

**MITCHUM ULTIMATE- 72 HOUR PROTECTION - WILD ENERGY SOLID - APDEO-
antiperspirant and deodorant stick
Revlon Consumer Products Corp.**

Mitchum Ultimate - 72 hour protection - Wild Energy Solid - APDEO

Active Ingredient

Aluminum Zirconium Tetrachlorohydrex GLY 20%

Purpose

Antiperspirant

Uses

Reduces underarm 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

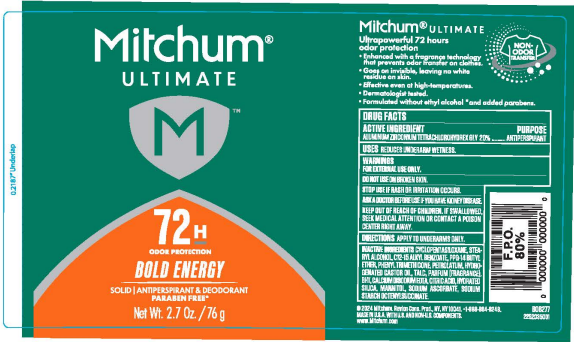
Keep out of reach of children, if swallowed, seek medical attention or conduct a poison center right away.

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



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antiperspirant and deodorant stick

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10967-790
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY (UNII: 80386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY - UNII:80386558JE)		ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY	20.2 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:10967-790-27	76 g in 1 BOTTLE, WTH 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(10967-790)

Revised: 6/2025

Revlon Consumer Products Corp.