

AXE SIGNATURE ISLAND ANTIPERSPIRANT- aluminum zirconium terachlorohydrex 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 Island Antiperspirant

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrex 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**

48HR ANTI MARKS PROTECTION

ANTI WHITE MARKS

ANTI YELLOW STAINS

KEEPS YOU (AND YOUR SHIRTS) FEELING CLEAN AND FRESH.

Packaging



AXE SIGNATURE ISLAND ANTIPERSPIRANT

aluminum zirconium tetrachlorohydrate gly stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64942-1473
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY (UNII: 8O386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY - UNII:8O386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY	11.4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	

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:64942-1473-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	10/19/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	10/19/2016	

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

Revised: 10/2016

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