

AXE- black antiperspirant and deodorant 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 Black 48H Dry Antiperspirant and Deodorant

AXE BLACK 48H DRY ANTIPERSPIRANT AND DEODORANT - aluminum zirconium tetrachlorohydrate stick

Axe Black 48H Dry Antiperspirant and Deodorant

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrate GLY (18.2 %)

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

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

Questions?

Call toll-free 1-800-450-7580

Packaging



AXE

black antiperspirant and deodorant stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1589
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM TETRACHLORO HYDREX GLY (UNII: 8O386558JE) (ALUMINUM ZIRCONIUM TETRACHLORO HYDREX GLY - UNII:8O386558JE)	ALUMINUM ZIRCONIUM TETRACHLORO HYDREX GLY	18.2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
TALC (UNII: 7SEV7J4R1U)	
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

STEARYL ALCOHOL (UNII: 2KR89I4H1Y)				
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)				
MINERAL OIL (UNII: T5L8T28FGP)				
STEARETH-100 (UNII: 4OH5W9UM87)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1589-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	01/01/2020	
2	NDC:64942-1589-2	2 in 1 PACKAGE	01/01/2020	
2	NDC:64942-1589-1	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 final		part350	01/01/2020	

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

Revised: 11/2020

Conopco Inc. d/b/a/ Unilever