ANTI-PERSPIRANT DEODORANT- aluminum chlorohydrate spray
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 Spray

Drug Facts

Active ingredient

Aluminum Chlorohydrate, 13% - 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

Hold two inches from underarm and spray. Use daily for best results.

Inactive ingredients

Purified Water, Isopropyl Alcohol, Propylene Glycol, Peg-40 Hydrogenated Castor Oil, Fragrance.

MFG BY:
HYDROX LABORATORIES
825 Tollgate Rd. • Elgin, IL 60123

PRINCIPAL DISPLAY PANEL

Personal Care

FreshMoment

Anti-Perspirant
Deodorant Spray

Helps Reduce Wetness
Spring Fresh Fragrance

MADE IN USA

2 FL. OZ. (59mL)

ANTI-PERSPIRANT DEODORANT

aluminum chlorohydrate spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10565-067
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Isopropyl Alcohol (UNII: ND2M416302)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Polyoxyl 40 Hydrogenated Castor Oil (UNII: 7YC686GQ8F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10565-067-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/11/2011	
2	NDC:10565-067-04	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/11/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	11/11/2011	

Labeler - Hydrox Laboratories (025164302)

Registrant - Hydrox Laboratories (025164302)

Establishment

Name	Address	ID/FEI	Business Operations
Hydrox Laboratories		025164302	MANUFACTURE(10565-067) , label(10565-067) , pack(10565-067)

Revised: 3/2020 Hydrox Laboratories