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

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.

Hand Antiperspirant Daily Use

Active Ingredient Purpose

Aluminum Zirconium Tetrachlorohydrex 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 (Suflower) 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

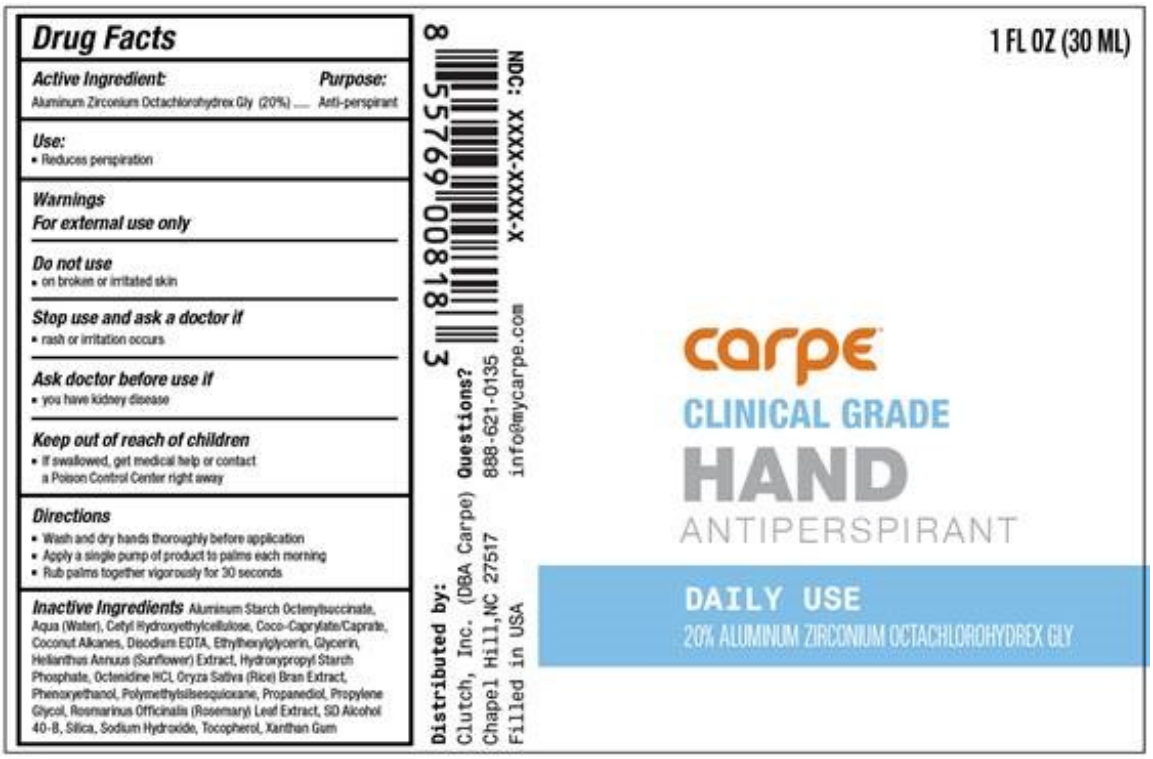
Carpe

Clinical Grade HAND

Antiperspirant

Daily Use

20% Aluminum Zirconium Octachlorohydrex GLY



HAND ANTIPERSPIRANT DAILY USE

aluminum zirconium octachlorohydrex 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 OCTACHLOROHYDREX GLY (UNII: P9D3YP29MY) (ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY - UNII:P9D3YP29MY)	ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY	20 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
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: 059QF0KO0R)	
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 final	part350	06/01/2020	

Labeler - Clutch Inc (080214892)

Establishment

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

Revised: 6/2020 Clutch Inc