

**AXE NIGHT 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 Night Antiperspirant and Deodorant

AXE NIGHT ANTIPERSPIRANT AND DEODORANT - aluminum zirconium tetrachlorohydrate gly stick

Axe Night Antiperspirant and Deodorant

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrate GLY (11.4 %)

Purpose

antiperspirant

Uses

- reduces underarm wetness

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, PPG-14 Butyl Ether, Stearyl Alcohol, Polyethylene, Hydrogenated Castor Oil, PEG-8 Distearate, Fragrance (Parfum), Silica, BHT.

Questions?

Call toll-free **1-800-450-7580**

Packaging



AXE NIGHT ANTIPERSPIRANT AND DEODORANT

aluminum zirconium tetrachlorohydrate gly stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1542
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	11.4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
PEG-8 DISTEARATE (UNII: 7JNC8VN07M)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1542-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	09/27/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	09/27/2017	

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

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