**DAYLOGIC ANTIBACTERIAL FOAMING WASH- benzalkonium chloride soap**
**Rite Aid Corporation**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Drug Facts**

**Active ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Uses**

for hand washing to decrease bacteria on the skin.

**Warnings**

For external use only.

**When using this product**

avoid contact with eyes. In case of contact, rinse thoroughly with water.

Stop use and ask a doctor if

irritation and redness develops and lasts

*Keep out of reach of children.*

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

**Directions**

- apply onto dry hands.
- lather and rinse thoroughly.

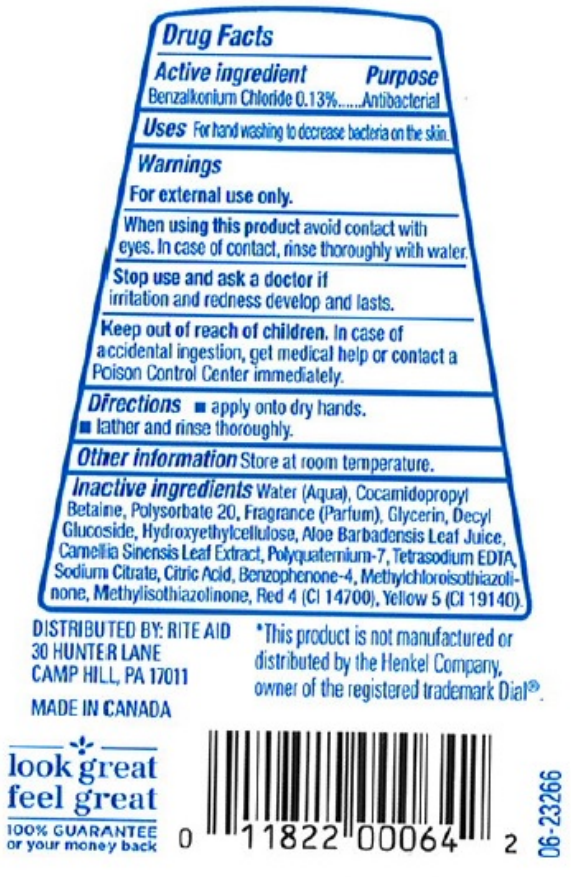
**Other information**

store at room temperature.

**Inactive ingredients**

Water (Aqua), Cocamidopropyl Betaine, Polysorbate 20, Fragrance (Parfum), Glycerin, Decyl Glucoside, Hydroxyethylcellulose, Aloe Barbadensis Leaf Juice, Camellia Sinensis Leaf Extract, Polyquaternium-7, Tetrasodium EDTA, Sodium Citrate, Citric Acid, Benzophenone-4, Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 5 (CI 19140).

**Label Copy**


# DAYLOGIC ANTIBACTERIAL FOAMING WASH

benzalkonium chloride soap

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:11822-1240 |
| **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) | |
| **POLYSORBATE 20** (UNII: 7T1F30V5YH) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **DECYL GLUCOSIDE** (UNII: Z17H97EA6Y) | |
| **HYDROXYETHYL CELLULOSE (5000 CPS AT 1%)** (UNII: X70SE62ZAR) | |
| **ALOE VERA LEAF** (UNII: ZY81Z83H0X) | |

| | |
|---|---|
| **GREEN TEA LEAF** (UNII: W2ZU1RY8B0) | |
| **POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW)** (UNII: 0L414VCS5Y) | |
| **EDETATE SODIUM** (UNII: MP1J8420LU) | |
| **SODIUM CITRATE** (UNII: 1Q73Q2JULR) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **SULISOBENZONE** (UNII: 1W6L629B4K) | |
| **METHYLCHLOROISOTHIAZOLINONE** (UNII: DEL7T5QRPN) | |
| **METHYLISOTHIAZOLINONE** (UNII: 229D0E1QFA) | |
| **FD&C RED NO. 4** (UNII: X3W0AM1JLX) | |
| **FD&C YELLOW NO. 5** (UNII: I753WB2F1M) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:11822-1240-8 | 222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/28/2017 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333E | 04/28/2017 | |

## Labeler - Rite Aid Corporation (014578892)

## Registrant - Apollo Health and Beauty Care Inc. (201901209)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Apollo Health and Beauty Care Inc. | | 201901209 | manufacture(11822-1240) |

Revised: 5/2017 Rite Aid Corporation