MITCHUM ULTIMATE- 72 HOUR PROTECTION - CLEAN CONTROL - APDEO- antiperspirant and deodorant stick
Revlon Consumer Products Corp.

Mitchum Ultimate - 72 hour protection - Clean Control - APDEO

Active Ingredient

Aluminum Zirconium Tetrachlorohydrex GLY 20%

Purpose

Antiperspirant

Uses

Reduces underam wetness

Warnings

For external use only.

Do Not Use

Do not use on broken skin.

Stop Use

Stop use if rash or irritation occurs.

Ask a doctor

Ask a doctor before use if you have kidney disease.

Keep out of reach of children

Keep out of reach of children, if swallowed, seek medical attention or conduct a poison center right away.

Directions

Directions Apply to underarms only.

Inactive ingredients

Cyclopentasiloxane, Stearyl Alcohol, C12-15 Alkyl Benzoate, PPG-14 Butyl Ether, Phenyl Trimethicone, Petrolatum, Hydrogenated Castor Oil, Parfum (Fragrance), Talc, BHT, Calcium Disodium EDTA, Citric Acid, Hydrated Silica, Mannitol, Sodium Ascorbate, Sodium Starch Octenylsuccinate

MITCHUM ULTIMATE- 72 HOUR PROTECTION - CLEAN CONTROL - APDEO

antiperspirant and deodorant stick

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10967-791
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	15.3653 g in 76 g

Inactive Ingredients

Ingredient Name	Strength
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
PETROLATUM (UNII: 4T6H12BN9U)	
CALCIUM DISODIUM EDTA (UNII: 8U5D034955)	
MODIFIED CORN STARCH (1-OCTENYL SUCCINIC ANHYDRIDE) (UNII: 461P5CJN6T)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
MANNITOL (UNII: 3OWL53L36A)	
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	
C12-15 ALKYL BENZOATE (UNII: A9EJ3J61HQ)	
CITRIC ACID (UNII: 2968PHW8QP)	
TALC (UNII: 7SEV7J4R1U)	
BHT (UNII: 1P9D0Z171K)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
SODIUM ASCORBATE (UNII: S033EH8359)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10967-791-27	76 g in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	10/17/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	10/17/2025	

Labeler - Revlon Consumer Products Corp. (788820165)

Registrant - Revlon, Inc. (188442578)

Establishment

Name	Address	ID/FEI	Business Operations
Revlon, Inc.		809725570	manufacture(10967-791)

Revised: 6/2025 Revlon Consumer Products Corp.