**HI AND DRI ANTIPERSPIRANT ROLL-ON POWDER FRESH- 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.*

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**Hi & Dri Roll-On Antiperspirant - Powder Fresh 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 irriation 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.

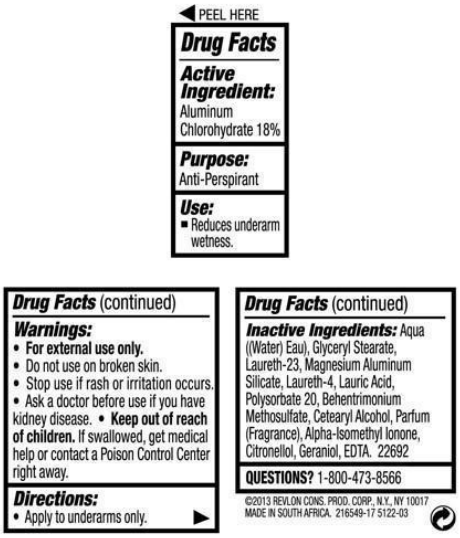
**Inactive Ingredients**

aqua((water) eau), glyceryl stearate, laureth-23, magnesium aluminum silicate, laureth-4, lauric acid, cetearyl alcohol, behentrimonium methosulfate, edta, alpha-isomethyl ionone, citronellol, limonene, hydroxycitronellol, benzyl alcohol, benzyl salicylate, linalool, citral

**Questions**

1-800-473-8566

**Principal Display Panel - 1.7 fl oz bottle**


# HI AND DRI ANTIPERSPIRANT ROLL-ON POWDER FRESH

aluminum chlorohydrate liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:10967-057 |
| **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 METHOSULFATE** (UNII: 5SHP745C61) | |
| **EDETIC ACID** (UNII: 9G34HU7RV0) | |
| **ISOMETHYL-.ALPHA.-IONONE** (UNII: 9XP4LC555B) | |
| **.BETA.-CITRONELLOL, (R)-** (UNII: P01OUT964K) | |

| | |
|---|---|
| **GERANIOL** (UNII: L837108USY) | |
| **LIMONENE, (+)-** (UNII: GFD7C86Q1W) | |
| **HYDROXYCITRONELLAL** (UNII: 8SQ0VA4YUR) | |
| **BENZYL ALCOHOL** (UNII: LKG8494WBH) | |
| **BENZYL SALICYLATE** (UNII: WAO5MNK9TU) | |
| **LINALOOL, (+)-** (UNII: F4VNO44C09) | |
| **CITRAL** (UNII: T7EU0O9VPP) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:10967-057-97 | 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(10967-057) |

Revised: 12/2013 Revlon Consumer Products Corp