NORTH WOODS SUPREME ANTISEPTIC HAND CLEANSER- chloroxylenol 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.

North Woods Supreme Antiseptic Hand Cleanser

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Active Ingredient

Chloroxylenol 0.375%

North Woods Supreme Antiseptic Hand Cleanser

Uses

- Antibacterial hand cleaner.
- Use in daycare, hospitals, nursing homes, physicians offices, dental offices and clinics

North Woods Supreme Antiseptic Hand Cleanser

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**.

North Woods Supreme Antiseptic Hand Cleanser

Directions

- **Read the entire label before using this product.**
- Dispense 1-2 pumps of product onto wet palm.
- Lather and rinse hands with clean water

North Woods Supreme Antiseptic Hand Cleanser

Inactive Ingredients

Water, Sodium Lauryl lSulfate, Cocamide DEA, Cocamidopropyl betadine, Phenoxyethanol, Sodium Laureth Sulfate, Propylene Glycol, Fragrance, DMDM

Hydantoin, Glycol Stearate, Laurimide DEA, Glycerine, Tocopheryl Acetate, D&C Green #5, FD&C Yellow #5.

North Woods Supreme Antiseptic Hand Cleanser

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

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

North Woods Supreme Antiseptic Hand Cleanser

Purpose

Antibacterial

North Woods Supreme Antiseptic Hand Cleanser

KEEP OUT OF REACH OF CHILDREN

North Woods by Superior Supreme Antiseptic Hand Cleanser

SUperior 141

North Woods by Superior Supreme Antiseptic Hand Cleanser

Lotionized Antibacterial and pH Balanced

Excellence Since 1964

Superior Chemical Corporation

Sheboygan, WI 53081

NORTH WOODS SUPREME ANTISEPTIC HAND CLEANSER

chloroxylenol soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53125-708
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	3.75 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIMETHYL BENZYL CARBINYL BUTYRATE (UNII: 3Q0C60547R)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
LAURIC DIETHANOLAMIDE (UNII: I29I2VHG38)	
BENZYL BENZOATE (UNII: N863NB338G)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ETHYL ACETOACETATE (UNII: IZP61H3TB1)	
DIMETHYL BENZYL CARBINYL ACETATE (UNII: 6Y9488RL8H)	
SODIUM CARBONATE (UNII: 45P3261C7T)	
.GAMMA.-DECALACTONE (UNII: 7HLS05KP9O)	
ETHYL METHYLPHENYLGLYCIDATE (UNII: UD51D5KR4A)	
METHYL ANTHRANILATE (UNII: 981I0C1E5W)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM ALUMINIUM SILICATE (UNII: 058TS43PSM)	
CALCIUM SILICATE (UNII: S4255P4G5M)	
SODIUM FERROCYANIDE (UNII: 5HT6X21AID)	
LAURIC ISOPROPANOLAMIDE (UNII: 82DUX3RRVU)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
METHYL ALCOHOL (UNII: Y4S76JWI15)	
ETHYLENE OXIDE (UNII: JJH7GNN18P)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
CHLOROACETIC ACID (UNII: 5GD84Y125G)	
ETHYL BUTYRATE (UNII: UFD2LZ005D)	
MYRCENE (UNII: 3M39CZS25B)	
DIOXANE (UNII: J8A3S10O7S)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53125-708-19	900 mL in 1 BAG; Type 0: Not a Combination Product	11/12/2012	
2	NDC:53125-708-04	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/12/2012	

Labeler - Superior Chemical Corporation (023335086)

Registrant - Betco corporation, Ltd. (024492831)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corpo, Ltd.		024492831	manufacture(53125-708) , label(53125-708)

Revised: 2/2023

Superior Chemical Corporation