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

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: T7EU0O9VPP)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
WATER (UNII: 059QF0KO0R)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
PANTHENOL (UNII: WV9CM0O67Z)	
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

CHEMCO CORPORATION		032495954	manufacture(49283-110)