WOMENS MITCHUM ADVANCED INVISIBLE ROLL-ON ANTIPERSPIRANT DEODORANT POWDER FRESH- aluminum zirconium tetrachlorohydrex gly 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.

Womens Mitchum Advanced Roll On Antiperspirant Deodorant

Drug Facts

Active Ingredient

Aluminum zirconium tetrachlorohydrex gly 20%

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, polysorbate 20, tocopheryl acetate, aloe barbadensis leaf extract, hydrogen peroxide, silica dimethicone silylate, laureth-4, behentrimonium methosulfate, cetearyl alcohol, lauric acid, EDTA, fragrance, benzyl salicylate, linalool, limonene, hexyl cinnamal, coumarin, citronellol, geraniol

Questions

1-888-8-MITCHUM

Principal Display Panel - 1.7 fl oz bottle

Inside

Base

WOMENS MITCHUM ADVANCED INVISIBLE ROLL-ON ANTIPERSPIRANT DEODORANT POWDER FRESH

aluminum zirconium tetrachlorohydrex gly liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10967-583
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY (UNII: 8O386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY - UNII:8O386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY	0.20 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)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)	
LAURETH-4 (UNII: 6HQ855798J)	
BEHENTRIMONIUM METHOSULFATE (UNII: 5SHP745C61)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
LAURIC ACID (UNII: 1160N9NU9U)	
EDETIC ACID (UNII: 9G34HU7RV0)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
LINALOOL, (+)- (UNII: F4VNO44C09)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
COUMARIN (UNII: A4VZ22K1WT)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
GERANIOL (UNII: L837108USY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10967-583-97	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	12/15/2013	

Labeler - Revlon Consumer Products Corp (788820165)

Establishment

Name	Address	ID/FEI	Business Operations
Revlon South Africa (PTY) Ltd		637155859	manufacture(10967-583)

Revised: 12/2013 Revlon Consumer Products Corp