

**MITCHUM CLINICAL SOFT SOLID POWDER FRESH- aluminum zirconium tetrachlorohydrate gly 20.0% stick  
Revlon Consumer Products Corp.**

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**Mitchum Clinical Soft Solid - Powder Fresh**

**Active Ingredients**

ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY 20.0%

**Keep out of reach of children**

If swallowed, get medical help or call Poison Control Center right away

**Inactive Ingredient section**

Cyclopentasiloxane; Tribehenin; Dimethicone; Petrolatum; Perfume (Fragrance);  
Trisiloxane; C18-36 Acid Triglyceride; Aloe Barbadensis Leaf Extract; Tocopheryl Acetate;  
Sodium Starch Octenylsuccinate, Citric Acid, BHT, Sodium Ascorbate,

Calcium Disodium EDTA, Hydrated Silica, Hexyl Cinnamal, Limonene, Linalool, Mannitol,

Benzyl Salicylate, Alpha-Isomethyl Ionone, Citronellol, Hydroxycitronellal, Benzyl  
Benzoate,

Geraniol, Cinnamyl Alcohol.

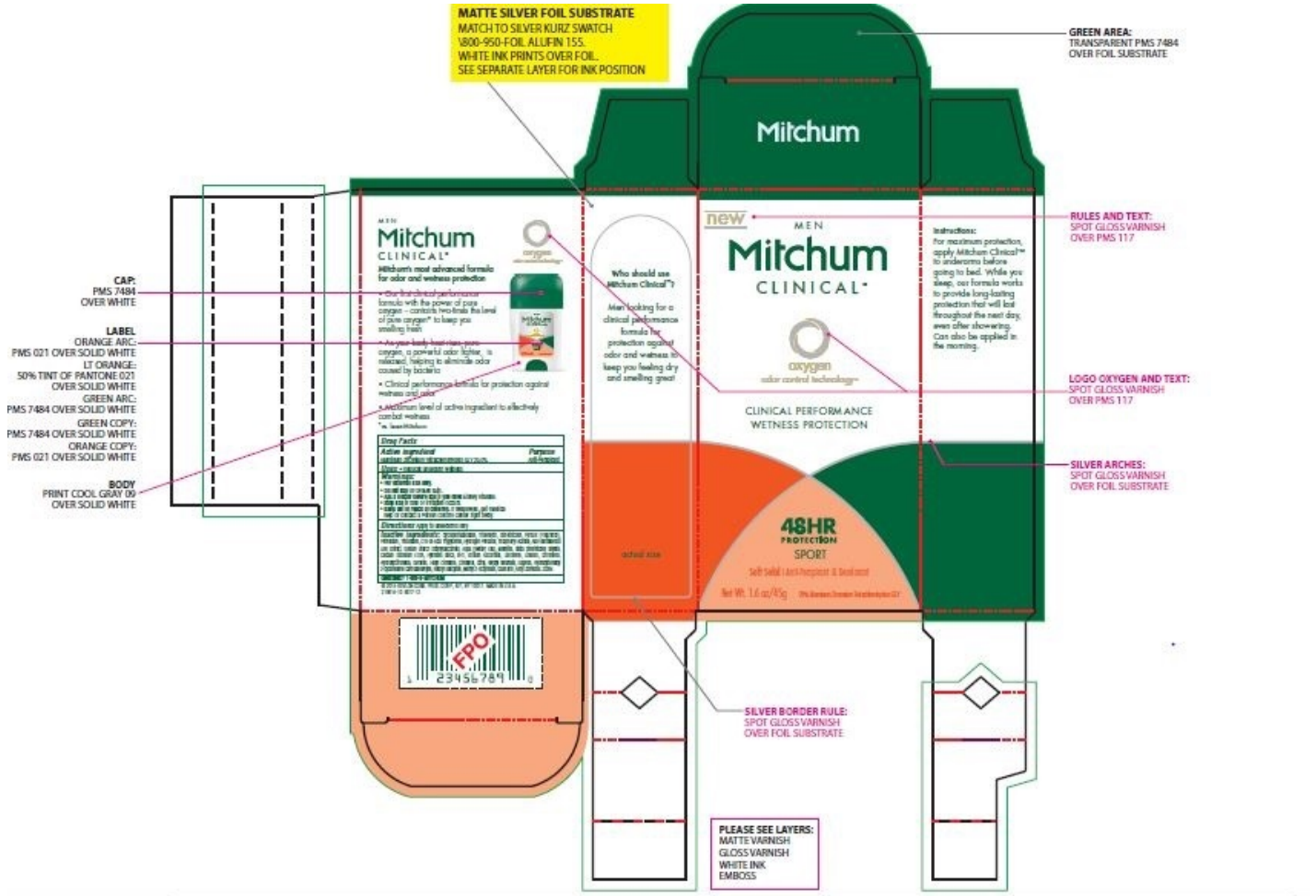
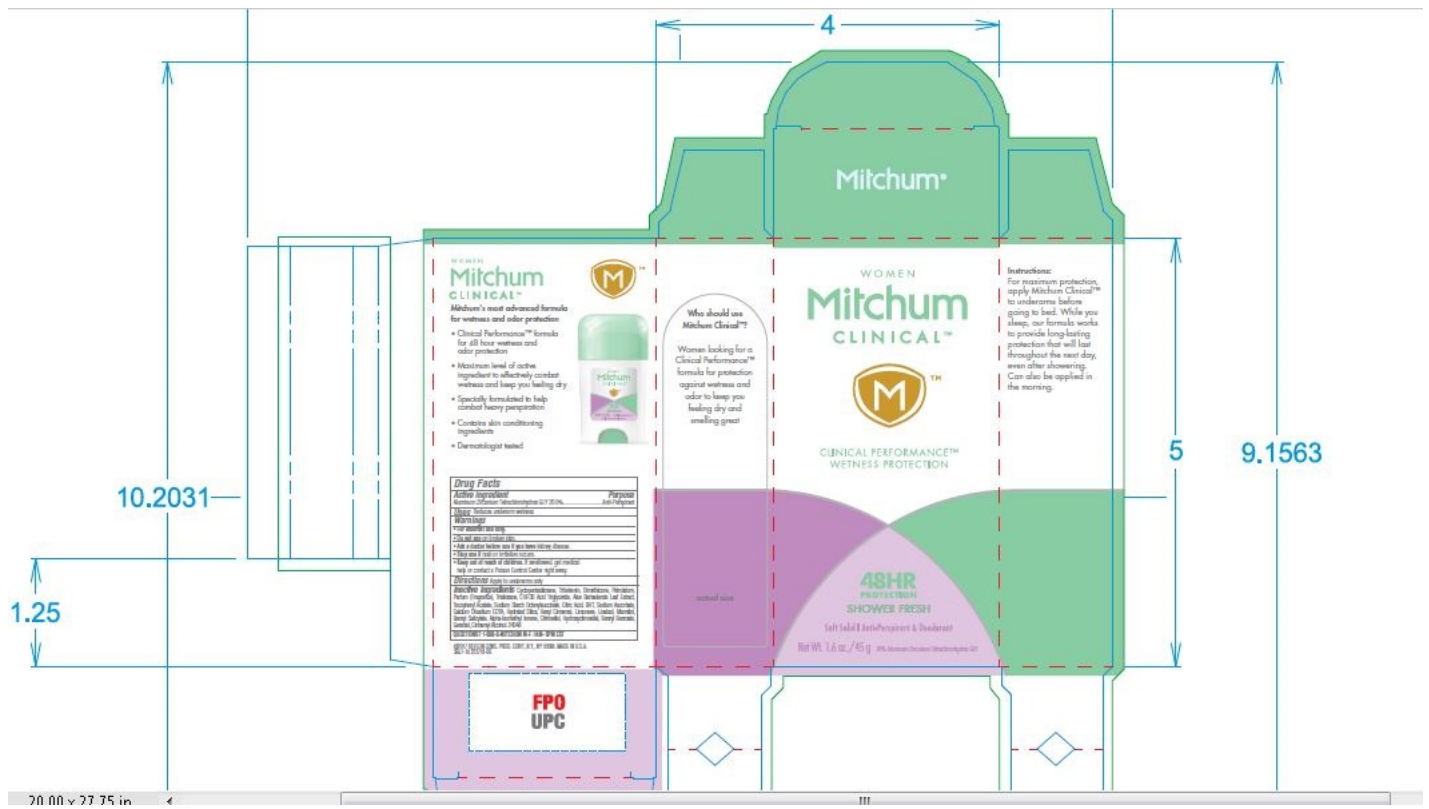
**Uses**

