

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

49283-110-07

Benzalkonium Chloride 0.1%

Antimicrobial

USES:

To decrease bacteria on the skin.

For external use only.

Do not use on wounds or damaged skin, with a heating pad or on a child under 12 years of age.

- Avoid contact with the eyes.
- Do not apply to the irritated skin or if excessive irritation develops.

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:

Apply a full-palm amount or as needed to cover the affected area (hands, feet, body). Rub until absorbed.

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

Water (Aqua), Paraffinum Liquidum, Stearic Acid, Cetyl Alcohol, Fragrance (Parfum), Stearyl Alcohol, DMDM Hydantoin, Triethanolamine, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Methylparaben, Propylparaben, Cocos Nucifera Oil (Organic), Argania Spinosa Kernel Oil (Organic), Lavandula Angustifolia (Lavender) Extract, Rosmarinus Officinalis (Rosemary) Extract, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, D&C Violet No.2 (CI 60730), Benzyl Salicylate, Linalool, Hexyl Cinnamal, Alpha Isomethyl Ionone, Citronellol, Geraniol.

Call (305) 623-4445

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

NDC#:49283-110-07

Gelle-n-Detox™

spaREDI™
pedicure therapy

CONTAINS
CERTIFIED
ORGANIC
INGREDIENTS

ICY MINT
CALMING - SOOTHING

MASSAGE LOTION

□ □ □ ■
Body • Feet • Hands

***ANTI-MICROBIAL**
Kills 99.9% of Germs & Bacteria *

LUXURIOUS PEDICURE

Drug Facts

Active ingredient Benzalkonium chloride 0.1% **Purpose** Antimicrobial

Uses
To decrease bacteria on the skin.

Warnings
• For external use only. • Do not use on wounds or damaged skin, with a healing pad or on a child under 12 years of age.
• Ask a doctor before use if you have any redness over affected area.

When using this product
• Avoid contact with the eyes.
• Do not apply to the irritated skin or if excessive irritation develops.
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
Apply a full-palm amount or as needed to cover the affected area (hands, feet, body). Rub until absorbed.

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

Questions?
Call (855) 523-4445

  

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

Net: 0.7 FL oz e (20 ml)

GELLE-N-DETOX ICE MINT AND EUCALYPTUS MASSAGE

benzalkonium chloride lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49283-110
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
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y)	

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)
CITRAL (UNII: T7EU009VPP)
LIMONENE, (+)- (UNII: GFD7C86Q1W)
WATER (UNII: 059QF0KO0R)
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)
METHYL SALICYLATE (UNII: LAV5U5022Y)
PANTHENOL (UNII: WW9CM0067Z)
EUCALYPTUS OIL (UNII: 2R04ONI662)
MINERAL OIL (UNII: T5L8T28FGP)
STEARIC ACID (UNII: 4ELV7Z65AP)
TROLAMINE (UNII: 9O3K93S3TK)
PROPYLPARABEN (UNII: Z8IX2SC1OH)
ARGAN OIL (UNII: 4V59G5UW9X)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)
COCONUT OIL (UNII: Q9L0O73W7L)
METHYLPARABEN (UNII: A2I8C7HI9T)
DMDM HYDANTOIN (UNII: BYR0546TOW)
HYDROXYPROLINE (UNII: RMB44WO89X)
GERANIOL (UNII: L837108USY)
LINALOOL, (+)- (UNII: F4VNO44C09)

Product Characteristics

Color	green (Light green)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-110-07	20 g in 1 POUCH; Type 1: Convenience Kit of Co-Package	01/19/2023	12/31/2027

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/19/2023	12/31/2027

Labeler - CHEMCO CORPORATION (032495954)

Registrant - CHEMCO CORPORATION (032495954)

Establishment

Name	Address	ID/FEI	Business Operations
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Revised: 12/2025

CHEMCO CORPORATION