

**SKIN BRIGHTENING ANTIPERSPIRANT DEODORANT ROLL-ON JAFRA DAILY-**

**aluminum chlorohydrate liquid**

**Jafra cosmetics International**

*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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**Jafra Daily Skin Brightening antiperspirant roll-on**

**Active ingredients**

**Purpose**

Aluminum Chlorohydrate 12.3%    Antiperspirant

**Uses**

reduces underarm wetness

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

**Stop use** if rash or irritation occurs

**Warnings**

For external use only

Do not use on broken skin

Ask a doctor before use if you have kidney disease

**Directions**

Apply to underarms only

Water/Aqua, PEG-40 Stearate, PEG-25 Propylene Glycol Stearate, Cetyl Alcohol, Magnesium Aluminum Silicate, Sorbitan Sesquioleate, Fragrance/parfum, Phenoxyethanol, Ehtylhexylhexyl Glycerin, Mica, Titanium Dioxide/CI 77891, Azadirachta Indica Extract, Curcuma Longa, Hemisdesmus Indicus, Sphaeranthus Indicus Extract, Phyllanthus Emblica Extract

Jafra

Daily

Skin Brightening

antiperspirant deodorant roll-on

60 ml 2 FL OZ

FORMULA: JLD144-10497001  
.SM



493 c



Black



### Drug Facts

**Active Ingredients**      **Purpose**  
Aluminum Chlorohydrate 12.3%....Antiperspirant

**Uses**  
■ reduces underarm wetness

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Made in Mexico for:  
JAFRA Cosmetics International, Inc.  
Westlake Village, CA 91361  
Distributed in EU: JAFRA Cosmetics  
D-80636 Munich  
[www.JAFRA.com](http://www.JAFRA.com)  
Hecho en México para:  
Distribuidora Comercial JAFRA, S.A. de C.V.  
Blvd. Adolfo López Mateos 2273 (antes 515),  
C.P. 01710, Distrito Federal, México.  
JAFRA es Marca Registrada  
LOS ANGELES - ZURICH - MUNICH  
MEXICO CITY - MILAN - MOSCOW

JAFRA  
DAILY

skin  
brightening  
antiperspirant deodorant  
roll-on

60 ml e 2 FL OZ (US)

<b>SKIN BRIGHTENING ANTIPERSPIRANT DEODORANT ROLL-ON JAFRA DAILY</b>			
aluminum chlorohydrate liquid			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68828-199
<b>Route of Administration</b>	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	12.3 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PEG-25 PROPYLENE GLYCOL STEARATE (UNII: X21KPH4633)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
MICA (UNII: V8A1AW0880)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
AZADIRACHTA INDICA BARK (UNII: G580B439YI)	
TURMERIC (UNII: 856YO1Z64F)	
HEMIDESMUS INDICUS ROOT (UNII: Y5CFT48S90)	
SPHAERANTHUS INDICUS FLOWERING TOP (UNII: 1O5Y93LB44)	
PHYLLANTHUS EMBLICA FRUIT (UNII: YLX4CW2576)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68828-199-02	1 in 1 CARTON	09/03/2014	
1	NDC:68828-199-01	60 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	09/03/2014	

**Labeler** - Jafra cosmetics International (041676479)

**Registrant** - Jafra cosmetics International (041676479)

## Establishment

Name	Address	ID/FEI	Business Operations
Jafra Manufacturing, S.A. de C.V.		814732061	manufacture(68828-199)