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

Babaria Oat 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 Posion Control Center right away.

Directions

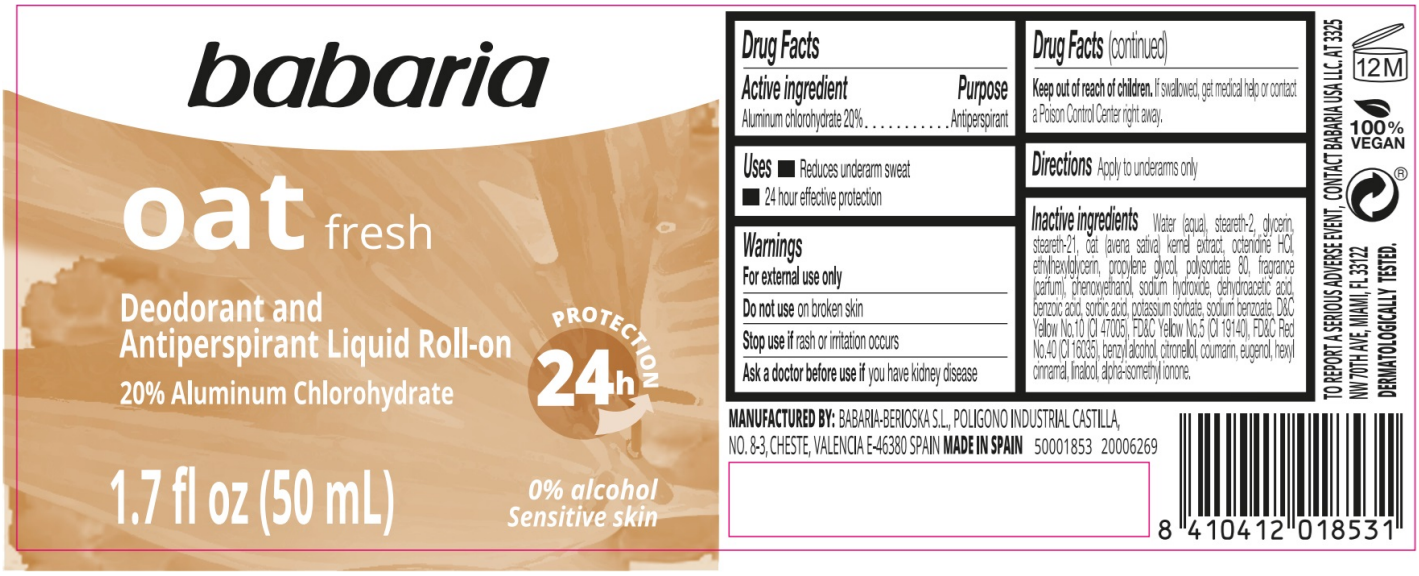
Apply to underarms only

Inactive ingredients

Water (aqua), steareth-2, glycerin, steareth-21, oat (avena sativa) kernel extract,

octenidine HCL, ethylhexylglycerin, propylene glycol, Polysorbate 80, fragrance (parfum), phenoxyethanol, sodium hydroxide, dehydroacetic acid, benzoic acid, sorbic acid, potassium sorbate, sodium benzoate, D&C Yellow No.10 (CI 47005), FD&C Yellow No.5 (CI 19140), FD&C Red No.40 (CI 16035), benzyl alcohol, citronellol, coumarin, eugenol, hexyl cinnamal, linalool, alpha-isomethyl ionone.

Package Labeling:



BABARIA OAT FRESH DEODORANT AND ANTIPERSPIRANT LIQUID ROLL-ON

aluminum chlorohydrate emulsion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78283-005
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEARETH-21 (UNII: 53J3F32P58)	
AVENA SATIVA FLOWERING TOP (UNII: MA9CQJ3F7F)	
OCTENIDINE HYDROCHLORIDE (UNII: U84956NU4B)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SORBIC ACID (UNII: X045WJ989B)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
COUMARIN (UNII: A4VZ22K1WT)	
EUGENOL (UNII: 3T8H1794QW)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
ISOMETHYL-.ALPHA.-IONONE (UNII: 9XP4LC555B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78283-005-01	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