panel

1893 mL NDC: 74401-500-64



Drug Facts

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400 mL NDC: 74401-500-40





MOISTURIZING FOAM

UNSCENTED

13.5 fl oz. (400 mL)

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MADE IN THE U.S.A.

Volu-Sol 800-821-2495 Salt Lake City, UT 84120

NDC: 74401-500-40



3785 mL NDC: 74401-500-12

Foaming Hand Sanitizer Moisturizing Foam, Unscented

ID: WHF-128 NDC: 74401-500-12

3.79 L / 1 gal

LOT

11182020



11/2025



Hand Sanitizer, Antiseptic

See SDS for further information



Volu-Sol 5095 W 2100 S Salt Lake City, UT 84120 Phone: 1-800-821-2495 www.volusol.com customerservice@volusol.com





Drug Facts

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Uses

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MOISTURIZING FOAM HAND SANITIZER

alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:74401-500

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	74.3 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
PEG-10 ACRYLATE/PERFLUOROHEXYLETHYL ACRYLATE COPOLYMER (UNII: D76Z87928N)	0.165 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.35 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
2-(PERFLUOROHEXYL)ETHANOL (UNII: G2R5YO5N3V)	0.165 mL in 100 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74401- 500-33	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74401- 500-12	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	
3	NDC:74401- 500-40	400 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/30/2020	
4	NDC:74401- 500-64	1893 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	03/30/2020	03/31/2023	

Labeler - Volu-Sol (050173424)

Registrant - Volu-Sol (050173424)

Establishment			
Name	Address	ID/FEI	Business Operations
Volu-Sol		050173424	manufacture(74401-500)

Revised: 1/2025 Volu-Sol