**NORTH WOODS DERMA FOAM E-2- benzalkonium chloride soap**
**Superior Chemical Corporation**

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## North Woods Derma Foam E-2

### *Active Ingredient*

Benzalkonium Chloride 0.13%

### *Uses*

- For handwashing to decrease the bacteria on the skin.
- Recommended for repeated use.

### *Warnings*

- **For external use only.**
- When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.
- Stop use and ask a doctor If irritation persists or redness deelops, or if 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*

- **Read the entire label before using this product.**
- Dispense 2 pumps of product onto palm of hand and scrub thoroughly over all surfaces of both hands for 15 seconds. Rinse with clean water.

### *Inactive Ingredients*

Water, coco-glucoside, laurtrimonium chloride, cocamidopropylamine oxide, citric acid.

## Superior Derma Foam E2

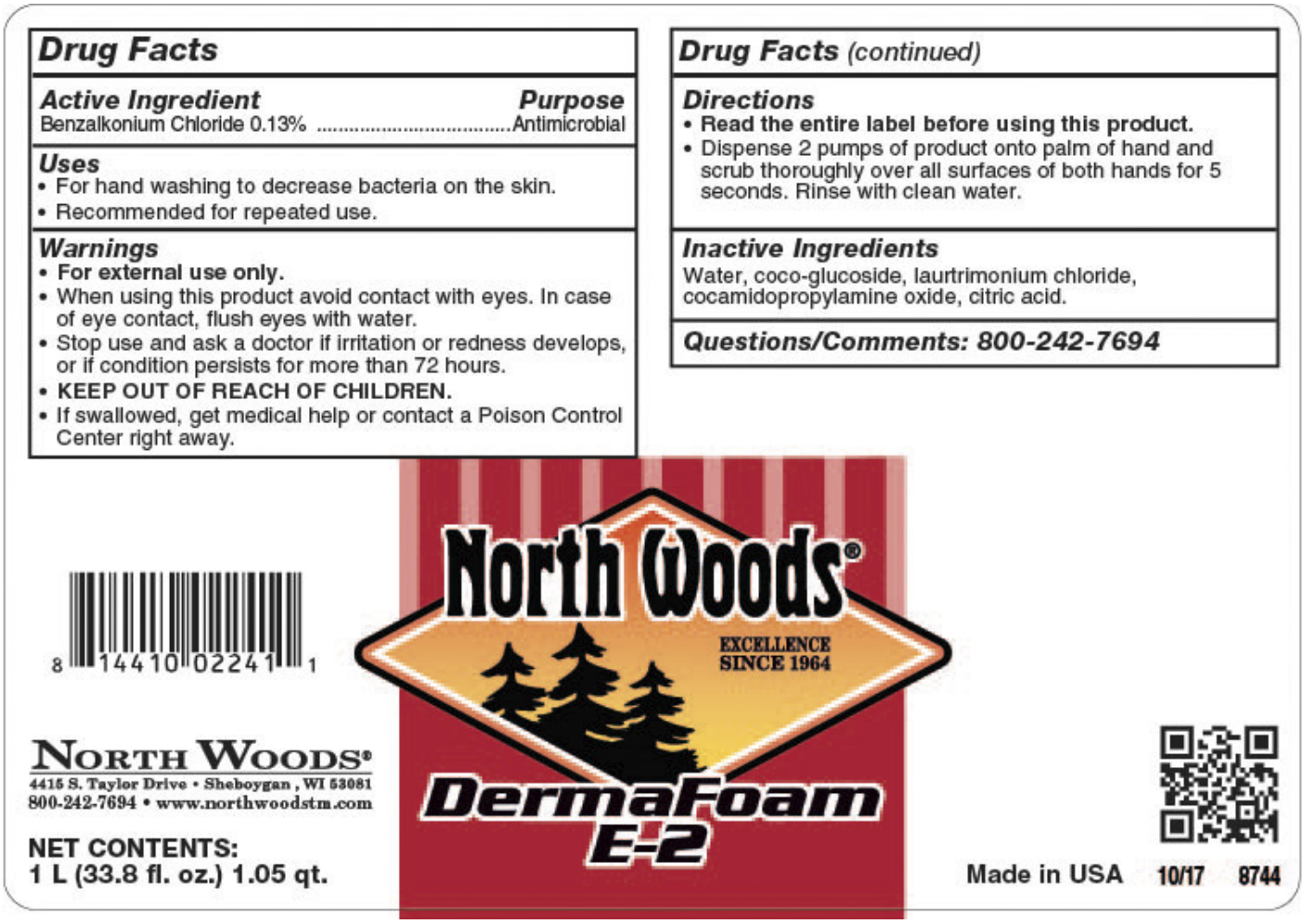
### *Purpose*

Antimicrobial

## Superior Derma Foam E2

KEEP OUT OF REACH OF CHILDREN

## Superior Derma Foam E2




71729-00 Superior Derma Foam E2

# NORTH WOODS DERMA FOAM E-2

benzalkonium chloride soap

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:53125-817 |
| **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** |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **ALCOHOL** (UNII: 3K9958V90M) | |
| **TETRASODIUM EDTA** (UNII: MP1J8420LU) | |

| | |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **LAURTRIMONIUM CHLORIDE** (UNII: A81MSI0FIC) | |
| **CAPRYLYL/CAPRYL OLIGOGLUCOSIDE** (UNII: E00JL9G9K0) | |
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:53125-817-29 | 1000 mL in 1 BAG; Type 0: Not a Combination Product | 09/15/2016 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 09/15/2016 | |

## Labeler - Superior Chemical Corporation (023335086)

## Registrant - Betco corporation, Ltd. (024492831)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Betco Corporation, Ltd. | | 005050158 | manufacture(53125-817) , label(53125-817) , pack(53125-817) |

Revised: 11/2024 Superior Chemical Corporation