

DERMAFOAM PREMIUM ANTIBACTERIAL- benzalkonium chloride soap
Superior Chemical Corporation

North Woods DermaFoam Premium Antibacterial

Active Ingredient

Triclosan 0.30%

Uses

- Antibacterial hand cleaner.
- Use in a variety of public facilities including daycare centers, hospitals, nursing homes, physicians offices.

Warnings

- **For external use only.**
- Avoid contact with eyes.
- Children under the age of 6 should be supervised by an adult when using this product.
- Discontinue use if irritation or redness develops.
- If irritation persists for more than 72 hours, consult a physician.
- **KEEP OUT OF REACH OF CHILDREN.**

Directions

- **Read the entire label before using this product.**
- Dispense 0.8 mL of product onto wet palm.
- Rub hands together to distribute product, then rinse hands with clean.

Inactive Ingredients

Water, Potassium cocoate (contains coconut), Propylene Glycol, Glycerine, DMDM Hydantoin, Fragrance, FD&C Yellow #5, Aloe Barbadensis Leaf Juice. FD&C Red #40.

Questions or Comments?Phone: (800) 777-9343

MDS information:(800) 891-4965

Purpose

Antibacterial

KEEP OUT OF REACH OF CHILDREN

Drug Facts

Active Ingredient **Purpose**
Benzalkonium Chloride 0.13% Antibacterial

Uses

- Antibacterial skin cleanser.
- Use in a variety of public facilities including daycare centers, hospitals, nursing homes, physicians offices.

Warnings

- **For external use only.** • Avoid contact with eyes.
- Discontinue use if irritation or redness develops.
- If irritation persists for more than 72 hours, consult a physician.
- **KEEP OUT OF REACH OF CHILDREN.**
- If swallowed, get medical help or contact a Poison Control Center right away.

Drug Facts (continued)

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.
- Rinse with clean water.

Inactive Ingredients

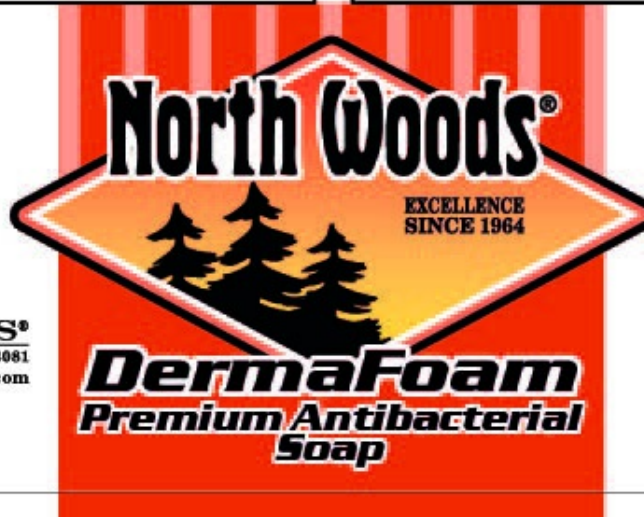
Water, Potassium Cocoate, Propylene Glycol, Glycerin, Tetrasodium EDTA, Fragrance, Bronopol, Sodium Citrate, Magnesium Nitrate, Sodium Hydroxyacetate, Ethanol, Aloe Barbadensis Leaf Juice, Trisodium Nitroacetate, FD&C Yellow #5, Sodium Hydroxide, Methylchlorisothiazolinone, Magnesium Chloride, Methylisothiazolinone, FD&C Red #40.

Questions/Comments: 800-242-7694



NORTH WOODS®
4415 S. Taylor Drive • Sheboygan, WI 53081
800-242-7694 • www.northwoodstm.com

NET CONTENTS:
1 L (33.8 fl. oz.) 1.05 qt.



Made in USA 04/17 7637

75129-00_Derma Foam Premium Antibacterial

DERMAFOAM PREMIUM ANTIBACTERIAL

benzalkonium chloride soap

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:53125-709 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-----------------------|-----------------|
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 0.13 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| WATER (UNII: 059QF0KO0R) | |
| POTASSIUM COCOATE (UNII: F8U72V8ZXP) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |

| | |
|---|--|
| EDETATE SODIUM (UNII: MP1J8420LU) | |
| DMDM HYDANTOIN (UNII: BYR0546TOW) | |
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN) | |
| METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA) | |
| FD&C RED NO. 40 (UNII: WZB9127XOA) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:53125-709-39 | 1000 mL in 1 BAG; Type 0: Not a Combination Product | 11/12/2012 | 01/01/2026 |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | 505G(a)(3) | 11/12/2012 | 01/01/2030 |

Labeler - Superior Chemical Corporation (023335086)

Registrant - Betco corporation, Ltd. (005050158)

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|---|
| Betco Corporation, Ltd. | | 005050158 | label(53125-709) , manufacture(53125-709) |

Revised: 11/2024

Superior Chemical Corporation