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

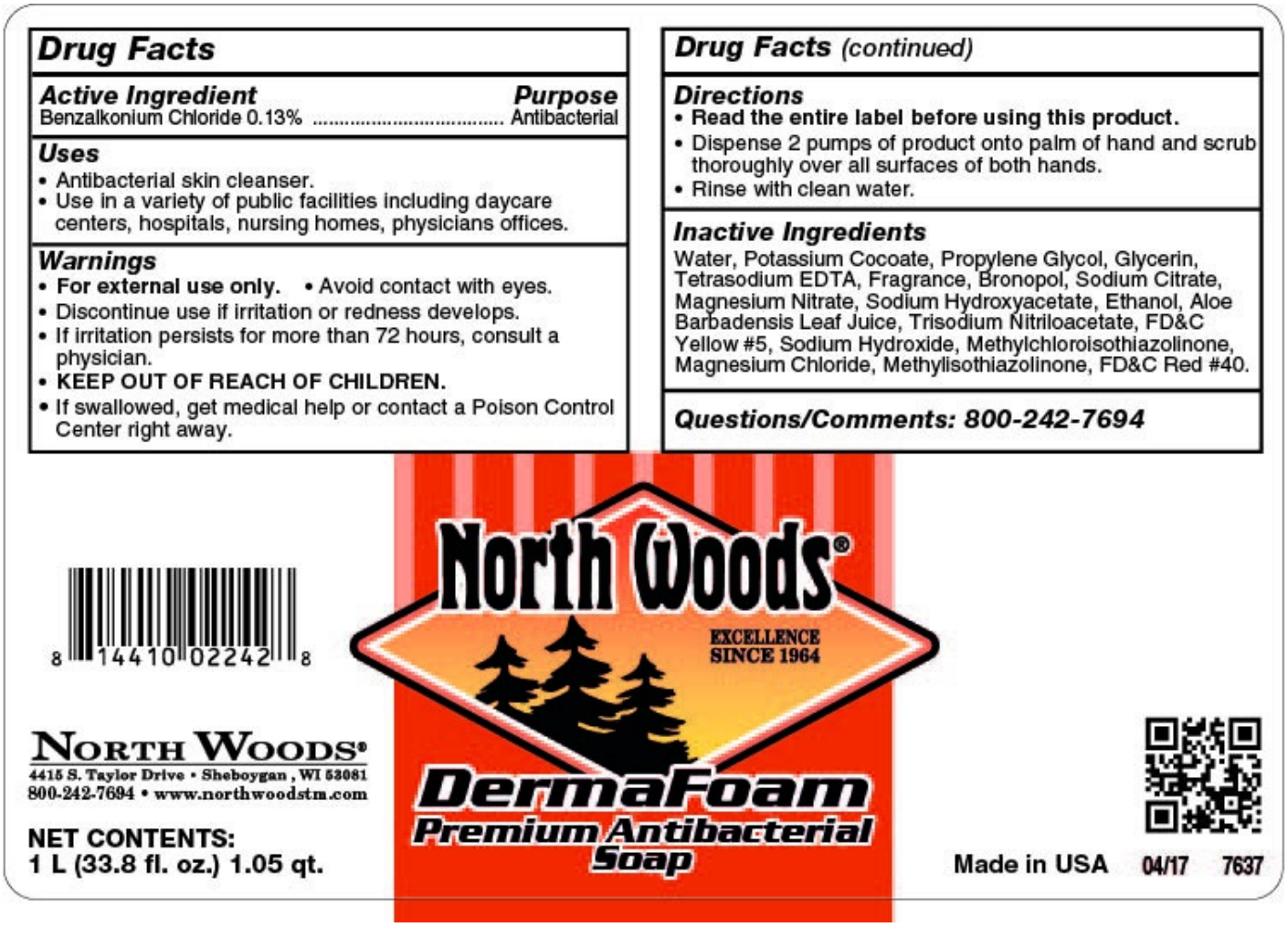
Questions or Comments?**Phone: (800) 777-9343**

MDS information:**(800) 891-4965**

Purpose

Antibacterial

KEEP OUT OF REACH OF CHILDREN



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