HAND ANTIPERSPIRANT NIGHTLY USE- aluminum chloride hexahydrate 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 Nightly Use

Active Ingredient Purpose

Aluminum Chloride Hexahydrate (15%) Anti-perspirant

Use

- reduces perspiration

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

Stop use and ask a doctor 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 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 (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

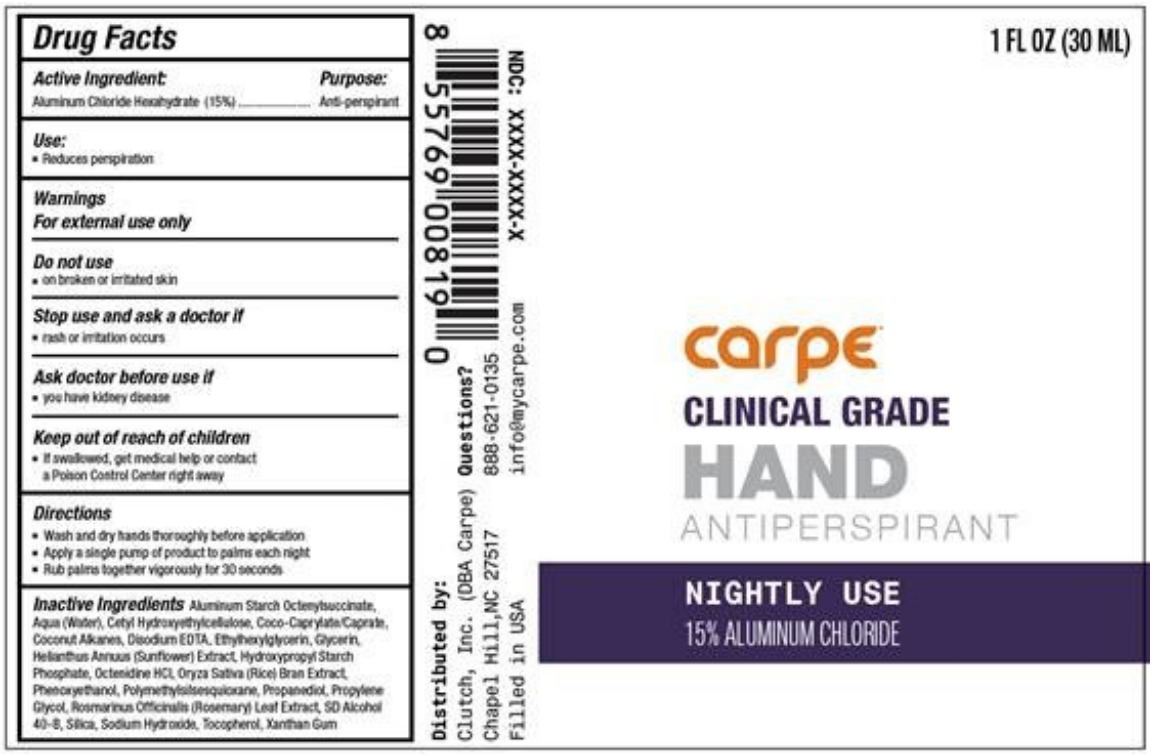
Carpe

Clinical Grade HAND

Antiperspirant

Nightly Use

15% Aluminum Chloride



HAND ANTIPERSPIRANT NIGHTLY USE

aluminum chloride hexahydrate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74307-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLORIDE (UNII: 3CYT62D3GA) (ALUMINUM CATION - UNII:3XHB1D032B)	ALUMINUM CHLORIDE	15 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
OCTENIDINE HYDROCHLORIDE (UNII: U84956NU4B)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
COCOYL CAPRYLOCAPRATE (UNII: 8D9H4QU99H)	
WATER (UNII: 059QF0KO0R)	
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)	
TOCOPHEROL (UNII: R0ZB2556P8)	
XANTHAN GUM (UNII: TTV12P4NEE)	

ROSEMARY (UNII: IJ67X351P9)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
RICE BRAN (UNII: R60QEP13IC)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPANEDIOL (UNII: 5965N8W85T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74307-008-01	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(74307-008)

Revised: 6/2020 Clutch Inc