**ELIMISHIELD CLINICAL HAND SANTIZER- benzalkonium chloride liquid**
**Bryson Industries Inc.**

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**ElimiShield Clinical Hand Santizer**

## *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-trihydroxysily) propyl chloride, Aloe Barbadenis leaf extract, Aqua, Caprylyl glucoside, Citric acid, Laureth-4, Methylparaben, Polyaminopropyl Biguanide, Silk protein

**Package Labeling:**


## ELIMISHIELD CLINICAL HAND SANTIZER

benzalkonium chloride liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:71853-001 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 13 mg in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |
| **CAPRYLYL GLUCOSIDE** (UNII: V109WJT6RL) | |
| **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:71853-001-01 | 50 mL in 1 BOTTLE; Type 0: Not a Combination Product | 11/02/2017 | |
| **2** | NDC:71853-001-07 | 207 mL in 1 BOTTLE; Type 0: Not a Combination Product | 11/02/2017 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 11/02/2017 | |

**Labeler -** Bryson Industries Inc. (040363256)

Revised: 11/2023 Bryson Industries Inc.