

**BABARIA DOUBLE EFFECT DEODORANT AND ANTIPERSPIRANT LIQUID ROLL-ON- aluminum chlorohydrate emulsion
BERIOSKA SL**

Babaria Double Effect 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, glycerin, steareth-21, tocopherol acetate octenidine HCL,

creosote bush (*Larrea tridentata* var, *tridentata*) extract, tocopherol, dimethicone fragrance (parfum), biotin, lecithin, phenoxyethanol, sodium hydroxide, propylene glycol, ethylhexylglycerin, dehydroacetic acid, benzoic acid, sorbic acid, butylene glycol, ascorbic acid, sodium citrate, potassium sorbate, citric acid, sodium metabisulfite, limonene, linalool, hexyl cinnamal.

Package Labeling:

babaria

double effect

24h PROTECTION

Deodorant and Antiperspirant Liquid Roll-on
20% Aluminum Chlorohydrate

1.7 fl oz (50 mL)

0% alcohol
Silky skin
With Vitamin E

Drug Facts

Active ingredient Purpose
Aluminum chlorohydrate 20% 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

Drug Facts (continued)
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, glycerin, steareth-21, tocophenyl acetate, octenidine HCl, creosote bush (*Larrea tridentata* var. *tridentata*) extract, tocopherol, dimethicone, fragrance (parfum), biotin, lecithin, phenoxyethanol, sodium hydroxide, propylene glycol, ethylhexylglycerin, dehydroacetic acid, benzoic acid, sorbic acid, butylene glycol, ascorbic acid, sodium citrate, potassium sorbate, citric acid, sodium metabisulfite, limonene, linalool, hexyl cinnamal.

MANUFACTURED BY: BABARIA-BERIOSKA S.L., POLIGONO INDUSTRIAL CASTILLA, NO. 8-3, CHESTE, VALENCIA E-46380 SPAIN MADE IN SPAIN
TO REPORT A SERIOUS ADVERSE EVENT, CONTACT BABARIA USA LLC. AT 3325 NW 70TH AVE, MIAMI, FL 33122
DERMATOLOGICALLY TESTED.

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BABARIA DOUBLE EFFECT DEODORANT AND ANTIPERSPIRANT LIQUID ROLL-ON

aluminum chlorohydrate emulsion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78283-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZ W13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZ W13M)	ALUMINUM CHLOROHYDRATE	200 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
STEARETH-2 (UNII: V56DFE46J5)	
LARREA TRIDENTATA WHOLE (UNII: B755J144H1)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEARETH-21 (UNII: 53J3F32P58)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
OCTENIDINE HYDROCHLORIDE (UNII: U84956NU4B)	
TOCOPHEROL (UNII: R0ZB2556P8)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
BIOTIN (UNII: 6SO6U10H04)	
PHENOXYETHANOL (UNII: HIE492Z Z3T)	

SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SORBIC ACID (UNII: X045WJ989B)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6037OK2I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78283-003-01	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