

ESIKA TOTAL SEC DELICATE SKIN- 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.

ESIKA Total Sec Delicate Skin Alcohol-Free Women's Roll-On Deodorant and Antiperspirant All Day Extra Effective Protection

Drug Facts

Active Ingredient

Aluminum sesquichlorohydrate 16.2 %

Purpose

Antiperspirant

Uses

- Reduces underarm perspiration
- All day extra effective protection

Warnings

For external use only

Do not use on broken skin

Ask a doctor before use if you have Kidney disease

Stop use and ask a doctor 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, ALUMINUM SESQUICHLOROXYDRATE, STEARETH-2, STEARETH-21, PPG-15 STEARYL ETHER, CYCLOPENTASILOXANE, FRAGRANCE, CYCLOHEXASILOXANE, DICAPRYLYL CARBONATE, BISABOLOC, TRICLOSAN, BENZALKONIUM CHLORIDE, METHYLPARABEN, BHT, PROPYLENE GLYCOL, TETRASODIUM EDTA, BAMBUSIA ARUNDINACEA LEAF EXTRACT

Dist. By Ventura Corporation, Ltd. San Juan, Puerto Rico 00926.

PRINCIPAL DISPLAY PANEL - 50 ml Bottle Label

ēsika
total
sec

DELICATE SKIN

Alcohol-free
Women's Roll-on
Deodorant and
Antiperspirant
All day extra
effective protection

36

50 ml e (1.7 fl.oz.)

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Directions ■ Apply to underarms only.

Inactive Ingredients
aqua (water), steareth-2, steareth-21, ppg-15 stearyl ether, cyclopentasiloxane, parfüm (fragrance), cyclohexasiloxane, dicaprylyl carbonate, bisabolol, triclosan, benzalkonium chloride, methyparaben, bht, propylene glycol, tetrasodium edta, bambusa arundinacea leaf extract.

• DERMATOLOGICALLY TESTED • HYPOALLERGENIC
P.R.: Dist. by Veneria Corporation, Ltd. San Juan, Puerto Rico 00926.
Made in Colombia.

ESIKA TOTAL SEC DELICATE SKIN

aluminum sesquichlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:13537-157
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM SESQUICHLORO HYDRATE (UNII: UCN889409 V) (ALUMINUM SESQUICHLORO HYDRATE - UNII:UCN889409 V)	ALUMINUM SESQUICHLORO HYDRATE	0.162 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARETH-15 (UNII: O6V041E38J)	
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
LEVOMENOL (UNII: 24WE03BX2T)	
TRICLOSAN (UNII: 4NM5039 Y5X)	

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	
BAMBUSA ARUNDINACEA LEAF (UNII: HW86D1FGSS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13537-157-01	50 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part350	01/24/2012	

Labeler - Ventura Corporation, Ltd. (602751344)

Establishment

Name	Address	ID/FEI	Business Operations
Bel Star S.A. (Colombia)		880160197	MANUFACTURE(13537-157)

Revised: 1/2012

Ventura Corporation, Ltd.