**NORTH WOODS DERMA FOAM E-2- benzalkonium chloride soap**
**Superior Chemical 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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**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** |
| **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 not final | part333A | 09/15/2016 | |

## Labeler - Superior Chemical Corporation (023335086)

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

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Betco Corporation, Ltd. | | 024492831 | manufacture(53125-817) , label(53125-817) , pack(53125-817) |

Revised: 7/2020 Superior Chemical Corporation