NORTH WOODS DERMA FOAM HAND SANITIZER- 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.

North Woods Derma Foam Hand Sanitizer

Active Ingredient

Benzalkonium Chloride 0.13%

Uses

- Use in a variety of public facilities.
- Use this product when soap and water are not available.

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 product onto dry hands. Rub hands together until hands are dry.
- Use as needed between hand washes to reduce bacteria on the skin.

Inactive Ingredients

Deionized IWater, Sodium PCA, PEG/PPG-8/3 Laurate, Dimethicone, DMDM Hydantoin, PEG-3 Cocamide, Fragrance, Iodoproynyl Btylcarbamate, D&C Green #5.

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

MDS information: (800) 891-4965

Purpose

Antiseptic

KEEP OUT OF REACH OF CHILDREN

Superior Derma Foam Hand Sanitizer.jpg

Drug Facts

Active Ingredient

Purpose

Benzalkonium Chloride 0.13%

. Antiseptic

Uses

- · Use in a variety of public facilities.
- . Use this product when soap and water are not available.

Warnings

- · For external use only.
- Avoid contact with eyes. If contact occurs, rinse thoroughly with water.
- 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 rub thoroughly over all surfaces of both hands until dry.

Inactive Ingredients

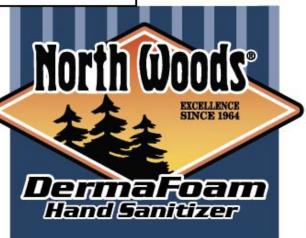
Deionized Water, Sodium PCA, PEG/PPG-8/3 Laurate, Dimethicone, PEG-3 Cocamide, Fragrance, Magnesium Salts, Methyl Chloro Isothiazolinone, Methyl Isothiazolinone, D&C Green #5.

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

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NORTH WOODS DERMA FOAM HAND SANITIZER

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benzalkonium chloride soap

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:53125-700

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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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 DIO XANE (UNII: J8 A3S 10 O7S) METHYLCHLO RO ISO THIAZO LINO NE (UNII: DEL7T5QRPN) CYCLO METHICO NE 4 (UNII: CZ227117JE) HEXYL SALICYLATE (UNII: 8F78 EY72YL) ALCO HOL (UNII: 3K9958 V90 M)

ISOPROPYL ALCOHOL (UNII: ND2M416302)
ACETALDEHYDE (UNII: GO 1N1ZPR3B)
LIME OIL (UNII: UZH29 XGA8G)
LEMON OIL (UNII: 19 GRO 8 2 4 LL)
MAGNESIUM NITRATE (UNII: 77CBG3UN78)
HEXAMETHYLINDANO PYRAN (UNII: 14170060 AT)
MYRCENE (UNII: 3M39 CZS25B)
DIHYDROMYRCENOL (UNII: 46L1B02ND9)
METHYL DIHYDRO JASMO NATE (SYNTHETIC) (UNII: 3GW44CIE3Y)
LINALOOL, (+/-)- (UNII: D81QY6I88E)
BUTYLPHENYL METHYLPROPIONAL (UNII: T7540 GJV69)
WATER (UNII: 059QF0KO0R)
GERANIOL (UNII: L837108USY)
SODIUM PIDOLATE (UNII: 1V74VH163T)
GRAPEFRUIT OIL (UNII: YR377U58W9)
ORANGE OIL (UNII: AKN3KSD11B)
PEG/PPG-15/15 ALLYL ETHER ACETATE (UNII: 8 RP39 FN7AJ)
PEG-6 CO CAMIDE (UNII: YZ6 NLA4O 1E)
N-ALKYL DIMETHYL BENZYL AMMONIUM CHLORIDE (C12-C18) (UNII: 9U1Q4T4ZYS)
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)
.BETACITRONELLOL, (+/-)- (UNII: 565OK72VNF)
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)

l	Packaging			
	# Item Code Package Description		Marketing Start Date	Marketing End Date
	1 NDC:53125-700- 88	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/01/2015	

Marketing Information					
Marketing Category Application Number or Monograph		Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	10/01/2015			

Labeler - Superior Chemical Corporation (023335086)

Registrant - Betco corporation, Ltd. (005050158)

Establishment				
Name	Address	ID/FEI	Business Operations	
Betco Corporation, Ltd.		005050158	manufacture(53125-700), pack(53125-700), label(53125-700)	