BABARIA ALOE FRESH DEODORANT AND ANTIPERSPIRANT LIQUID ROLL-ON-
aluminum chlorohydrate emulsion
BERIOSKA SL

Babaria Aloe Fresh Deodorant and Antiperspirant Liquid Roll-On

Drug Facts

Active ingredient

Aluminum chlorohydrate 20%

Purpose

Antiperspirant

Uses

- Reduces underarm sweat
- 24 hour effective protection

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

Water (aqua), steareth-2, steareth-21, glycerin, aloe barbadensis leaf extract, octenidine

HCL, polysorbate 80, ethylhexylglycerin, fragrance (parfum), phenoxyethanol, sodium hydroxide, propylene glycol, dehydroacetic acid, benzoic acid, sorbic acid, citric acid, ascorbic acid, sodium sulfite, sodium benzoate, potassium sorbate.

Package Labeling:

BABARIA ALOE FRESH DEODORANT AND ANTIPERSPIRANT LIQUID ROLL-ON

aluminum chlorohydrate emulsion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78283-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	200 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
OCTENIDINE HYDROCHLORIDE (UNII: U84956NU4B)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SORBIC ACID (UNII: X045WJ989B)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78283-001-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/18/2024	01/01/2030

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	03/18/2024	01/01/2030

Labeler - BERIOSKA SL (462392556)

Revised: 12/2025 BERIOSKA SL