HI AND DRI ROLL ON- aluminum chlorohydrate liquid 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

Aluminum chlorohydrate 18%

Purpose

Antiperspirant

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

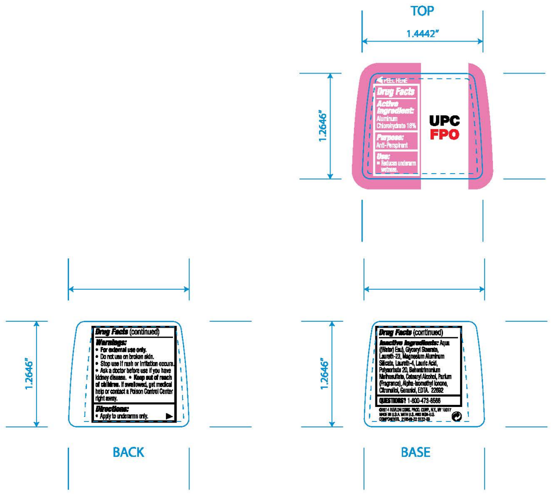
If swallowed, get medical help or contact a Poison Control Center right away.

Inactive Ingredients

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

Reduces underarm wetness

HI & DRI ROLL ON 1.7OZ POWDER FRESH
BACK LABEL
TR05967 / 5122-03
1.4442"W x 1.2646"H



ENLARGED 200% FOR PROOFREADING

◀ 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.
- Ask a doctor before use if you have kidney disease. • **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions:
- Apply to underarms only. ▶

Drug Facts (continued)

Inactive Ingredients: Aqua ((Water) Eau), Glyceryl Stearate, Laureth-23, Magnesium Aluminum Silicate, Laureth-4, Lauric Acid, Polysorbate 20, Behentrimonium Methosulfate, Cetearyl Alcohol, Parfum (Fragrance), Alpha-Isomethyl Ionone, Citronellol, Geraniol, EDTA. 22692

QUESTIONS? 1-800-473-8566

MADE IN U.S.A. WITH U.S. AND NON-U.S. COMPONENTS. 216549-22 5122-03

UPC: 309975122039

Mechanical Approval

Disk Release: 03-11-2014

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SCHAWK!
1600 East Sherwin Avenue
Des Plaines, IL 60018
Phone: 847-296-6000
800-621-1909
Fax: 847-296-9466

For support please contact the following:
Level 1: Customer Service: Susan Gojney
susan.gojney@schawk.com - 847.759.7045
Level 2: Customer Service: Tim Dawson
tim.dawson@schawk.com - 847.759.7070
Package Developer: Donna Lisowski
donna.lisowski@revlon.com - 732.287.7644

Client: Revlon **Brand:** Hi & Dri **Job #:** 525335A **Anthem #:** 40262
Description: Hi & Dri 1.7oz Roll On Powder Fresh Back Label
Component: Back Label **Program #:** **Project #:** TR05967
Stock #: 216549-22 **MLI #:** 22692 **Product #:** 5122-03
Size: 1.4442"W x 1.2646"H **Die #:**
File Name: REV_525335A02_TR05967.ai **Software:** 15
Op/Date: spgao 03-3-2014, ZCL 03-11-2014

QC1____
QC2____
QC3____

100%	75%	50%	25%	10%	3%	2%	1%	Color
								PMS211
								Black
								DIESTRIKE
								REVLON CALLOUTS

ARTWORK COMPONENTS

CHECKED BY:	Copy	Trademarks	Registration Marks	Weight Claim	Die Vinyls/ Blister Placement	Int'l Symbols	UPC #	UPC Size/Loc	Ctry Origin	Color Breaks	Component Numbers	Product Numbers	Merch Codes	Claims	MLI's	Shades	Label Pos	Translations	Make-up Credits (displays)	Pricing (WM displays)
MARKETING																				
LEGAL																				
PACKAGING DEVELOPMENT																				
R&D																				

Reason for change:
1. **Release** Date: 03-11-14
2. **Change description** Date: 00-00-00
3. **Change description** Date: 00-00-00
4. **Change description** Date: 00-00-00

SUBSTRATE IS WHITE

BACKGROUND PRINTS:
100% PMS 211

BLACK TEXT, RULES & UPC PRINT:
100% BLACK

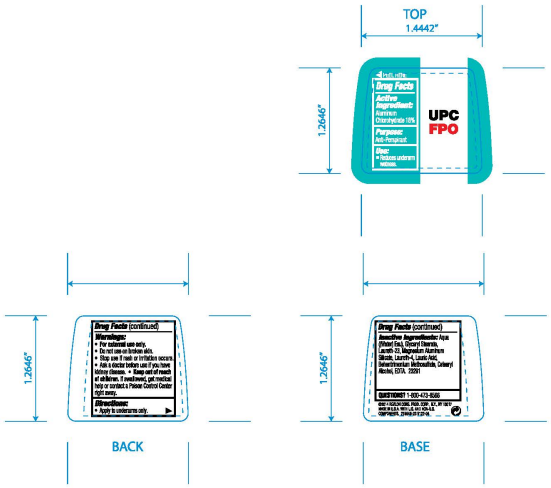
WHITE TEXT & RULES:
K/O WHITE

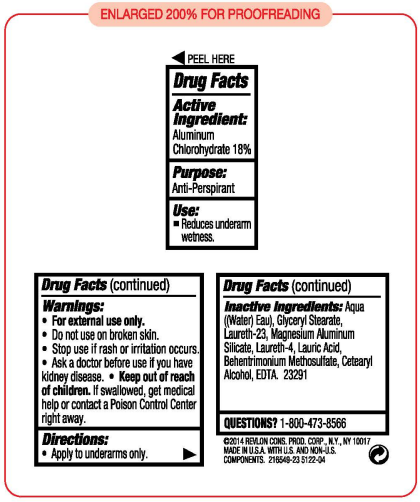
CYAN RULES REPRESENT DIE AND MEASUREMENTS:
DO NOT PRINT

CYAN DASHED RULES REPRESENT LIVE AREA:
DO NOT PRINT

Hi & Dri Roll-On Antiperspirant

HI & DRI ROLL ON 1.7OZ UNSCENTED
BACK LABEL
TR05968 / 5122-04
1.4442"W x 1.2646"H





HI AND DRI ROLL ON

aluminum chlorohydrate liquid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10967-602
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)	
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)	
LINALOOL, (+)- (UNII: F4VNO44C09)	
CITRAL (UNII: T7EU0O9VPP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
LAURETH-23 (UNII: N72LMW566G)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10967-602-17	50 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	01/01/2014	

HI AND DRI ROLL ON

aluminum chlorohydrate liquid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10967-600
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)	
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)	
LINALOOL, (+)- (UNII: F4VNO44C09)	
CITRAL (UNII: T7EU0O9VPP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
LAURETH-23 (UNII: N72LMW566G)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
HYDROXYCITRONELLAL (UNII: 8SQ0VA4YUR)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10967-600-17	50 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	01/01/2014	

Labeler - Revlon Consumer Products Corp (788820165)

Registrant - REVLON, INC. (188442578)

Establishment

Name	Address	ID/FEI	Business Operations
Revlon South Africa (PTY) Ltd		637155859	manufacture(10967-600, 10967-602)