ACTIVE ARGAN 02.3(TM) PLASMA PRO 50 FEATHERWEIGHT DAY BROAD SPECTRUM SPF 20 WITH PROPRIETARY A.P.I. AND ACTIVATED ARGAN COMPLEX(TM)- zinc oxide cream SIBORG

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.

ACTIVE ARGAN 02.3(TM) PLASMA PRO 50+ FEATHERWEIGHT DAY CREAM BROAD SPECTRUM SPF 20 WITH PROPRIETARY A.P.I. & ACTIVATED ARGAN COMPLEX(TM)

Active ingredients

Zinc oxide 14%

Purpose

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

Warnings

Warnings

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.

Stop use and ask a doctor if rash occurs

Keep out of reach of children.

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

Directions

- apply liberally and evenly 15 minutes before sun exposure
- Reapply at least every 2 hours
- Use a water resistant sunscreen if swimming or sweating
- **Sun Protection Measures**: Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a broad spectrum SPF value of 15 or higher and other sun protection measures including:
- Limit time in the sun, especially from 10 a.m. 2 p.m.
- Wear long-sleeved shirts, pants, hats, and sunglasses
- Children under 6 months of age: Ask a doctor

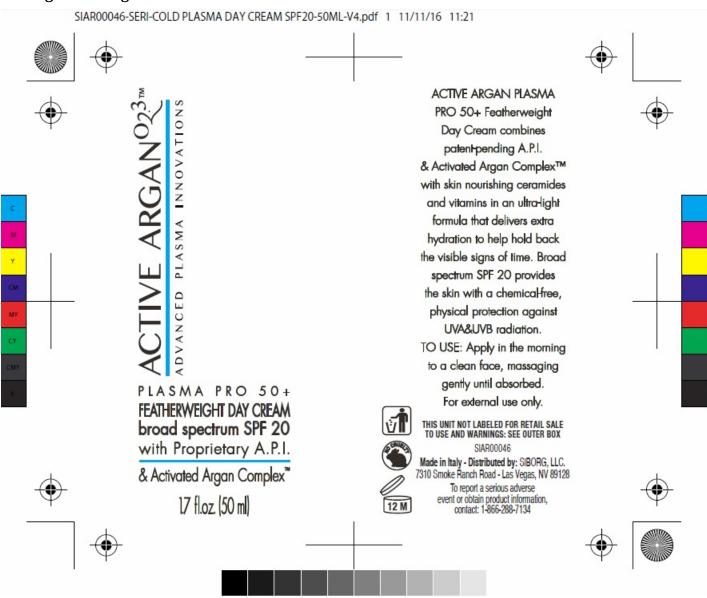
Other information

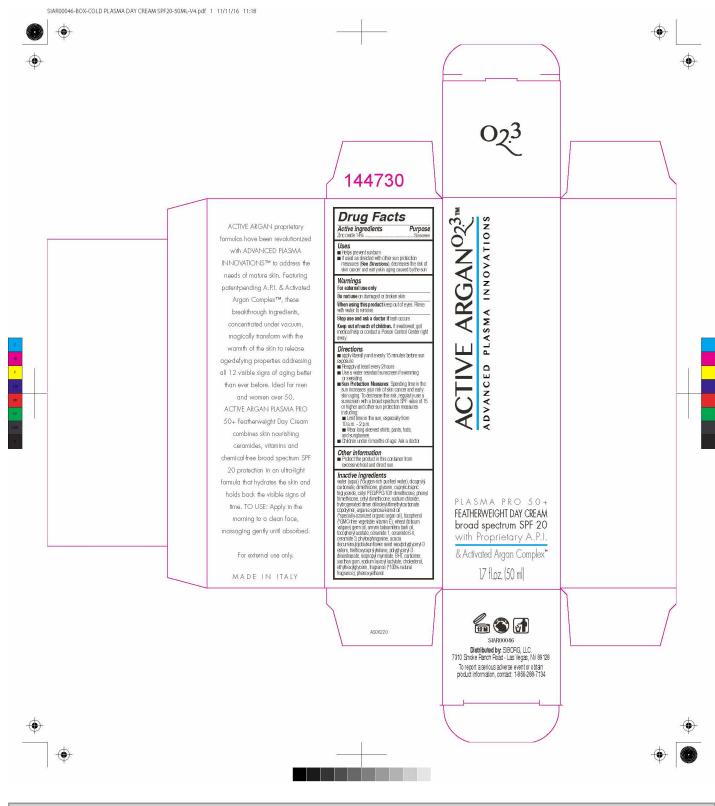
Protect the product in this container from excessive heat and direct sun.

Inactive ingredients

water (aqua) (*oxygen-rich purified water), dicaprylyl carbonate, dimethicone, glycerin, caprylic/capric triglyceride, cetyl PEG/PPG-10/1 dimethicone, phenyl trimethicone, cetyl dimethicone, sodium chloride, hydrogenated dimer dilinoleyl/dimethylcarbonate copolymer, argania spinosa kernel oil (*specially-ozonized organic argan oil), tocopherol (*GMO-free vegetable vitamin E), wheat (triticum vulgare) germ oil, amyris balsamifera bark oil, tocopheryl acetate, ceramide 1, ceramide 6-II, ceramide 3, phytosphingosine, acacia decurrens/jojoba/sunflower seed wax/polyglyceryl-3 esters, triethoxycaprylylsilane, polyglyceryl-3 diisostearate, isopropyl myristate, BHT, carbomer, xanthan gum, sodium lauroyl lactylate, cholesterol, ethylhexylglycerin, fragrance (*100% natural fragrance), phenoxyethanol

Package Labeling:





ACTIVE ARGAN 02.3(TM) PLASMA PRO 50 FEATHERWEIGHT DAY BROAD SPECTRUM SPF 20 WITH PROPRIETARY A.P.I. AND ACTIVATED ARGAN COMPLEX(TM)

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70803-023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	140 mg in 1 mL	

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DICAPRYLYL CARBONATE (UNII: 609 A3V1SUA)	
DIMETHICO NE (UNII: 92RU3N3Y1O)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PHENYL TRIMETHICO NE (UNII: DR0 K5NOJ4R)	
SO DIUM CHLO RIDE (UNII: 451W47IQ8 X)	
ARGAN OIL (UNII: 4V59G5UW9X)	
TOCOPHEROL (UNII: R0ZB2556P8)	
WHEAT GERM OIL (UNII: 14C97E680P)	
AMYRIS BALSAMIFERA O IL (UNII: 11BJ961J2E)	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	
CERAMIDE 1 (UNII: 5THT33P7X7)	
CERAMIDE NP (UNII: 4370 DF0 50 B)	
PHYTOSPHINGOSINE (UNII: GIN46U9Q2Q)	
TRIETHO XYCAPRYLYLSILANE (UNII: LDC331P08E)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46 P231IQ V8)	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)	
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)	
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM307FC)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM LAURO YL LACTYLATE (UNII: 7243K85WFO)	
CHOLESTEROL (UNII: 97C5T2UQ7J)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENO XYETHANOL (UNII: HIE492ZZ3T)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:70803-023-50	1 in 1 BOX	12/0 1/20 16	
1	50 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part352	12/0 1/20 16		

Labeler - SIBORG (102875148)

Revised: 2/2018 SIBORG