

CC CREAM COMPLEXION CORRECTOR LIGHT MEDIUM BROAD SPECTRUM SPF 15 JAFRA- avobenzone, octinoxate, octisalate, octocrylene cream

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.

Jafra CC Creme Complexion Corrector Broad Spectrum SPF 15 Light Medium

Active Ingredients	Purpose
Avobenzone 2%	Sunscreen
Octinoxate 5%	Sunscreen
Octisalate 4%	Sunscreen
Octocrylene 1.8%	Sunscreen

Uses

helps prevent sunburn

if used as directed with other sun protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun

Keep out of reach of children

Stop use and ask a doctor if rash occurs

For External use only

Do not use on damaged or broken skin

When using this product, keep out of eyes. Rinse with water to remove.

If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

apply liberally 15 minutes before sun exposure

use a water resistant sunscreen if swimming or sweating

reapply at least every 2 hours

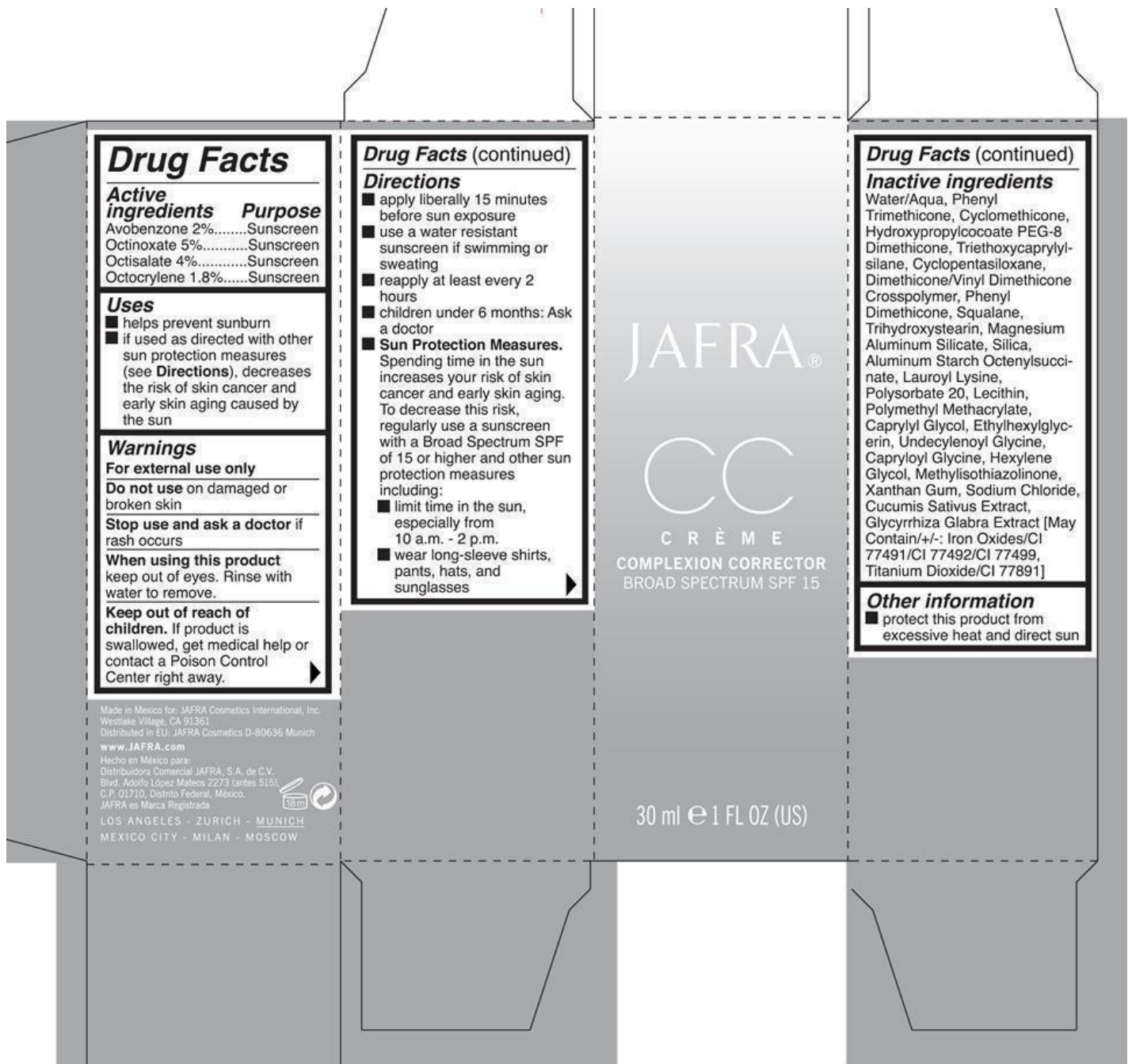
children under 6 months

Sun Protection Measures

Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk of 15 or higher and other sun protection measures including:

- limit time in the sun, especially from 10 a.m. - 2p.m.
- wear long sleeve-shirts, pants, hats, and sunglasses

Water/Aqua, Phenyl Trimethicone, Cyclomethicone, Hydroxypropylcocoate PEG-8 Dimethicone, Triethoxycaprylylsilane, Cyclopentasiloxane, Dimethicone/Vinyl Dimethicone Crosspolymer, Phenyl Dimethicone, Squalane, Trihydroxystearin, Magnesium Aluminum Silicate, Silica, aluminum Starch Octenylsuccinate, Lauryl Lysine, Polysorbate 20, Lecithin, Polymethyl Methacrylate, Caprylyl Glycol, Ethylhexylglycerin, Undecylenoyl Glycine, Hexylene Glycol, Methylisothiazolinone, Xanthan Gum, Sodium Chloride, Cucumis Sativus Extract, Glycyrrhiza Glabra Extract [May contain+/-: Iron Oxides/CI 77491/CI 77492/77499, Titanium Dioxide/CI 77891]



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68828-190
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2 g in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	5 g in 100 mL

OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	4 g in 100 mL
OCTOCRYLENE (UNII: 5A68 WGF6 WM) (OCTOCRYLENE - UNII:5A68 WGF6 WM)	OCTOCRYLENE	1.8 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
CYCLOMETHICONE (UNII: NMQ347994Z)	
HYDROXYPROPYLCOCOATE PEG-8 DIMETHICONE (UNII: 8TE0BZU36S)	
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)	
CYCLOMETHICONE 5 (UNII: 0THI5PCI0R)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
SQUALANE (UNII: GW89575KF9)	
TRIHYDROXYSTEARIN (UNII: 06YD7896S3)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
LAUROYL LYSINE (UNII: 113171Q70B)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
POLY(METHYL METHACRYLATE; 450000 MW) (UNII: Z47NNT4J11)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
UNDECYLENOYL GLYCINE (UNII: 4D20464K2J)	
CAPRYLOYL GLYCINE (UNII: 8TY5YO42NJ)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CUCUMBER (UNII: YY7C30VXJT)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68828-190-02	1 in 1 CARTON	03/03/2014	
1	NDC:68828-190-01	30 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/03/2014	

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(68828-190)

Revised: 1/2019

Jafra cosmetics International