

AXE- black 48h fresh and dry antiperspirant stick
Conopco Inc. d/b/a/ Unilever

Axe Black 48H Fresh & Dry Antiperspirant

AXE BLACK 48H FRESH & DRY ANTIPERSPIRANT - aluminum zirconium tetrachlorohydrate gly stick

Axe Black 48H Anti Sweat Antiperspirant

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrate GLY (19.0 %)

Purpose

antiperspirant

Uses

- reduces underarm wetness
- 48 Hour Protection

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 irritation 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

Isopropyl Palmitate, Stearyl Alcohol, Cyclopentasiloxane, PPG-14 Butyl Ether, Mineral Oil, Talc, Hydrogenated Castor Oil, Fragrance (Parfum), Steareth-100, BHT.

Questions?

Call toll-free 1-800-450-7580



AXE

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-2070
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	19 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
TALC (UNII: 7SEV7J4R1U)	

CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
MINERAL OIL (UNII: T5L8T28FGP)	
STEARETH-100 (UNII: 4OH5W9UM87)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-2070-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	10/22/2022	
2	NDC:64942-2070-2	2 in 1 PACKAGE	10/22/2022	
2	NDC:64942-2070-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product		
3	NDC:64942-2070-3	2 in 1 PACKAGE	10/22/2022	
3	NDC:64942-2070-2	2 in 1 PACKAGE		
3	NDC:64942-2070-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	10/22/2022	

Labeler - Conopco Inc. d/b/a/ Unilever (001375088)

Revised: 11/2024

Conopco Inc. d/b/a/ Unilever