

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 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 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 ☐Water, Sodium PCA, PEG/PPG-8/3 Laurate, Dimethicone, DMDM Hydantoin, PEG-3 Cocamide, Fragrance, Iodopropynyl 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

75229-00_Superior Derma Foam Hand Sanitizer.

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

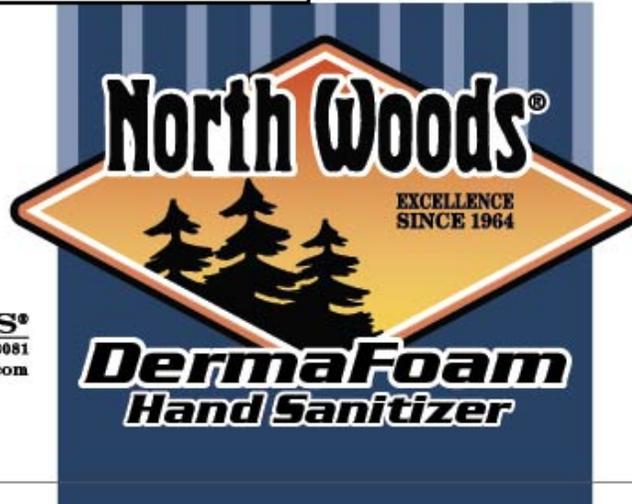
Deionized Water, Sodium PCA, PEG/PPG-8/3 Laurate, Dimethicone, PEG-3 Cocamide, Fragrance, Magnesium Salts, Methyl Chloro Isothiazolinone, Methyl Isothiazolinone, D&C Green #5.

Questions/Comments: 800-242-7694



NORTH WOODS®
4416 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 10/17 8750

NORTH WOODS DERMA FOAM HAND SANITIZER

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53125-700
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: 059QF0K00R)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
PEG-3 COCAMINE (UNII: KTM00873VC)	

IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)

D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53125-700-57	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/01/2015	
2	NDC:53125-700-54	1000 mL in 1 BAG; Type 0: Not a Combination Product	10/01/2015	

Marketing Information

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

Labeler - Superior Chemical Corporation (023335086)

Registrant - Betco corporation, Ltd. (024492831)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd.		024492831	manufacture(53125-700) , pack(53125-700) , label(53125-700)

Revised: 4/2019

Superior Chemical Corporation