**FOAMING HAND- benzalkonium chloride soap**
**UpLift Brands, LLC**

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**Germ-X 224.000/224AA**
**Antibacterial Foaming Hand Soap**

## Active ingredient

Benzalkonium chloride 0.13%

## Purpose

Antibacterial

## Use

for handwashing to decrease bacteria on the skin

## Warnings

**For external use only: hands only**

## When using this product

avoid contact with eyes. If contact occurs, rinse eyes with water.

## Stop use and ask a doctor if

- irritation or redness develop
- condition persists for more than 72 hours

## Keep out of reach of children.

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

## Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

## Inactive ingredients

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, citric acid, tetrasodium EDTA, sodium benzoate

**Adverse event**

Manufactured By: Vi-Jon, Inc.

St. Louis, Mo 63114

**Principal display panel**

germ-X

Professional

ANTIBACTERIAL

FOAMING

HAND SOAP

1000 ML (33.8 FL OZ)


## FOAMING HAND

benzalkonium chloride soap

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:83986-224 |
| **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 |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX) | |
| **LAURAMIDOPROPYLAMINE OXIDE** (UNII: I6KX160QTV) | |
| **LAURAMINE OXIDE** (UNII: 4F6FC4MI8W) | |
| **MYRISTAMIDOPROPYLAMINE OXIDE** (UNII: 3HSF539C9T) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **EDETATE SODIUM** (UNII: MP1J8420LU) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:83986-224-08 | 3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/11/2024 | |
| **2** | NDC:83986-224-45 | 1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/11/2024 | |
| **3** | NDC:83986-224-86 | 1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product | 04/11/2024 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 04/11/2024 | |

## Labeler - UpLift Brands, LLC (119091527)

## Registrant - Consumer Product Partners, LLC (119091520)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Consumer Product Partners, LLC | | 119091520 | manufacture(83986-224) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Consumer Product Partners, LLC | | 119091514 | manufacture(83986-224) |

Revised: 4/2024 UpLift Brands, LLC