MICROSURE HAND SANITIZER- benzalkonium chloride solution Strategia Project Management, Inc.

Microsure Hand Sanitizer

DRUG FACTS

ACTIVE INGREDIENT

Benzalkonium Chloride 0.13%

PURPOSE

Antimicrobial

USES:

Sanitizes hands to helps reduce bacteria that potentially cause diseases.

WARNINGS:

For external use only.

Do not use

in eyes. If contact occurs, flush thoroughly 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:

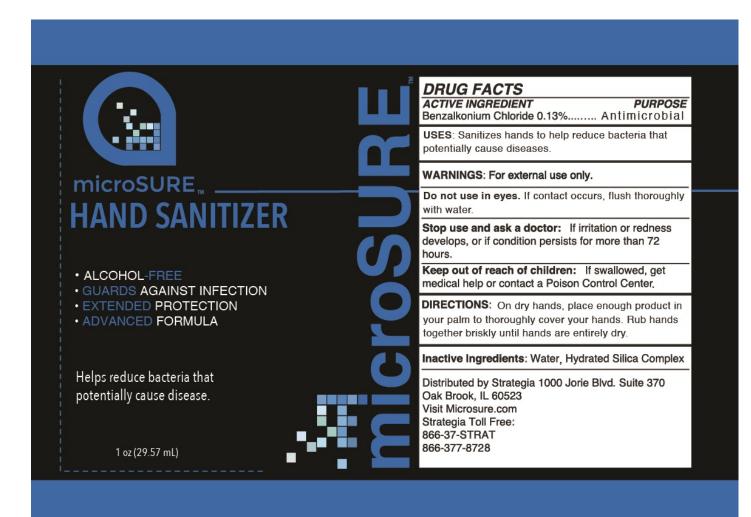
If swallowed, get medical help or contact a Poison Control Center.

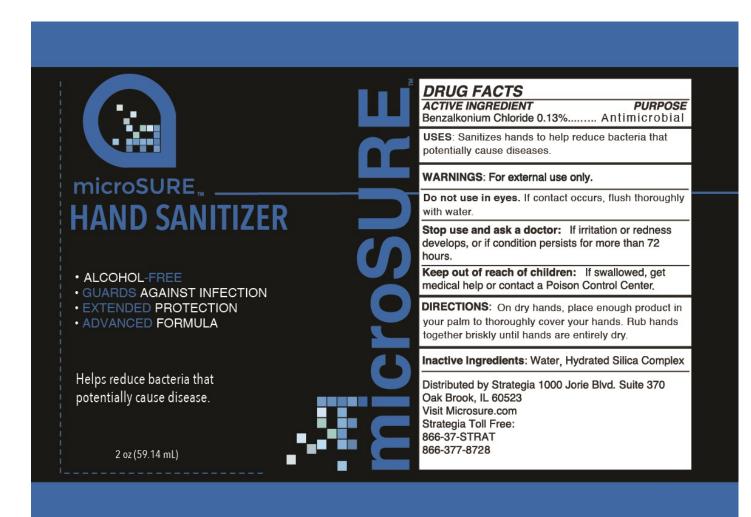
DIRECTIONS:

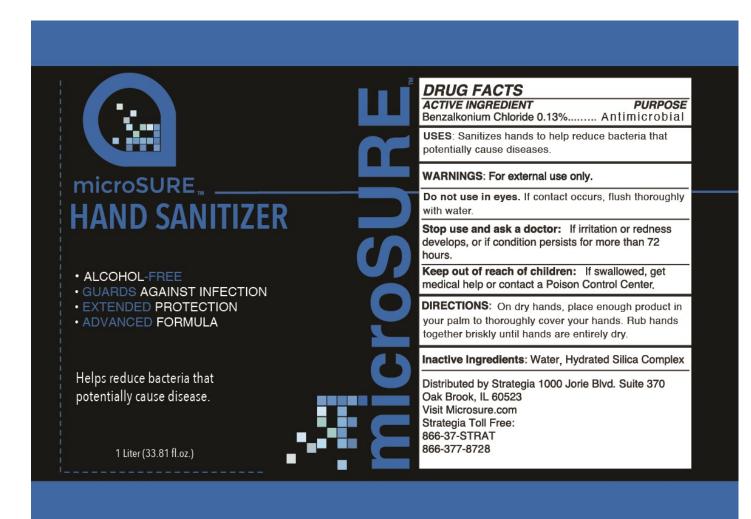
Wipe on both hands to disinfect palms, between fingers, back of hands, and wrists. 8 hours of continues protection once skin is fully dry.

