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

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	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/19/2023	

Labeler - CHEMCO CORPORATION (032495954)

Registrant - CHEMCO CORPORATION (032495954)

Establishment

Name	Address	ID/FEI	Business Operations
CHEMCO CORPORATION		032495954	manufacture(49283-110)

Revised: 9/2023

CHEMCO CORPORATION