AXE- black eclipse 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 Eclipse 48H Dry Antiperspirant and Deodorant

AXE BLACK ECLIPSE 48H DRY ANTIPERSPIRANT AND DEODORANT - aluminum zirconium tetrachlorohydrex gly stick

Axe Black Eclipse 48H Dry Antiperspirant and Deodorant

Drug Facts

Active ingredient

Aluminum Zirconium Tetrachlorohydrex 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

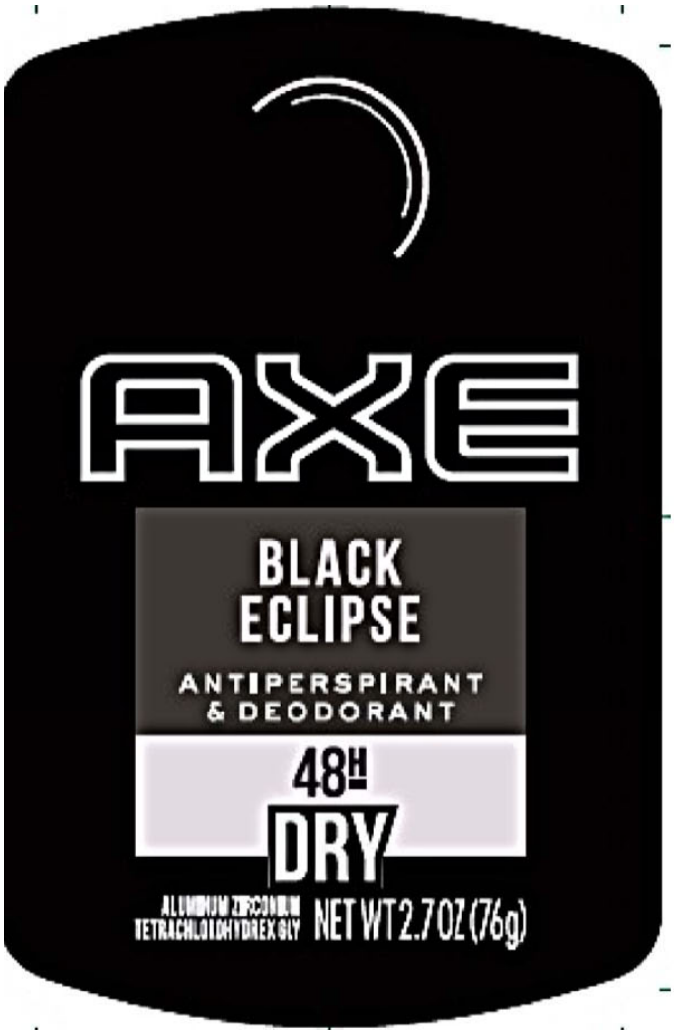
Inactive ingredients

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

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64942-1591
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**
TALC (UNII: 7SEV7J4R1U)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
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:64942-1591-1	76 g in 1 CONTAINER; Type 0: Not a Combination Product	01/01/2020	

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