

ANTIPERSPIRANT ANTISUDORIFIQUE- antiperspirant antisudorifique spray
Archifox GmbH & Co. KG

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

DIRECTIONS Underarm use only.

Decreases excessive sweating.

Dimethicone, Stearyl Alcohol, Kaolin, PPG-14 Butyl Ether, Hydrogenated Castor Oil, Octyldodecanol, Tocopherol, BHT, Parfum.

For external use only. Do not use on broken skin. Stop use if rash or irritation occurs. 'Aska doctor before use if you have kidney disease. Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center.

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PURPOSE: Antiperspirant

ACTIVE INGREDIENT: Aluminum Chloride 20%

Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center.

ANTIPERSPIRANT ANTISUDORIFIQUE

odaban@ daily DISTRIBUTED BY: Archifox GmbH & Co. KG Im Kitzwinkel 2, 34327 Kérle, Germany www.odaban.com 025381 5 ANTIPERSPIRANT deo stick DRUG FACTS ACTIVE INGREDIEN' PURPOSE: Aluminum Chloride 20%... Antiperspirant re Decreases excessive sweating. WARNINGS For external use only. Do not use on broken skin. Stop use if rash or irritation occurs. 'Aska doctor before use if you have kidney disease. Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center. DIRECTIONS Underarm use only. reat olan eS) Dimethicone, Stearyl Alcohol, Kaolin, PPG-14 Butyl Ether, Hydrogenated Castor Oil, Octyldodecanol, Tocopherol, BHT, Parfum. QUESTIONS? Net Wt. 2.1 oz

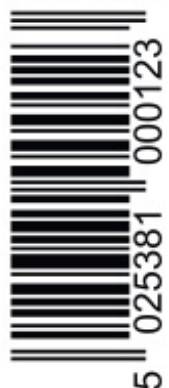
odaban 
daily



**ANTI
PERSPIRANT**
ANTISUDORIFIQUE

odaban 
daily **ANTIPERSPIRANT**
deo stick

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DRUG FACTS

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PURPOSE: Antiperspirant

USES

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INACTIVE INGREDIENTS

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QUESTIONS?

Call +1 (646) 566-6506

Net Wt. 2.1 oz

ANTIPERSPIRANT ANTISUDORIFIQUE

antiperspirant antisudorifique spray

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:84724-0002(NDC:84788-0002)
Route of Administration		TOPICAL		
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ALUMINUM CHLORIDE (UNII: 3CYT62D3GA) (ALUMINUM CATION - UNII:3XHB1D032B)			ALUMINUM CHLORIDE	12.4 mg in 62 mL
Inactive Ingredients				
Ingredient Name				Strength
TOCOPHEROL (UNII: R0ZB2556P8)				
PPG-14 BUTYL ETHER (UNII: R199TJT95T)				
KAOLIN (UNII: 24H4NWX5CO)				
DIMETHICONE (UNII: 92RU3N3Y1O)				
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)				
OCTYLDODECANOL (UNII: 461N1O614Y)				
BHT (UNII: 1P9D0Z171K)				
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84724-0002-1	62 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2025	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved drug other			12/01/2025	

Labeler - Archifox GmbH & Co. KG (315107452)

Registrant - Archifox GmbH & Co. KG (315107452)

Establishment			
Name	Address	ID/FEI	Business Operations
Archifox GmbH & Co. KG		315107452	relabel(84724-0002)