**PRIME SOURCE ANTIBACTERIAL FOAM HANDWASH- benzalkonium chloride liquid**
**BUNZL**

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**PRIME SOURCE® Antibacterial Foam Handwash**

## Active ingredient

Benzalkonium Chloride 0.5%

## Purpose

Antimicrobial

## Uses

- Handwash to help decrease bacteria on the skin
- Recommended for repeated use

## Warnings

**For external use only**

**When using this product** do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

**Stop use and ask a doctor if** irritation or rash appears and lasts

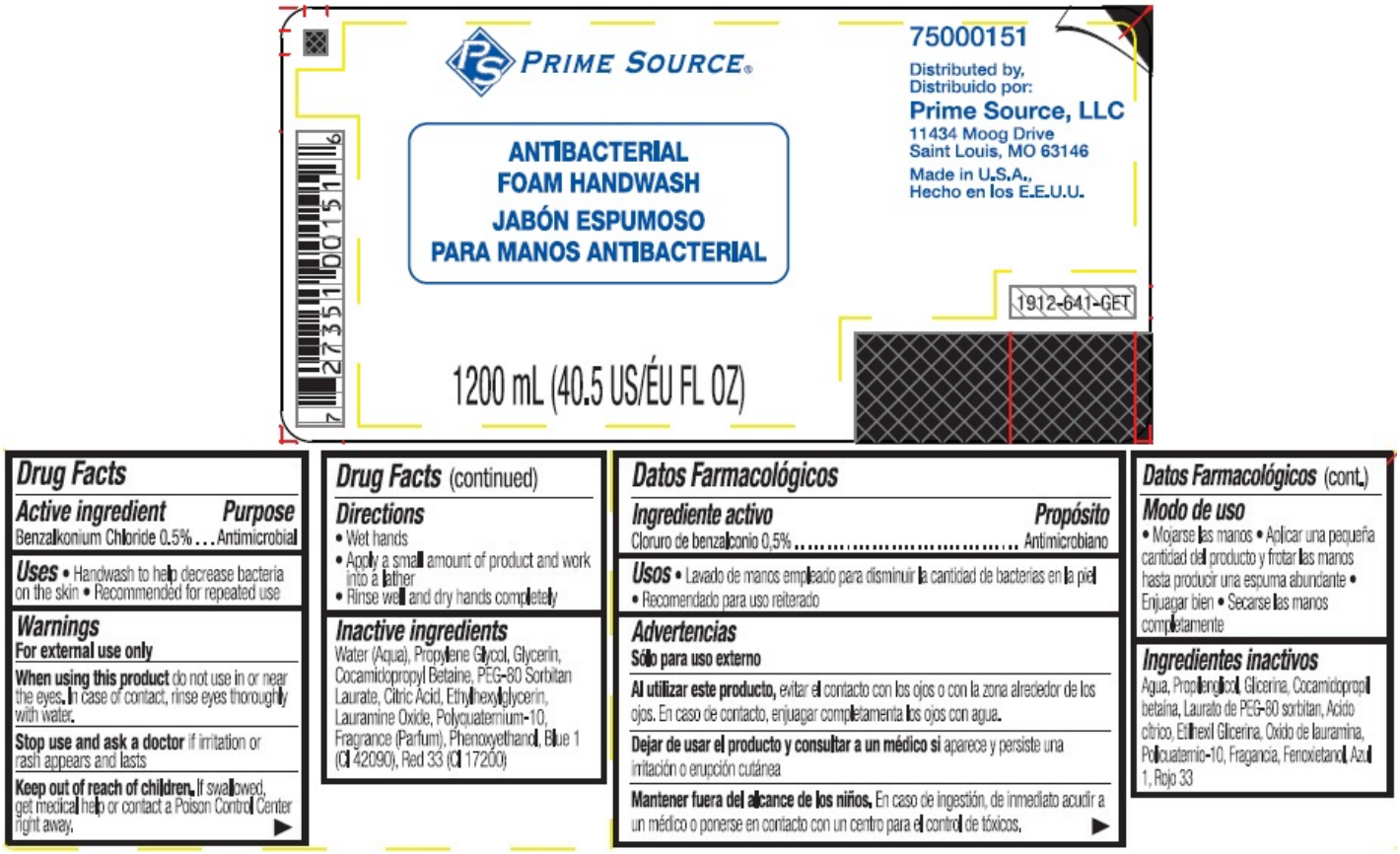
**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

## Directions

- Wet hands
- Apply a small amount of product and work into a lather
- Rinse well and dry hands completely

## Inactive Ingredients

Water (Aqua), Propylene Glycol, Glycerin, Cocamidopropyl Betaine, PEG-80 Sorbitan Laurate, Citric Acid, Ethylhexylglycerin, Lauramine Oxide, Polyquaternium-10, Fragrance (Parfum), Phenoxyethanol, Blue 1 (CI 42090), Red 33 (CI 17200)




# PRIME SOURCE ANTIBACTERIAL FOAM HANDWASH

benzalkonium chloride liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:66294-400 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 0.005 mg in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX) | |
| **PEG-80 SORBITAN LAURATE** (UNII: 239B50Y732) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **ETHYLHEXYLGLYCERIN** (UNII: 147D247K3P) | |
| **LAURAMINE OXIDE** (UNII: 4F6FC4MI8W) | |
| **POLYQUATERNIUM-10 (10000 MPA.S AT 2%)** (UNII: PI1STR9QYH) | |

| | |
|---|---|
| **PHENOXYETHANOL** (UNII: HIE492ZZ3T) | |
| **FD&C BLUE NO. 1** (UNII: H3R47K3TBD) | |
| **D&C RED NO. 33** (UNII: 9DBA0SBB0L) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:66294-400-23 | 700 mL in 1 PACKAGE; Type 0: Not a Combination Product | 04/30/2017 | |
| **2** | NDC:66294-400-40 | 1200 mL in 1 PACKAGE; Type 0: Not a Combination Product | 04/30/2017 | |
| **3** | NDC:66294-400-42 | 1250 mL in 1 PACKAGE; Type 0: Not a Combination Product | 04/30/2017 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 04/30/2017 | |

## Labeler - BUNZL (799540588)

Revised: 12/2024 BUNZL