ROLL ON ANTI PERSPIRANT ALCOHOL FREE- aluminum chlorohydrate solution
Hydrox Laboratories

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.

Anti-Perspirant
Deodorant Roll-On
Alcohol-Free

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, Fragrance, Tetrasodium EDTA

PRINCIPAL DISPLAY PANEL

Personal Care

*Fresh**Moment***

Anti-Perspirant
Deodorant Roll-On
Alcohol-Free

Helps Reduce Wetness

Spring Fresh Fragrance

MADE IN USA

1.5 FL. OZ. (45mL)

ROLL ON ANTI PERSPIRANT ALCOHOL FREE

aluminum chlorohydrate solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10565-066
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	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:10565-066-02	45 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	10/26/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part350	10/26/2011	

Labeler - Hydrox Laboratories (025164302)

Registrant - Hydrox Laboratories (025164302)

Establishment

Name	Address	ID/FEI	Business Operations
Hydrox Laboratories		025164302	MANUFACTURE(10565-066) , label(10565-066) , pack(10565-066)

Revised: 3/2020 Hydrox Laboratories