

HAND ANTIPERSPIRANT DAILY USE- aluminum zirconium octachlorohydrate gly cream
Clutch Inc

HAND ANTIPERSPIRANT DAILY USE

Active Ingredient Purpose

Aluminum Zirconium Tetrachlorohydrate Gly (20%) Antiperspirant

Use

reduces perspiration

Keep out reach of children. If swallowed get medical help or contact a Poison Control Center right away

Stop use if rash or irritation occurs

Warnings

For external use only.

Do not use on broken or irritated skin

Ask a doctor before use if you have kidney disease.

Directions

- wash and dry hands thoroughly before application
- Apply a single pump of product to palms each morning
- Rub Balms together vigorously for 30 seconds

Inactive Ingredients Aluminum Starch Octenylsuccinate, Aqua (water), Cetyl Hydroxyethylcellulose, Coco-Caprylate/Caprate, Coconut Alkanes, Disodium EDTA, Ethylhexylglycerin, Glycerin, Helianthus Annuus (Sunflower) Extract, Hydroxypropyl Starch Phosphate, Octenidine HCL, Oryza SATiva (Rice) Bran Extract, Phenoxyethanol, Polymethylsilsesquioxane, Propanediol, Propylene Glycol, Rosmarinus Officinalis (Rosemary) Leaf Extract, SD Alcohol 40-B, Silica, Sodium Hydroxide, Tocopherol, Xanthan Gum

Carpe

Clinical Grade HAND

Antiperspirant

Daily Use

20% Aluminum Zirconium Octachlorohydrate GLY

Drug Facts		1 FL OZ (30 ML)	
Active Ingredient: Aluminum Zirconium Octachlorohydrate Gly (20%)	Purpose: Anti-perspirant	 NDC :007-74307-007 X-XXXX-XXXX :008 info@mycarpe.com 888-621-0135 Chapel Hill, NC 27517 Filled in USA	 carpe CLINICAL GRADE HAND ANTIPERSPIRANT DAILY USE 20% ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY
Use: • Reduces perspiration			
Warnings <i>For external use only</i>			
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Stop use and ask a doctor if • rash or irritation occurs			
Ask doctor before use if • you have kidney disease			
Keep out of reach of children • if swallowed, get medical help or contact a Poison Control Center right away			
Directions • Wash and dry hands thoroughly before application • Apply a single pump of product to palms each morning • Rub palms together vigorously for 30 seconds			
Inactive Ingredients Aluminum Starch Octenylsuccinate, Aqua (Water), Cetyl Hydroxyethylcellulose, Coco-Caprylate/Caprate, Coconut Alkanes, Disodium EDTA, Ethylhexylglycerin, Glycerin, Helianthus Annuus (Sunflower) Extract, Hydroxypropyl Starch Phosphate, Octenidine HCl, Oryza Sativa (Rice) Bran Extract, Phenoxyethanol, Polymethylsilsesquioxane, Propanediol, Propylene Glycol, Rosmarinus Officinalis (Rosemary) Leaf Extract, SD Alcohol 40-B, Silica, Sodium Hydroxide, Tocopherol, Xanthan Gum			

HAND ANTIPERSPIRANT DAILY USE

aluminum zirconium octachlorohydrate gly cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74307-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY (UNII: P9D3YP29MY) (ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY - UNII:P9D3YP29MY)		ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY	20 g in 100 mL

Inactive Ingredients		Strength
Ingredient Name		
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)		
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)		
COCONUT ALKANES (UNII: 1E5KJY107T)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
PHENOXYETHANOL (UNII: H1E492ZZ3T)		
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)		
XANTHAN GUM (UNII: TTV12P4NEE)		
PROPANEDIOL (UNII: 5965N8W85T)		
OCTENIDINE HYDROCHLORIDE (UNII: U84956NU4B)		
RICE BRAN (UNII: R60QEP13IC)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		

ROSEMARY (UNII: IJ67X351P9)	
TOCOPHEROL (UNII: R0ZB2556P8)	
GLYCERIN (UNII: PDC6A3C0OX)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
COCOYL CAPRYLOCAPRATE (UNII: 8D9H4QU99H)	
WATER (UNII: 059QF0K00R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74307-007-01	30 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	06/01/2020	

Labeler - Clutch Inc (080214892)

Establishment

Name	Address	ID/FEI	Business Operations
Wasatch Product Development, LLC.		962452533	manufacture(74307-007)