

**GELLE-N-DETOX ICE MINT AND EUCALYPTUS MASSAGE- benzalkonium chloride lotion**  
**CHEMCO CORPORATION**

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**49283-014-07**

Benzalkonium Chloride 0.1%

Antiseptic

**Uses**

First aid to help protect against bacterial contamination.

- For external use only.
- Do not use in the eyes or apply over large areas of the body.
- In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.
- Stop use and ask a doctor if the condition persists or gets worse.
- Do not use longer than 1 week unless directed by a doctor

Ask a doctor before use if you have redness over affected area.

Ask a health professional before use.

If swallowed, get medical help or contact a Poison Control Center right away

**Directions**

Adults and children 12 years of age or older: • Clean the affected area. • Apply a small amount of this product on the area 1 to 3 times daily. • May be covered with a sterile bandage. Children under 12 years of age: do not use, consult a doctor.

Store tightly closed in a dry place at controlled room temperature between 59°-86° F (15°-30° C).

Water (Aqua), Paraffinum Liquidum, Cetyl Alcohol, Stearic Acid, Stearyl Alcohol, Fragrance (Parfum), DMDM Hydantoin, Triethanolamine, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Methylparaben, Propylparaben, Cocos Nucifera Oil (Organic), Argania Spinosa Kernel Oil (Organic), Methyl Salicylate, Eucalyptus Globulus Leaf Oil, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, FD&C Yellow No.5 (CI 19140), FD&C Blue No.1 (CI 42090), Citral, D-Limonene, Geraniol, Linalool.

Call (305) 623-4445

**GELLE-N-DETOX ICE MINT AND EUCALYPTUS MASSAGE LOTION 0.7 oz**

NDC#:49283-014-07

Gelle-n-Detox™



ICY MINT

CALMING - SOOTHING

MASSAGE LOTION



Body • Feet • Hands

**\*ANTISEPTIC**

Protects Against Bacteria Reducing the Risk of Skin Infection with Each Use

LUXURIOUS PEDICURE

Drug Facts

Active ingredient	Purpose
Benzalkonium chloride 0.1%	Antiseptic

Uses

First aid to help protect against bacterial contamination.

Warnings

• For external use only. Do not use in the eyes or apply over large areas of the body. In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor. • Stop use and ask a doctor if the condition persists or gets worse. • Do not use longer than 1 week unless directed by a doctor.

Ask a doctor before use if you have any redness over affected area.

If pregnant or breast – feeding

Ask a health professional before use.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 12 years of age or older:

• Clean the affected area. • Apply a small amount of this product on the area 1 to 3 times daily. • May be covered with a sterile bandage.

Children under 12 years of age: do not use, consult a doctor.

Other information

Store tightly closed in a dry place at controlled room temperature between 59°-86° F (15°-30° C).

Inactive ingredients

Water (Aqua), Paraffinum Liquidum, Cetyl Alcohol, Stearic Acid, Stearyl Alcohol, Fragrance (Parfum), DMDM Hydantoin, Triethanolamine, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Methylparaben, Propylparaben, Cocos Nucifera Oil (Organic), Argania Spinosa Kernel Oil (Organic), Methyl Salicylate, Eucalyptus Globulus Leaf Oil, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, FD&C Yellow No.5 (CI 19140), FD&C Blue No.1 (CI 42090), Citral, D-Limonene, Geraniol, Linalool.

Questions?

Call (305) 623-4445



Net: 0.7 fl oz e (20 mL)

Chemco Corp.  
4920 NW 165 St.  
Miami, FL 33014  
www.chemco.com  
Manufactured in USA

GELLE-N-DETOX ICE MINT AND EUCALYPTUS MASSAGE

benzalkonium chloride lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49283-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	

<b>TRIDECETH-6</b> (UNII: 3T5PCR2H0C)
<b>HYALURONATE SODIUM</b> (UNII: YSE9PPT4TH)
<b>SODIUM PYRROLIDONE CARBOXYLATE</b> (UNII: 469OTG57A2)
<b>AMINO ACIDS, WHEAT</b> (UNII: 0370GZL32F)
<b>SYMPHYTUM OFFICINALE WHOLE</b> (UNII: H8FJJ6KX5Y)
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)
<b>CITRAL</b> (UNII: T7EU009VPP)
<b>LIMONENE, (+)-</b> (UNII: GFD7C86Q1W)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>STEARYL ALCOHOL</b> (UNII: 2KR89I4H1Y)
<b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y)
<b>PANTHENOL</b> (UNII: WW9CM0067Z)
<b>EUCALYPTUS OIL</b> (UNII: 2R04ONI662)
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)
<b>ARGAN OIL</b> (UNII: 4V59G5UW9X)
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)
<b>COCONUT OIL</b> (UNII: Q9L0O73W7L)
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)
<b>DMDM HYDANTOIN</b> (UNII: BYR0546TOW)
<b>HYDROXYPROLINE</b> (UNII: RMB44WO89X)
<b>GERANIOL</b> (UNII: L837108USY)
<b>LINALOOL, (+)-</b> (UNII: F4VNO44C09)

### Product Characteristics

<b>Color</b>	green (Light green)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-014-07	20 g in 1 POUCH; Type 1: Convenience Kit of Co-Package	12/04/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/04/2025	

**Labeler** - CHEMCO CORPORATION (032495954)

**Registrant** - CHEMCO CORPORATION (032495954)

**Establishment**

Name	Address	ID/FEI	Business Operations
CHEMCO CORPORATION		032495954	manufacture(49283-014)

Revised: 12/2025

CHEMCO CORPORATION