

**MITCHUM ROLL ON- aluminum zirconium tetrachlorohydrex gly liquid**  
**Revlon Consumer Products Corp**

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**Mitchum Roll-On**

Aluminum Zirconium Tetrachlorohydrex Gly 20%

**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 irritation occurs

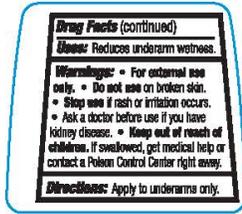
If swallowed, get medical help or contact a Poison Control Center right away.

**Inactive Ingredients**

AQUA ((WATER) EAU), GLYCERYL STEARATE, LAURETH-23, MAGNESIUM ALUMINUM SILICATE, POLYSORBATE 20, PARFUM (FRAGRANCE), LAURETH-4, LAURIC ACID, CETEARYL ALCOHOL, EDTA, BENHENTRIMONIUM METHOSULFATE, BENZYL SALICYLATE, LINALOOL, SILICA DIMETHICONE SILYLATE, LIMONENE, HEXYL CINNAMAL, TOCOPHERYL ACETATE, ALOE BARBADENSIS LEAF EXTRACT, COUMARIN, HYDROGEN PEROXIDE, CITRONELLOL, GERANIOL

Reduces underarm wetness

Mitchum Roll On



BACK



BASE

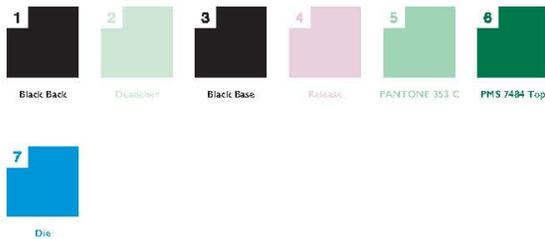


TOP

CUST. PART # 216549-30  
 ART # 93248-024A

PROOF INFORMATION

INKS USED ON JOB



**Customer:** REVLON Inc.

**Proof Date:** 1/21/15

**Size:** 1.2646" x 1.4442"

**Corner Radius:** N/A

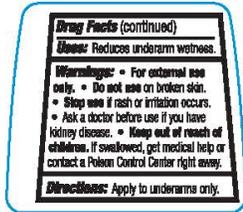
**Die Margin(printable area):** 1/16" WITHIN DIE LINE

**Substrate:** WHITE

**NOTES:**

Full Matte Varnish

**CCL LABEL**  
 YOUR GLOBAL PARTNER



BACK



BASE



TOP

CUST. PART # 216549-30  
 ART # 93248-024A

PROOF INFORMATION						
INKS USED ON JOB						
1	2	3	4	5	6	
Black Back	Darkblue	Black Base	Redlake	PANTONE 353 C	PMS 7494 Top	
7						
Die						
<b>Customer:</b> REVLON Inc. <b>Proof Date:</b> 1/21/15 <b>Size:</b> 1.2646" x 1.4442" <b>Corner Radius:</b> N/A <b>Die Margin(printable area):</b> 1/16" WITHIN DIE LINE <b>Substrate:</b> WHITE <b>NOTES:</b> Full Matte Varnish						<b>CCL LABEL</b> YOUR GLOBAL PARTNER

# MITCHUM ROLL ON

aluminum zirconium tetrachlorohydrate gly liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:10967-601
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ALUMINUM ZIRCONIUM TETRACHLOROXYDRATE GLY</b> (UNII: 80386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROXYDRATE GLY - UNII:80386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROXYDRATE GLY	0.2 g in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>WATER</b> (UNII: 059QF0K00R)	
<b>GLYCERYL STEARATE SE</b> (UNII: FCZ5MH785I)	
<b>LAURETH-23</b> (UNII: N72LMW566G)	
<b>MAGNESIUM ALUMINUM SILICATE</b> (UNII: 6M3P64V0NC)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>LAURETH-4</b> (UNII: 6HQ855798J)	
<b>LAURIC ACID</b> (UNII: 1160N9NU9U)	
<b>EDETIC ACID</b> (UNII: 9G34HU7RV0)	
<b>BENZYL SALICYLATE</b> (UNII: WAO5MKN9TU)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>.BETA.-CITRONELLOL, (R)-</b> (UNII: P01OUT964K)	
<b>GERANIOL</b> (UNII: L837108USY)	
<b>COUMARIN</b> (UNII: A4VZ22K1WT)	
<b>HYDROGEN PEROXIDE</b> (UNII: BBX060AN9V)	
<b>LIMONENE, (+/-)-</b> (UNII: 9MC3I34447)	
<b>LINALOOL, (+/-)-</b> (UNII: D81QY6I88E)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10967-601-34	100 mL in 1 CONTAINER; Type 0: Not a Combination Product	01/01/2014	
2	NDC:10967-601-17	50 mL in 1 CONTAINER; 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 Drug	M019	01/01/2014	

# MITCHUM ROLL ON

aluminum zirconium tetrachlorohydrate gly liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:10967-609
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY</b> (UNII: 8O386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY - UNII:8O386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY	0.2 g in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERYL STEARATE SE</b> (UNII: FCZ5MH785I)	
<b>LAURETH-23</b> (UNII: N72LMW566G)	
<b>MAGNESIUM ALUMINUM SILICATE</b> (UNII: 6M3P64V0NC)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>LAURETH-4</b> (UNII: 6HQ855798J)	
<b>LAURIC ACID</b> (UNII: 1160N9NU9U)	
<b>EDETIC ACID</b> (UNII: 9G34HU7RV0)	
<b>BENZYL SALICYLATE</b> (UNII: WAO5MKN9TU)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>.BETA.-CITRONELLOL, (R)-</b> (UNII: P01OUT964K)	
<b>GERANIOL</b> (UNII: L837108USY)	
<b>COUMARIN</b> (UNII: A4VZ22K1WT)	
<b>HYDROGEN PEROXIDE</b> (UNII: BBX060AN9V)	
<b>LIMONENE, (+/-)-</b> (UNII: 9MC3I34447)	
<b>LINALOOL, (+/-)-</b> (UNII: D81QY6I88E)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10967-609-17	50 mL in 1 CONTAINER; 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 Drug	M020	01/01/2014	12/31/2024

**Labeler** - Revlon Consumer Products Corp (788820165)

**Establishment**

Name	Address	ID/FEI	Business Operations
REVLON, INC.		809725570	manufacture(10967-601, 10967-609)

Revised: 12/2024

Revlon Consumer Products Corp