SPAREDI MINT AND EUCALYPTUS MASK- benzalkonium chloride cream
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.

49283-101-07

Benzalkonium chloride 0.1%

Antibacterial

USES:

To decrease bacteria on the skin.

DIRECTIONS:

Apply a thin layer to cover the foot. Leave for 10 minutes, then rinse thoroughly with warm water.

For external use only.

•Keep out of eyes. In case of contact with eyes, rinse thoroughly with water.• Do not apply to wounds or damaged skin.
•Do not apply to the irritated skin or if excessive irritation develops.

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

Aqua, Cetyl Alcohol, Stearyl Alcohol, Methyl Salicylate, Eucalyptus Globulus Leaf Oil, DMDM Hydantoin, Methylparaben, Propylparaben, Mentha Piperita (Peppermint) Oil, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, Cetrimonium Chloride, Argania Spinosa Kernel Oil (Organic), Cocos Nucifera Oil (Organic), FD&C Yellow #5 (CI 19140), FD&C Blue #1 (Cl 42090).

SPAREDI - MINT AND EUCALYPTUS CREAM MASK 0.7 oz

SPAREDI MINT AND EUCALYPTUS MASK

benzalkonium chloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49283-101
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**
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
ARGANIA SPINOSA WHOLE (UNII: 83K6O4FR76)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
MENTHA PIPERITA LEAF (UNII: A389O33LX6)	
CETYL ALCOHOL (UNII: 936JST6JCN)	

SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
COCOS NUCIFERA WHOLE (UNII: 245J88W96L)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PANTHENOL (UNII: WV9CM0O67Z)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SODIUM COCOYL WHEAT AMINO ACIDS (UNII: JW3VT57I11)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y)	

Product Characteristics

Color	green	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-101-07	20 g in 1 POUCH; Type 0: Not a Combination Product	01/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/17/2023	

Labeler - CHEMCO CORPORATION (032495954)

Registrant - CHEMCO CORPORATION (032495954)

Establishment

Name	Address	ID/FEI	Business Operations
CHEMCO CORPORATION		032495954	manufacture(49283-101)