Reduces Underarm Wetness

Apply to underarms only

- 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

Reduces Underarm wetness



**MITCHUM CLINICAL SOFT SOLID POWDER FRESH**  
aluminum zirconium tetrachlorohydrate 20.0% stick

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:10967-678
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)	
<b>TRIBEHENIN</b> (UNII: 8OC9U7TQZ0)	
<b>C18-36 ACID TRIGLYCERIDE</b> (UNII: ZRA72DR3R7)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>ALPHA-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>EDETATE CALCIUM DISODIUM ANHYDROUS</b> (UNII: 8U5D034955)	
<b>SODIUM STARCH GLYCOLATE TYPE A CORN</b> (UNII: AG9B65PV6B)	
<b>HYDRATED SILICA</b> (UNII: Y6O7T4G8P9)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>CYCLOPENTASILOXANE</b> (UNII: 0THT5PCI0R)	
<b>TRISILOXANE</b> (UNII: 9G1ZW13R0G)	
<b>CITRIC ACID</b> (UNII: 2968PHW8QP)	
<b>SODIUM ASCORBATE</b> (UNII: S033EH8359)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:10967-678-16	45 g in 1 CANISTER; Type 0: Not a Combination Product	10/01/2018	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M019	10/01/2018	

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

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
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REVLON, INC.

809725570

manufacture(10967-678)

Revised: 12/2024

Revlon Consumer Products Corp.