

**HI AND DRI ANTIPERSPIRANT ROLL-ON UNSCENTED- aluminum chlorohydrate liquid
Revlon Consumer Products Corp**

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.

Hi & Dri Roll-On Antiperspirant - Unscented 1.7 fl. oz (50 ml)

Drug Facts

Active Ingredient

Aluminum chlorohydrate 18%

Purpose

Antiperspirant

Use

- Reduces underarm wetness

Warnings:

For external use 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.

Inactive Ingredients

aqua((water) eau), glyceryl stearate, laureth-23, magnesium aluminum silicate, laureth-4, lauric acid, cetearyl alcohol, behentrimonium methosulfate, edta

Questions

1-800-473-8566

Principal Display Panel - 1.7 fl oz bottle



◀ PEEL HERE

Drug Facts

Active Ingredient:
Aluminum Chlorohydrate 18%

Purpose:
Anti-Perspirant

Use:
• Reduces underarm wetness.

Drug Facts (continued)

Warnings:

- For external use only.
- Do not use on broken skin.
- Stop use if rash or irritation occurs.
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Directions:

- Apply to underarms only. ▶

Drug Facts (continued)

Inactive Ingredients: Aqua (Water) Eau, Glyceryl Stearate, Laureth-23, Magnesium Aluminum Silicate, Laureth-4, Lauric Acid, Behentrimonium Methosulfate, Cetearyl Alcohol, EDTA, 23291

QUESTIONS? 1-800-473-8566

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MADE IN SOUTH AFRICA. 216549-18 5122-04

HI AND DRI ANTIPERSPIRANT ROLL-ON UNSCENTED

aluminum chlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10967-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	0.18 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
LAURETH-23 (UNII: N72LMW566G)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
LAURETH-4 (UNII: 6HQ855798J)	
LAURIC ACID (UNII: 1160N9NU9U)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
BEHENTRIMONIUM METHO SULFATE (UNII: 5SHP745C61)	
EDETIC ACID (UNII: 9G34HU7RV0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10967-100-97	50 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	01/01/1982	

Labeler - Revlon Consumer Products Corp (788820165)

Establishment

Name	Address	ID/FEI	Business Operations
Revlon South Africa (PTY) Ltd		637155859	manufacture(10967-100)

Revised: 12/2013

Revlon Consumer Products Corp