

CLINIQUE ANTIPERSPIRANT DEODORANT ROLL-ON- aluminum chlorohydrate liquid
CLINIQUE LABORATORIES 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.

ACTIVE INGREDIENT:ALUMINUM CHLOROHYDRATE 20.00%

USES: DECREASES UNDERARM PERSPIRATION

WARNINGS:

- FOR EXTERNAL USE ONLY
- DO NOT USE ON BROKEN SKIN
- STOP USE IF RASH OR IRRITATION OCCURS
- ASK A 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: APPLY TO UNDERARMS ONLY.

inactive ingredients: water [] urea [] silica [] alcohol denat. [] glycine [] aluminum chloride [] butyl stearate [] sodium lauroyl sarcosinate [] alcloxa [] benzethonium chloride [] trisodium edta [] methylparaben [] propylparaben [] butylparaben iln27821

PRINCIPAL DISPLAY PANEL:

CLINIQUE
anti-perspirant deodorant roll-on

ALUMINUM CHLOROHYDRATE

2.5FL OZ./ 70ML

CLINIQUE LABORATORIES, DIST.
NEW YORK, NY 10022

6183
CLINIQUE.COM

Drug Facts

Active ingredient	Purpose
Aluminum chlorohydrate 20%	Antiperspirant

Use Reduces underarm wetness.

Warnings
For external use only.

Do not use on broken skin.

Stop use if rash or irritation occurs.

Ask a 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.



Drug Facts (continued)

Directions
Apply to underarms only.

Inactive ingredients water (aqua purificata) purified, urea, silica, alcohol denat., glycols, aluminum chloride, butyl stearate, sodium lauroyl sarcosinate, alicoua, benzethonium chloride, trisodium octa, methylparaben, propylparaben, butylparaben [In27821]

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Made in U.K. 6JC4
clinique.com




CLINIQUE ANTIPERSPIRANT DEODORANT ROLL-ON

aluminum chlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49527-571
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROXYDRATE (UNII: HPN8MZW13M) (aluminum cation - UNII:3XHB1D032B)	ALUMINUM CHLOROXYDRATE	70 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49527-571-01	70 mL in 1 BOTTLE, WITH APPLICATOR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	12/01/1992	

Labeler - CLINIQUE LABORATORIES INC (173047747)**Establishment**

Name	Address	ID/FEI	Business Operations
ESTEE LAUDER COSMETICS, LTD		205952385	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
ESTEE LAUDER N.V.		370151326	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Len-Ron Manufacturing Division of Aramis Inc.		809771152	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Aramis Inc.		042918826	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Northtec Bristol		949264774	manufacture, relabel, repack

Establishment

Name	Address	ID/FEI	Business Operations
Northtec Keystone		618107429	manufacture, relabel, repack

Establishment

Name	Address	ID/FEI	Business Operations
PADC 1		110482184	manufacture, relabel, repack

Establishment

Name	Address	ID/FEI	Business Operations
Estee Lauder Pennsylvania Distribution Center 2		828534516	manufacture, relabel, repack

Establishment

Name	Address	ID/FEI	Business Operations
Estee Lauder Cosmetics, Ltd.		255175580	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Estee Lauder Cosmetics, Ltd		253616536	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Estee Lauder Cosmetics Distribution Center		208579636	manufacture, label, relabel

Establishment

Name	Address	ID/FEI	Business Operations
Estee Lauder Kabushiki Kaisha		712808195	relabel, repack

Establishment

Name	Address	ID/FEI	Business Operations
Whitman Laboratories Ltd.		216866277	manufacture

Establishment

Name	Address	ID/FEI	Business Operations
Aveda Corporation		071352058	manufacture