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 Posion Control Center right away.

Directions

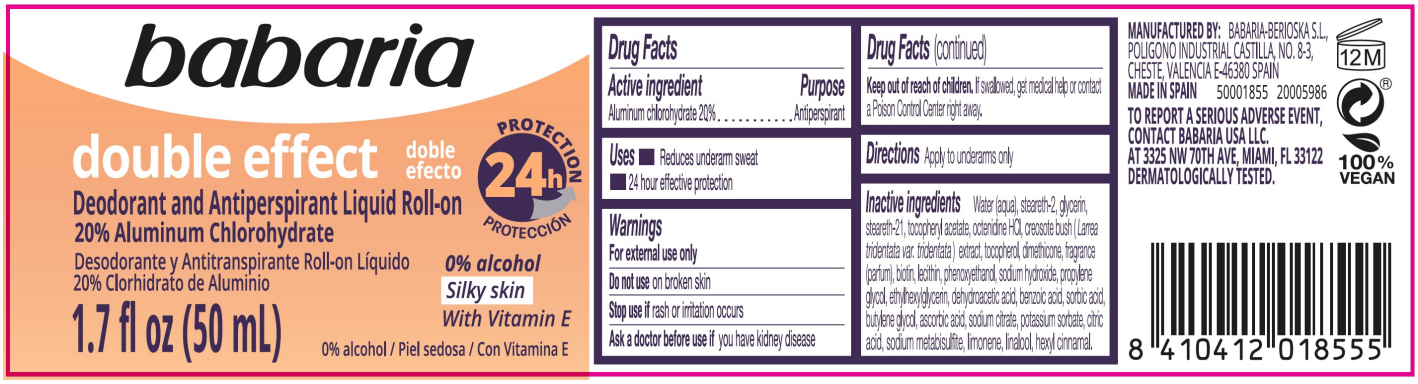
Apply to underarms only

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 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: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	200 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
LARREA TRIDENTATA WHOLE (UNII: B755J144H1)	
WATER (UNII: 059QF0KO0R)	
STEARETH-2 (UNII: V56DFE46J5)	
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: HIE492ZZ3T)	

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: 7X6O37OK2I)	

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	

Marketing Information

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

Labeler - BERIOSKA SL (462392556)

Revised: 12/2023 BERIOSKA SL