

**ESIKA EXUS ROLL-ON- aluminum sesquichlorohydrate liquid**  
**Ventura Corporation Ltd.**

*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.*

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**EXUS**  
**ROLL-ON DEODORANT**

***Drug Facts***

**Active Ingredients**

Aluminum sesquichlorohydrate (17%)

**Purpose**

Antiperspirant

**Uses**

- reduces underarm wetness.

**Warnings**

- **For external use only.**
- Do not use on broken skin.
- Stop use and ask a doctor if rash and 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**

Aqua (water), steareth-2, ppg-15 stearyl ether, steareth-21, fragrance, dicaprylyl carbonate, cyclopentasiloxane, triclosan, cyclohexasiloxane, benzalkonium chloride, tetrasodium edta, bisabolol, magnesium nitrate, methylchloroisothiazolinone, magnesium chloride, methylisothiazolinone.

**PR:** Distributed by Ventura Corporation, Ltd. San Juan, Puerto Rico 00926.

**PRINCIPAL DISPLAY PANEL - 50 ml Container Label**

**EXUS**

**ROLL-ON DEODORANT**

does not contain alcohol

**ēsika**

50 ml e (1.7 fl.oz.)

# EXUS

ROLL-ON DEODORANT  
does not contain alcohol

50 ml e (1.7 fl.oz.)

Controls excess perspiration and protects against odors, leaving a pleasant scent.

<b>Drug Facts</b>				
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## ESIKA EXUS ROLL-ON

aluminum sesquichlorohydrate liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:13537-062
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Aluminum Sesquichlorohydrate</b> (UNII: UCN889409V) (Aluminum Sesquichlorohydrate - UNII:UCN889409V)	Aluminum Sesquichlorohydrate	0.17 g in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0K00R)	
steareth-2 (UNII: V56DFE46J5)	
steareth-21 (UNII: 53J3F32P58)	
dicaprylyl carbonate (UNII: 609A3V1SUA)	
cyclomethicone 5 (UNII: 0THT5PCI0R)	
triclosan (UNII: 4NM5039Y5X)	
cyclomethicone 6 (UNII: XHK3U310BA)	
benzalkonium chloride (UNII: F5UM2KM3W7)	
edetate sodium (UNII: MP1J8420LU)	
levomenol (UNII: 24WE03BX2T)	
magnesium nitrate (UNII: 77CBG3UN78)	
methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
magnesium chloride (UNII: 02F3473H9O)	
methylisothiazolinone (UNII: 229D0E1QFA)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13537-062-01	50 mL in 1 BOTTLE		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part350	12/13/2011	

**Labeler** - Ventura Corporation Ltd. (602751344)**Establishment**

Name	Address	ID/FEI	Business Operations
Bel Star S.A. (Colombia)		880160197	MANUFACTURE(13537-062)

Revised: 12/2011

Ventura Corporation Ltd.