

**ANTI-PERSPIRANT DEODORANT ROLL-ON UNSCENTED- aluminum
chlorohydrate liquid
Hydrox Laboratories**

**Anti-Perspirant Deodorant Roll-On Unscented
Drug Facts**

Active ingredient

Aluminum Chlorohydrate, 10% - Anhydrous Basis

Purpose

Antiperspirant

Use

Reduces underarm wetness.

Warnings

FOR EXTERNAL USE ONLY.

KEEP OUT OF REACH OF CHILDREN.

Do not use on broken skin.

Stop use if rash or irritation occurs.

Ask a doctor before use if you have kidney disease. If swallowed, get medical help or contact a Poison Control Center immediately. Use only as directed.

Directions

Apply to underarms. Use daily for best results.

Inactive ingredients

Purified Water, Hydroxyethylcellulose, Glycerin, Polysorbate 20, Tetrasodium EDTA

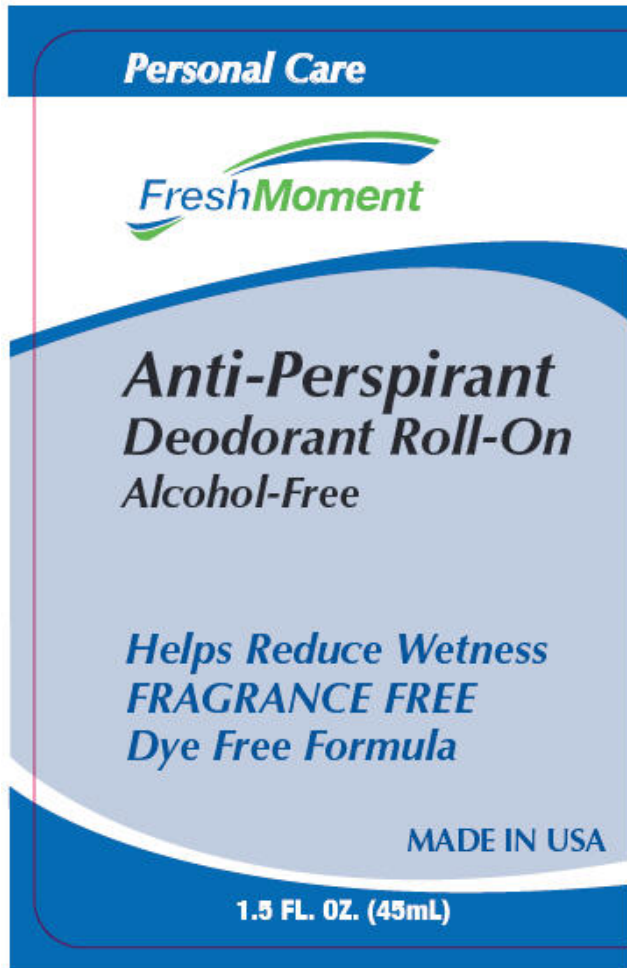
PRINCIPAL DISPLAY PANEL

Personal Care

FreshMoment

Anti-Perspirant

Deodorant Roll-On
 Alcohol Free
 Helps Reduce Wetness
 Fragrance Free
 Dye Free Formula
 Made in USA



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Purpose Antiperspirant	
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REF
K753C

MADE IN USA

MFG BY:
HYDROX LABORATORIES
825-B Tollgate Rd. • Elgin, IL 60123
1-07-K753C

0 21599 12332 1

ANTI-PERSPIRANT DEODORANT ROLL-ON UNSCENTED
 aluminum chlorohydrate liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10565-074
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZ W13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZ W13M)	ALUMINUM CHLOROHYDRATE	10 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10565-074-01	45 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	03/20/2012	
2	NDC:10565-074-02	59 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	03/20/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	03/20/2012	

Labeler - Hydrox Laboratories (025164302)

Registrant - Hydrox Laboratories (025164302)

Establishment

Name	Address	ID/FEI	Business Operations
Hydrox Laboratories		025164302	label(10565-074) , manufacture(10565-074) , pack(10565-074)

Revised: 10/2023

Hydrox Laboratories