**G AND H PROTECT HAND- benzalkonium chloride soap**
**Access Business Group LLC**

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**g&h Protect Hand Soap**

**Drug Facts**

**Active ingredient**

Benzalkonium Chloride 0.10%

***Purpose***

Antibacterial

**Uses**

Helps reduce bacteria on hands.

**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 develops. These may be signs of a serious condition.

**Keep out of reach of children**

except under adult supervision. If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

wash hands and rinse.

**Inactive ingredients:**

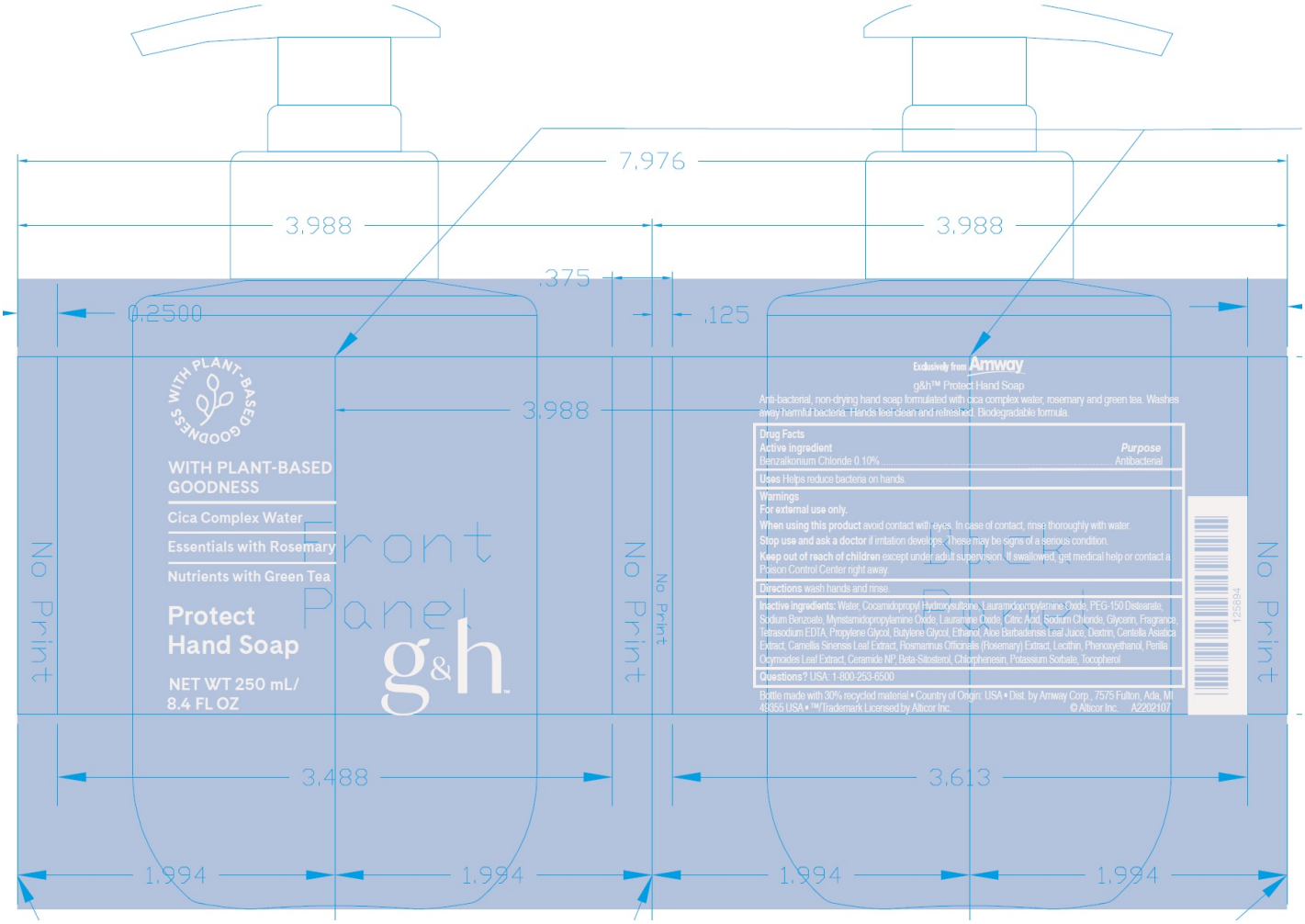
Water, Cocamidopropyl Hydroxysultaine, Lauramidopropylamine Oxide, PEG-150 Distearate, Sodium Benzoate, Myristamidopropylamine Oxide, Lauramine Oxide, Citric Acid, Sodium Chloride, Glycerin, Fragrance, Tetrasodium EDTA, Propylene Glycol, Butylene Glycol, Ethanol, Aloe Barbadensis Leaf Juice, Dextrin, Centella Asiatica Extract, Camellia Sinensis Leaf Extract, Rosmarinus Officinalis (Rosemary) Extract, Lecithin,

Phenoxyethanol, Perilla Ocymoides Leaf Extract, Ceramide NP, Beta-Sitosterol, Chlorphenesin, Potassium Sorbate, Tocopherol

## Questions?

USA: 1-800-253-6500

## Package Labeling:




# G AND H PROTECT HAND

benzalkonium chloride soap

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:10056-026 |
| **Route of Administration** | TOPICAL | | |

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

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **COCAMIDOPROPYL HYDROXYSULTAINE** (UNII: 62V75NI93W) | |
| **LAURAMIDOPROPYLAMINE OXIDE** (UNII: I6KX160QTV) | |
| **PEG-150 DISTEARATE** (UNII: 6F36Q0I0AC) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **MYRISTAMIDOPROPYLAMINE OXIDE** (UNII: 3HSF539C9T) | |
| **LAURAMINE OXIDE** (UNII: 4F6FC4MI8W) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **EDETATE SODIUM** (UNII: MP1J8420LU) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **BUTYLENE GLYCOL** (UNII: 3XUS85K0RA) | |
| **ALCOHOL** (UNII: 3K9958V90M) | |
| **ALOE VERA LEAF** (UNII: ZY81Z83H0X) | |
| **CENTELLA ASIATICA TRITERPENOIDS** (UNII: 4YS74Q4G4J) | |
| **GREEN TEA LEAF** (UNII: W2ZU1RY8B0) | |
| **ROSEMARY** (UNII: IJ67X351P9) | |
| **PHENOXYETHANOL** (UNII: HIE492ZZ3T) | |
| **PERILLA FRUTESCENS LEAF** (UNII: T4L5881Y68) | |
| **CERAMIDE NP** (UNII: 4370DF050B) | |
| **.BETA.-SITOSTEROL** (UNII: S347WMO6M4) | |
| **CHLORPHENESIN** (UNII: I670DAL4SZ) | |
| **POTASSIUM SORBATE** (UNII: 1VPU26JZZ4) | |
| **TOCOPHEROL** (UNII: R0ZB2556P8) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:10056-026-00 | 250 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/19/2023 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 04/19/2023 | |

## Labeler - Access Business Group LLC (839830713)

Revised: 10/2023 Access Business Group LLC