

1.8125"

3.2031"

ANTIPERSPIRANT & DEODORANT

Drug Facts

Active Ingredient Purpose
Aluminum zirconium trichlorohydrate 18% Antiperspirant

Use not use underarm perspiration

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.

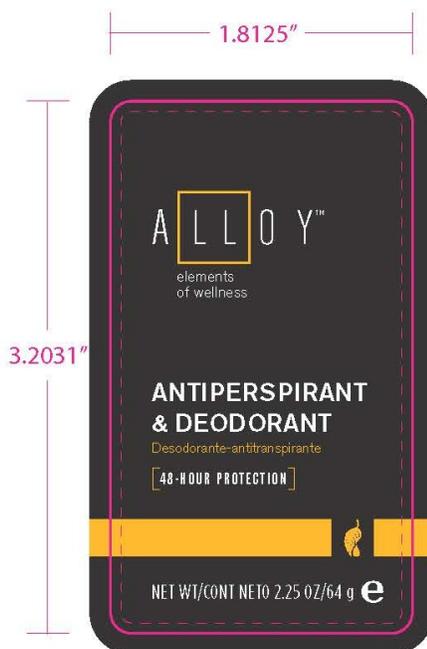
Directions Turn skin to rub product • apply to underarms only

Other information Do not freeze or store above 115°F (40°C)

Other ingredients
aluminum stearate, cocoyl lauric acid, sodium myristate (scent-free),
borax oil, benzoin extract, butyloctyl glycol, C12-15 alkyl
benzoate, C15-19 alkane, capryloyl glycolic acid, capryloyl
alcohol, dimethyl succinate, dimethylsiloxane, hydrogenated castor oil, perfume
(fragrance), silica, stearic alcohol, cocoyl myristate

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Product of Canada MP 20055 02/22 U
NOT FOR RESALE





ALLOY ANTIPERSPIRANT AND DEODORANT

aluminum zirconium trichlorohydrate gly stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54473-339
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY (UNII: T27D6T99LH) (ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY - UNII:T27D6T99LH)	ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY	10.24 g in 64 g

Inactive Ingredients

Ingredient Name	Strength
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.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
BAMBUSA ARUNDINACEA STEM (UNII: NRA4497HC5)
DIMETHICONE (UNII: 92RU3N3Y1O)
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)
C15-19 ALKANE (UNII: CI87N1IM01)
DIHEPTYL SUCCINATE (UNII: 057N7SS26Y)
WATER (UNII: 059QF0K00R)
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)
CETYL ALCOHOL (UNII: 936JST6JCN)
CAPRYLOYL GLYCERIN/SEBACIC ACID COPOLYMER (2000 MPA.S) (UNII: N7YC58165T)
BALANITES AEGYPTIACA SEED OIL (UNII: 75T177G59E)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54473-339-02	64 g in 1 APPLICATOR; Type 0: Not a Combination Product	04/20/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	04/20/2022	

Labeler - Melaleuca, Inc (139760102)

Registrant - Melaleuca, Inc (139760102)

Establishment

Name	Address	ID/FEI	Business Operations
KDC/ONE Development Corporation, Inc		204006464	manufacture(54473-339)

Revised: 2/2025

Melaleuca, Inc