

**AXE- essence black pepper and cedarwood scent 48h fresh and dry antiperspirant stick
Conopco Inc. d/b/a/ Unilever**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Axe Essence Black Pepper & Cedarwood Scent 48H Fresh & Dry Antiperspirant

AXE ESSENCE BLACK PEPPER & CEDARWOOD SCENT 48H FRESH & DRY ANTIPERSPIRANT - aluminum zirconium tetrachlorohydrate gly stick

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 1-800-761-3683

Packaging



AXE

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-2122
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY (UNII: 80386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY - UNII:80386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY	19 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
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)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-2122-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	07/19/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	07/03/2023	

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

Revised: 7/2023

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