RALPH LAUREN ROMANCE ANTIPERSPIRANT DEODORANT- aluminum zirconium tetrachlorohydrex gly stick
L'Oreal USA Products Inc

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.

Drug Facts

Active ingredient

Aluminum zirconium tetrachlorohydrex gly....16%

Use

reduces underarm wetness

Purpose

Antiperspirant

Warnings

For external use only

Do not use on broken skin

Stop use if rash or irritation occurs

Ask a doctor before use if you have kidney disease

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

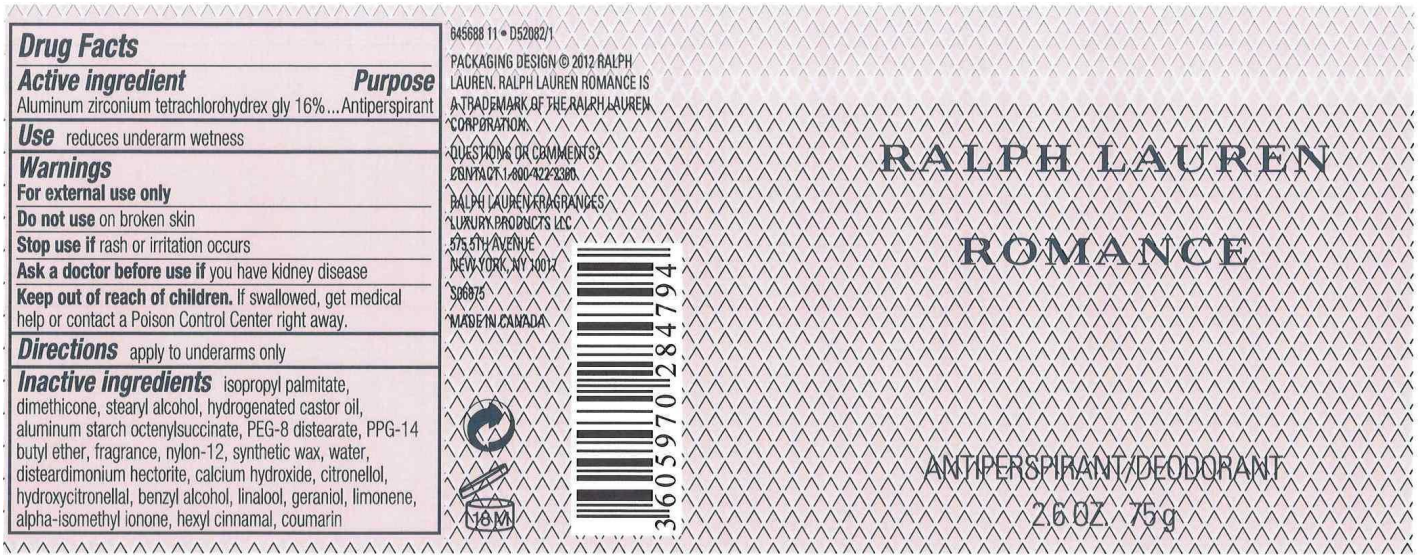
isopropyl palmitate, dimethicone, stearyl alcohol, hydrogenated castor oil, aluminum starch octenylsuccinate, PEG-8 distearate, PPG-14 butyl ether, fragrance, nylon-12, synthetic wax, water, disteardimonium hectorite, calcium hydroxide, citronellol, hydroxycitronellal, benzyl alcohol, linalool, geraniol, limonene, alpha-isomethyl ionone, hexyl cinnamal, coumarin

Questions or Comments?

Contact: 1-800-422-2360

Keep out of reach of children.

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



RALPH LAUREN ROMANCE ANTIPERSPIRANT DEODORANT

aluminum zirconium tetrachlorohydrex gly stick

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-479
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	1.2 g in 75 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
NYLON-12 (UNII: 446U8J075B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-479-01	75 g in 1 CONTAINER; Type 0: Not a Combination Product	07/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	07/01/2012	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-479)

Revised: 1/2020 L'Oreal USA Products Inc