

**AXE SIGNATURE NIGHT ANTIPERSPIRANT 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 Signature Night Antiperspirant Deodorant

Active ingredient

Aluminum Zirconium Tetrachlorohydrate GLY(15.2%)

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, Stearyl Alcohol, C12-15 Alkyl Benzoate, PPG-14 Butyl Ether, Hydrogenated Castor Oil, Fragrance (Parfum), Dimethicone, Polyethylene, Helianthus Annuus (Sunflower) Seed Oil, Steareth-100, BHT.

Questions?

Call toll-free 1-800-450-7580

2.7 oz PDP



AXE SIGNATURE NIGHT ANTIPERSPIRANT DEODORANT

aluminum zirconium tetrachlorohydrax gly stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1421
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Aluminum Zirconium Tetrachlorohydrax GLY (UNII: 8O386558JE) (Aluminum Zirconium Tetrachlorohydrax GLY - UNII:8O386558JE)	Aluminum Zirconium Tetrachlorohydrax GLY	15.2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PEG-8 DISTEARATE (UNII: 7JNC8VN07M)	
SUNFLOWER OIL (UNII: 3W1JG795Y)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64942-1421-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product		
2	NDC:64942-1421-2	14 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	12/01/2015	

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

Establishment

Name	Address	ID/FEI	Business Operations
Unilever Raeford HPCNA		131411576	manufacture(64942-1421)

Establishment

Name	Address	ID/FEI	Business Operations
Zolberg Corporation		720166446	manufacture(64942-1421)

Revised: 7/2015

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