NORTH WOODS DERMA GEL CLEAR- alcohol gel
Superior Chemical Corporation

North Woods Derma Gel Clear

Active Ingredient

Active Ingredient

Ethyl Alcohol 70%

Uses

Uses

• Hand sanitizer to reduce microorganisms on the skin.

• Use this product when soap and water are not available.

Purpose

Purpose

Antiseptic

Warnings

Warnings

• For external use only.

• Avoid contact with eyes. If contact occurs, rinse thoroughly with water.

• FLAMMABLE. This product contains ethyl alcohol. Keep away from sources of ignition.

• 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.

Directions

Directions

• Read the entire label before using this product.

• Place enough product on your palm to thoroughly cover your hands.

• Rub hands together briskly until dry.

Inactive Ingredients

Inactive Ingredients

Water, PEG/PPG-8/3 Laurate, Gycerin, Carbomer, Tetrahydroxypropylethylendiamine, Fragrance.

Warnings

KEEP OUT OF REACH OF CHILDREN.

North Woods Derma Gel Clear

Drug Facts

Active Ingredient — **Purpose**
Ethyl Alcohol 70% ..... Antiseptic

Uses
- Hand sanitizer to reduce microorganisms on the skin.
- Use this product when soap and water are not available.

Warnings
For external use only. FLAMMABLE. This product contains ethyl alcohol. Keep away from heat or flame.

When using this product avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs.

Keep out of the 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.**
- Place enough product on your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.

Other Information
- Store below 104°F (40°C).

Inactive Ingredients
Water, PEG/PPG-8/3 Laurate, Glycerin, Tetrahydroxypropylethylenediamine, Carbomer, Fragrance, Aloe Barbadensis Leaf Juice.

Questions/Comments: 800-242-7694

NDC# 53125-801-29

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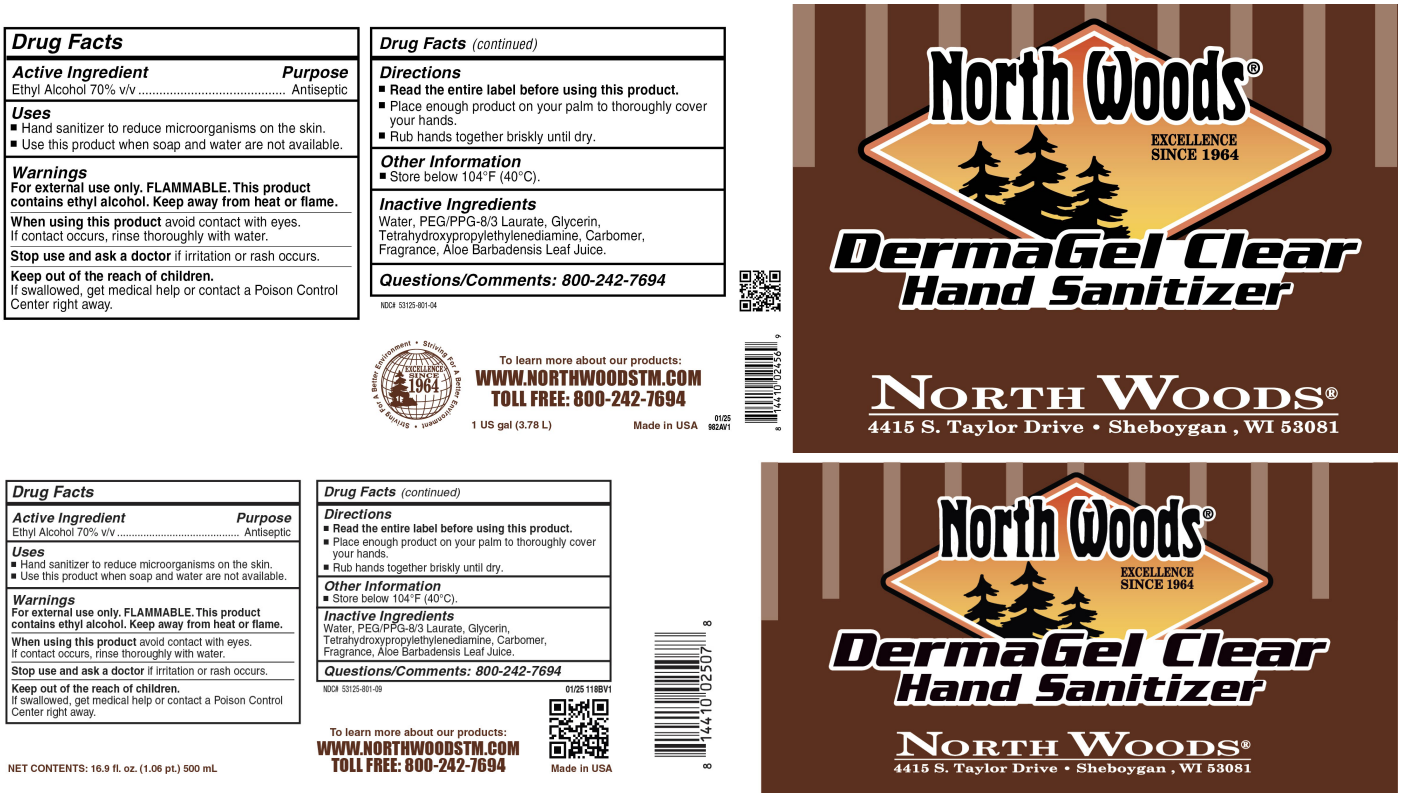
NORTH WOODS®
4415 S. Taylor Drive • Sheboygan , WI 53081
800-242-7694 • www.northwoodstm.com

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

North Woods®
EXCELLENCE SINCE 1964
DermaGel Clear Hand Sanitizer

Made in USA 01/25 090BV1



NORTH WOODS DERMA GEL CLEAR

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53125-801
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
PROPYLENE OXIDE (UNII: Y4Y7NYD4BK)	
ACRYLIC ACID (UNII: J94PBK7X8S)	
METHYL DIHYDROJASMONATE (SYNTHETIC) (UNII: 3GW44CIE3Y)	
LINALOOL (UNII: D81QY6I88E)	
HEXYL SALICYLATE (UNII: 8F78EY72YL)	
.ALPHA.-PINENE (UNII: JPF3YI7O34)	
CYCLOHEXANE (UNII: 48K5MKG32S)	
PEG-8 LAURATE (UNII: 762O8IWA10)	
WATER (UNII: 059QF0KO0R)	

CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETOL (UNII: Q4R969U9FR)	
GERANIOL (UNII: L837108USY)	
ALLYL CYCLOHEXANEPROPIONATE (UNII: H4W9H3L241)	
NEROL (UNII: 38G5P53250)	
OCTOXYNOL-13 (UNII: 480KVF3EBY)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
CITRONELLOL (UNII: P01OUT964K)	
.GAMMA.-TERPINENE (UNII: 4YGF4PQP49)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53125-801-57	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2019	
2	NDC:53125-801-04	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2019	
3	NDC:53125-801-09	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2019	08/20/2020
4	NDC:53125-801-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	08/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/01/2019	

Labeler - Superior Chemical Corporation (023335086)

Registrant - Betco Corporation, Ltd (005050158)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd		005050158	manufacture(53125-801) , label(53125-801)

Revised: 10/2025 Superior Chemical Corporation