

**AXE APOLLO ANTIPERSPIRANT AND DEODORANT- aluminum zirconium tetrachlorohydrate gly 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 Apollo Antiperspirant and Deodorant

AXE APOLLO ANTIPERSPIRANT AND DEODORANT - aluminum zirconium tetrachlorohydrate gly stick

Axe Apollo Antiperspirant and Deodorant

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrate GLY (18.2 %)

Purpose

antiperspirant

Uses

- reduces underarm wetness
- 24 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, Stearyl Alcohol, PPG-14 Butyl Ether, Hydrogenated Castor Oil, Talc, PEG-8 Distearate, Fragrance (Parfum), BHT.

Questions?

Call toll-free 1-800-450-7580

Packaging



AXE APOLLO ANTIPERSPIRANT AND DEODORANT

aluminum zirconium tetrachlorohydrate gly stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1532
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	18.2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
PEG-8 DISTEARATE (UNII: 7JNC8VN07M)	
TALC (UNII: 7SEV7J4R1U)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1532-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	09/21/2017	
2	NDC:64942-1532-2	2 in 1 PACKAGE	09/21/2017	
2	NDC:64942-1532-1	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	09/21/2017	

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

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Conopco Inc. d/b/a/ Unilever