DR SWEAT- aluminum chloride liquid
Conopco Inc. d/b/a/ Unilever

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.

Dr Sweat Clinical Strength Antiperspirant

DR SWEAT CLINICAL STRENGTH ANTIPERSPIRANT - aluminum chloride liquid

Dr Sweat Clinical Strength Antiperspirant

Drug Facts

Active ingredient

Aluminum Chloride (15.0%)

Purpose

Antiperspirant

Uses

• reduces underarm perspiration

Warnings

- **For externaluse only.**
- **Do not use** on broken skin **.**
- **Ask a doctor before use if you have** kidney disease.
- **Stop use** if rash or irritation occurs.
- **Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• Apply to underarms only
• Apply to clean dry underarms before going to bed.

Inactive ingredients

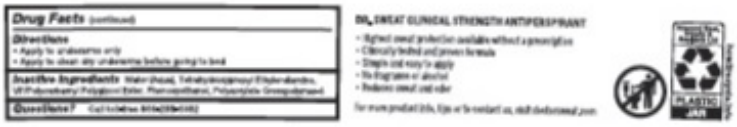
Water (Aqua), Tetrahydroxypropyl Ethylenediamine, VP/Polycarbamyl Polyglycol Ester, Phenoxyethanol, Polyacrylate Crosspolymer-6

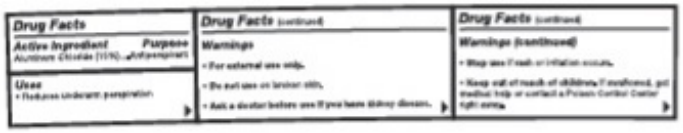
Questions?

Call toll-free 866-269-6082

- Highest sweat profession available without a prescription
- Clinically tested and proven formula
- Simply and easy to apply
- No fragrance or alcohol
- Reduces sweat and odor

Packaging





DR SWEAT

aluminum chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64942-1633
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 g

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
EDETOL (UNII: Q4R969U9FR)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1633-1	0.0284 g in 1 PACKET; Type 0: Not a Combination Product	03/29/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	03/29/2019	

Labeler - Conopco Inc. d/b/a/ Unilever (001375088)

Revised: 12/2022 Conopco Inc. d/b/a/ Unilever