CERTAIN DRI ANTIPERSPIRANT HAND-LOTION- aluminum chloride lotion
Clarion Brands, LLC

Drug Facts

Active ingredient

Aluminum chloride 15%

Purpose

Antiperspirant

Use

reduces perspiration

Warnings

For external use only

Do not use

- on broken or irritated skin
- immediately after shaving
- immediately after bathing

Ask a doctor before use if you have kidney disease

When using this product

do not use in or near eyes

Stop use and ask a doctor if rash or irritation develops

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 small, pea sized amount to the palms of your hands
- rub into palms vigorously for 15 seconds
- for best results, apply every night before bed and an additional two times per day for at least four weeks

Inactive ingredients

aloe barbadensis leaf juice, artemisia vulgarixs extract, behentrimonium methosulfate, bisaolol, bis-laurl cocaminopropylamine/HDI/PEG-100 copolymer, butylene glycol, cetearyl alcohol, cetyl alcohol, dimethicone, ethylhexyglycerin, hydroxyethylcellulose, jojoba esters, laminaria digitata extract, opuntia ficus-indica fruit extract, potassium hydroxide, saccharomyces cerevisiae extract, water, zingiber officinale (ginger) root extract

Questions?

call **1-844-923-7837**

PACKAGE LABEL

CERTAIN
DRI®
ANTIPERSPIRANT

QUICK DRY
SWEAT
GUARD

HAND
LOTION

CLINICALLY SHOWN
TO REDUCE SWEAT

NET WT 1.3 OZ (36 g)

Distributed by: Certain Dri LLC, 811 Broad Street, Suite 600
Chattanooga, TN 37402 ©2023

1000156

CERTAIN DRI ANTIPERSPIRANT HAND-LOTION

aluminum chloride lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69693-723
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Aluminum Chloride (UNII: 3CYT62D3GA) (Aluminum Cation -	Aluminum Chloride	.15 g

UNII:3XHB1D032B)	Aluminum Chloride	in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Aloe Vera Leaf (UNII: ZY81Z83H0X)	
Artemisia Vulgaris Root (UNII: 32MP823R8S)	
Behentrimonium Methosulfate (UNII: 5SHP745C61)	
Levomenol (UNII: 24WE03BX2T)	
Thiamine Bis-Laurylsulfate (UNII: 226A8HU328)	
Butylene Glycol (UNII: 3XUS85K0RA)	
Cetostearyl Alcohol (UNII: 2DMT128M1S)	
Cetyl Alcohol (UNII: 936JST6JCN)	
Dimethicone (UNII: 92RU3N3Y1O)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Hydroxyethyl Cellulose, Unspecified (UNII: T4V6TWG28D)	
Hydrolyzed Jojoba Esters (Acid Form) (UNII: UDR641JW8W)	
Laminaria Digitata (UNII: 15E7C67EE8)	
Opuntia Ficus-Indica Fruit Juice (UNII: 5ZC110ZY2H)	
Potassium Hydroxide (UNII: WZH3C48M4T)	
Saccharomyces Cerevisiae (UNII: 978D8U419H)	
Water (UNII: 059QF0KO0R)	
Ginger (UNII: C5529G5JPQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69693-723-13	1 in 1 CARTON	09/01/2024	
1		36 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	09/01/2024	

Labeler - Clarion Brands, LLC (079742703)

Revised: 7/2024 Clarion Brands, LLC