

**ELIMISHIELD CLINICAL CARE TECHNOLOGY ALCOHOL-FREE HAND SANITIZER-
benzalkonium chloride liquid
Bryson Industries Inc**

ElimiShield Clinical Care Technology Alcohol-Free Hand Sanitizer

Drug Facts

Active Ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic Hand Sanitizer

Uses

- Hand sanitizer to help decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

For external use only.

- Do not use in ears or mouth.

When using this product

- Avoid contact with the eyes
- In case of contact flush eyes with water

Stop use and ask a doctor if

redness or irritation develop and persist for more than 72 hours.

Keep out of reach of children.

Children should be supervised when using this product.

Directions

- Apply a small amount into palms of hands and spread on both hands.
- Rub into skin until dry.

Inactive Ingredients

1-Octadecanaminium NN dimethyl (3-trimethoxysilyl) propyl chloride, 1-

Octadecanaminium NN dimethyl(3-trihydroxysilyl)propylchloride, Aloe Barbadenis leaf extract, Aqua, Caprylyl glucoside, Citric acid, Laureth-4, Methylparaben, Polyaminopropyl biguanide, Silk protein

Package Labeling



Alcohol-Free **HAND SANITIZER**



**KILLS 99.99%
OF GERMS**



**NON-IRRITATING
FORMULA**



**LONG-LASTING
PROTECTION**

96 fl oz (2.84L)

← PRESS WEB DIRECTION



Apply into palms of hands
and rub hands together for
about 30 seconds, until skin is dry.
Provides long lasting protection.

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Inactive Ingredients 1-Octadecanaminium NN dimethyl (3-trimethoxysilyl) propyl chloride, 1-Octadecanaminium NN dimethyl (3-trihydroxysilyl) propyl chloride, Aloe Barbadenis leaf extract, Aqua,

Caprylyl glucoside, Citric acid, Laureth-4, Methylparaben, Polyaminopropyl biguanide, Silk protein

BELM0390



For More Information Regarding Safety
Of This Product: info@BrysonUSA.com



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MADE IN THE USA

← PRESS WEB DIRECTION

ELIMISHIELD CLINICAL CARE TECHNOLOGY ALCOHOL-FREE HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71853-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
DIMETHYLOCTADECYL(3-(TRIMETHOXYSILYL)PROPYL)AMMONIUM CHLORIDE (UNII: IQ36O85WQ4)	
OCTADECYLDIMETHYL(3-TRIHYDROXYSILYLPROPYL)AMMONIUM CHLORIDE (UNII: GLJ50K866T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
LAURETH-4 (UNII: 6HQ855798J)	
METHYLPARABEN (UNII: A2I8C7HI9T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71853-004-96	2840 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/22/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/22/2018	

Labeler - Bryson Industries Inc (040363256)

Revised: 11/2023

Bryson Industries Inc