

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

Mitchum Clinical Soft Solid - Pure 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.

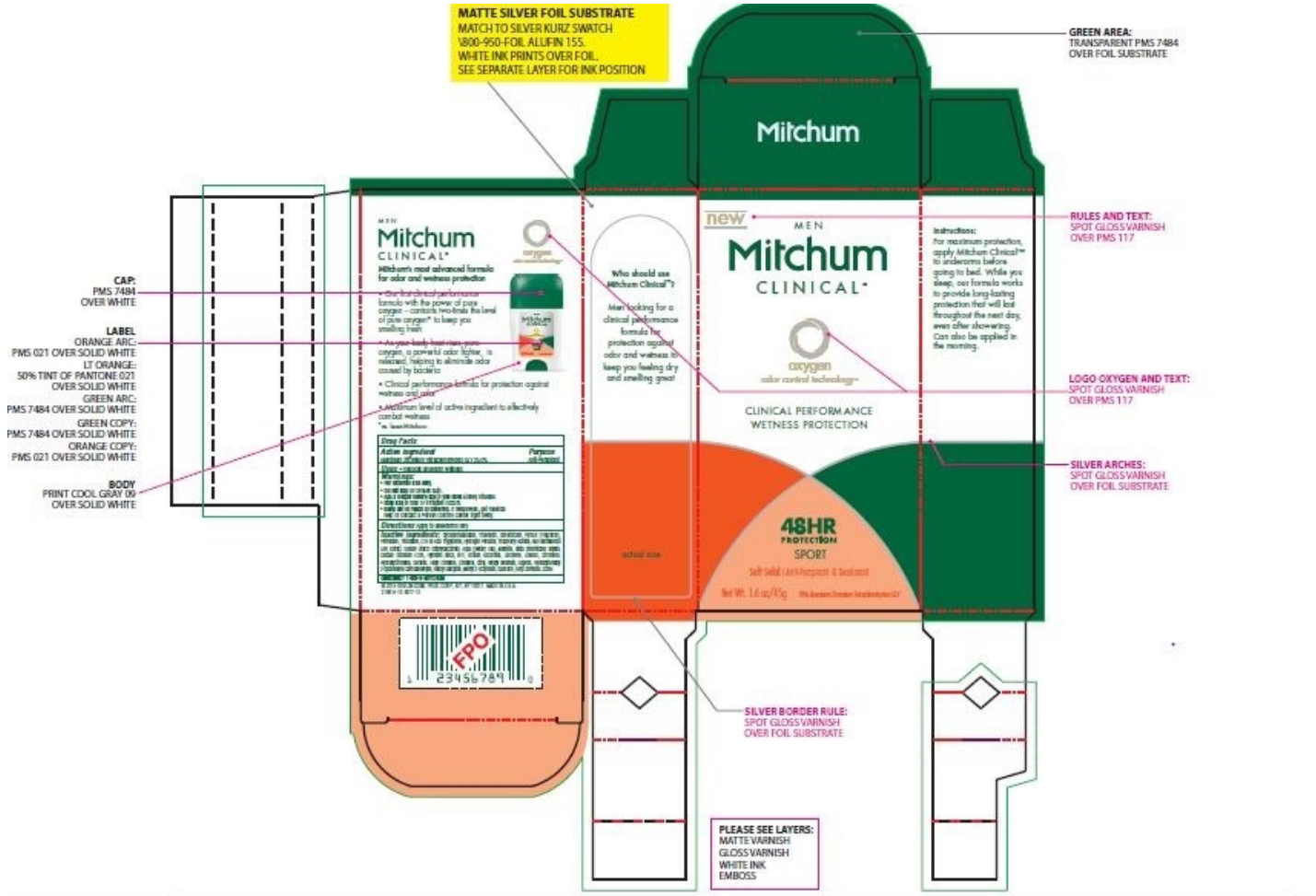
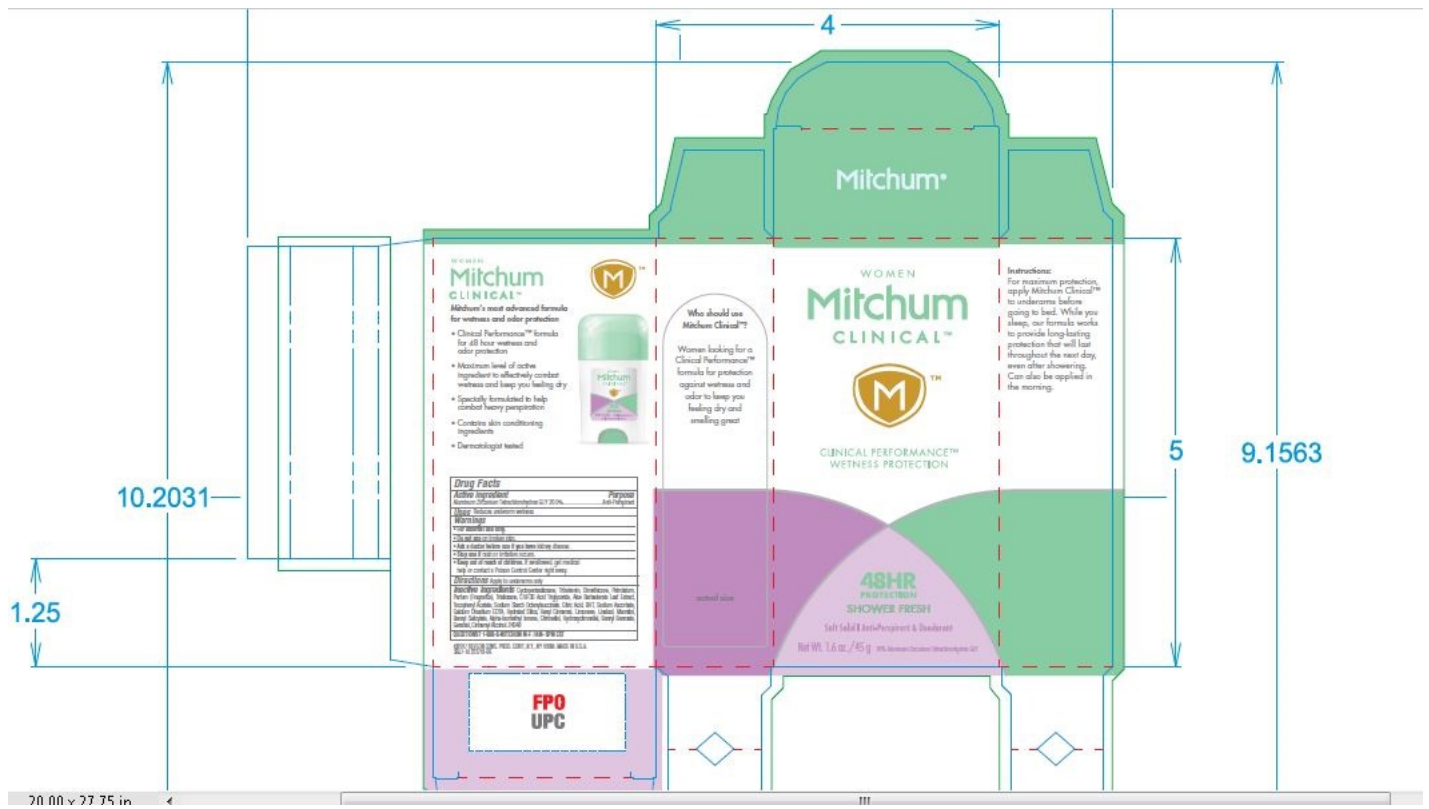
Uses

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 PURE FRESH
 aluminum zirconium tetrachlorohydrate gly 20.0% stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10967-680
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10967-680-16	45 g in 1 CANISTER; Type 0: Not a Combination Product	10/01/2018	

Marketing Information

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

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

Name	Address	ID/FEI	Business Operations
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REVLON, INC.

809725570

manufacture(10967-680)

Revised: 12/2024

Revlon Consumer Products Corp.