

**ESIKA TOTAL SEC NEUTRAL FRAGRANCE ANTIPERSPIRANT AND DEODORANT
UNISEX ROLL-ON ALL DAY EXTRA EFFECTIVE PROTECTION- aluminum
sesquichlorohydrate emulsion
BEL STAR S A**

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.

**esika total sec NEUTRAL FRAGRANCE Antiperspirant and deodorant Unisex
Roll-on All day extra effective protection**

Active Ingredient

Aluminum sesquichlorohydrate 15%

Purpose

Antiperspirant

Uses

Reduces underarm perspiration

Warnings

For external use only

Do not use

on broken skin

Ask a doctor before use if you have

Kidney disease

Stop use if

rash or irritation occurs

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, steareth-2, ppg-15 stearyl ether, cyclohexasiloxane, steareth-21, benzyl alcohol, dicaprylyl carbonate, 2-methyl 5-cyclohexylpentanol, bisabolol, hydroxyacetophenone, fragrance, tetrasodium edta, bht, propylene glycol, bambusa arundinacea leaf extract, potassium sorbate, sodium benzoate, hexyl cinnamal, limonene, linalool.

Product Packaging

total sec

NEUTRAL FRAGRANCE

Antiperspirant and deodorant

Unisex roll-on

All day extra effective protection

esika

1.6 fl. oz./50 ml



ESIKA TOTAL SEC NEUTRAL FRAGRANCE ANTIPERSPIRANT AND DEODORANT UNISEX ROLL-ON ALL DAY EXTRA EFFECTIVE PROTECTION

aluminum sesquichlorohydrate emulsion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:14141-962
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM SESQUICHLOROHYDRATE (UNII: UCN889409V) (ALUMINUM SESQUICHLOROHYDRATE - UNII:UCN889409V)	ALUMINUM SESQUICHLOROHYDRATE	15 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
2-METHYL 5-CYCLOHEXYLPENTANOL (UNII: 460837ILID)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
EDETATE SODIUM (UNII: MP1J8420LU)	
STEARETH-21 (UNII: 53J3F32P58)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
LEVOMENOL (UNII: 24WE03BX2T)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
STEARETH-2 (UNII: V56DFE46J5)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
BAMBUSA BAMBOS LEAF (UNII: HW86D1FGSS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:14141-962-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/07/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	06/07/2021	

Labeler - BEL STAR S A (880160197)

Establishment

Name	Address	ID/FEI	Business Operations
BEL STAR S A		880160197	manufacture(14141-962)

Revised: 3/2022

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