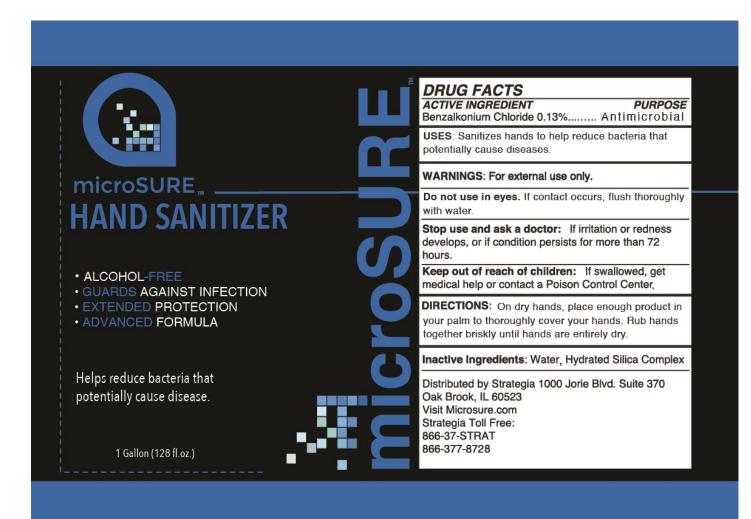
Inactive Ingredients:

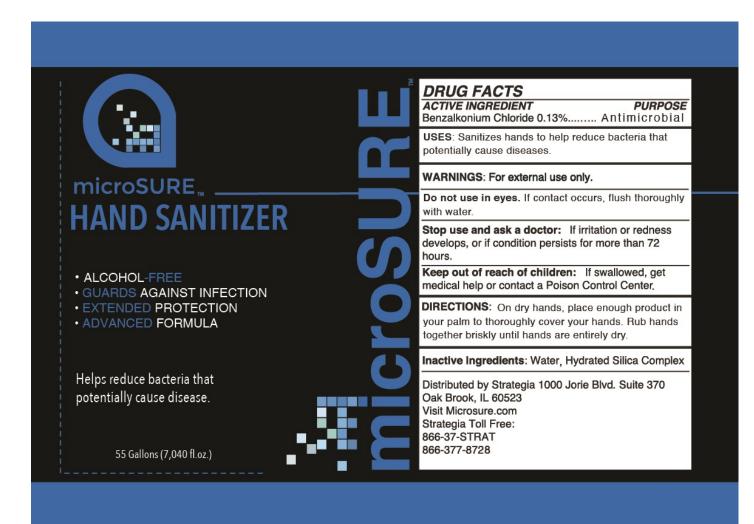
Water, Hydrated Silica Complex

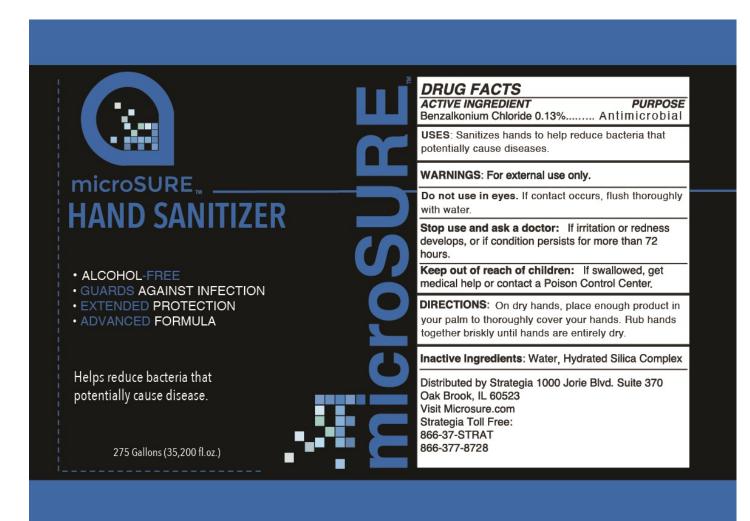














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ACTIVE INGREDIENT **PURPOSE** Benzalkonium Chloride 0.13%...... Antimicrobial

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WARNINGS: For external use only.

Do not use in eyes. If contact occurs, flush thoroughly with water.

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

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

DIRECTIONS: On dry hands, place enough product in your palm to thoroughly cover your hands. Rub hands together briskly until hands are entirely dry.

Inactive Ingredients: Water, Hydrated Silica Complex

Distributed by Strategia 1000 Jorie Blvd. Suite 370 Oak Brook, IL 60523 Visit Microsure.com Strategia Toll Free: 866-37-STRAT 866-377-8728

Strength

MICROSURE HAND SANITIZER

benzalkonium chloride solution

Product Information

HUMAN OTC DRUG Item Code (Source) NDC:72513-000 **Product Type**

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -**BENZ ALKONIUM** 1.3 mg UNII:7N6JUD5X6Y) **CHLORIDE** in 1 mL

Inactive Ingredients Ingredient Name

HYDRATED SILICA (UNII: Y6O7T4G8P9)

WATER (UNII: 059QF0KO0R)

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:72513- 000-01	29.57 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2020		
2	NDC:72513- 000-02	59.14 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2020		
3	NDC:72513- 000-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/20/2020		
4	NDC:72513- 000-04	3785.41 mL in 1 JUG; Type 0: Not a Combination Product	07/20/2020		
5	NDC:72513- 000-05	208198 mL in 1 DRUM; Type 0: Not a Combination Product	07/20/2020		
6	NDC:72513- 000-06	1040988.2 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/20/2020		
7	NDC:72513- 000-07	1249185.9 mL in 1 CONTAINER; Type 0: Not a Combination Product	07/20/2020		

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M016	07/20/2020				

Labeler - Strategia Project Management, Inc. (054047254)

Revised: 12/2023 Strategia Project Management, Inc.