# 6000X ELEMENTAL SCREEN BROAD SPECTRUM SPF30 SUNSCREEN- zinc oxide cream Allure Labs Inc

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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### **Drug Facts**

Active Ingredients Zinc Oxide 18.9%

Purpose

Sunscreen

### Uses

- helps to 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

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

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
- Children under 6 months: Ask a doctor

Reapply:

- after swimming or sweating
- immediately after towel drying
- at least every 2 hours
- 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 of 15 or higher and other sun protection measures including:
- limit time in the sun, especially from 10am 2pm
- wear long-sleeve shirts, pants, hats, and sunglasses

### **Inactive Ingredients**

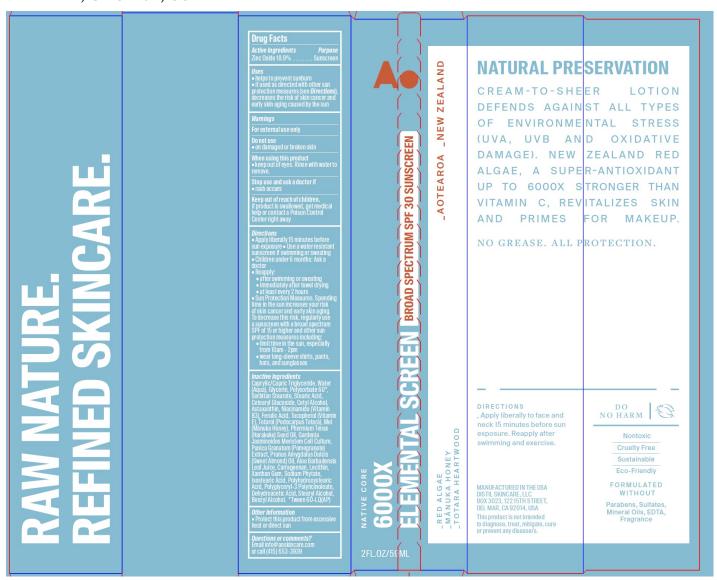
Caprylic/Capric Triglyceride, Water(Aqua), Glycerin, Polysorbate 60\*, Sorbitan Stearate, Stearic Acid, Cetearyl Glucoside, Cetyl Alcohol, Astaxanthin, Niacinamide (Vitamin B3), Ferulic Acid, Tocopherol (Vitamin E), Totarol (Podocarpus Totara), Mel (Manuka Honey), Phormium Tenax (Harakeke) Seed Oil, Gardenia Jasminoides Meristem Cell Culture, Punica Granatum (Pomegranate) Extract, Prunus Amygdalus Dulcis (Sweet Almond) Oil, Aloe Barbadensis Leaf Juice, Carrageenan, Lecithin, Xanthan Gum, Sodium Phytate, Isostearic Acid, Polyhydroxystearic Acid, Polyglyceryl-3 Polyricinoleate, Dehydroacetic Acid, Stearyl Alcohol, Benzyl Alcohol. \*Tween 60-LQ(AP)

### Other Information

• Protect this product from excessive heat or direct sun

Questions or comments? Email info@aoskincare.com or call (415) 653-3939

MANUFACTURED IN THE USA DISTIL SKINCARE, LLC BOX 3023, 122 15TH STREET, DEL MAR, CA 92104, USA



# 6000X ELEMENTAL SCREEN BROAD SPECTRUM SPF30 SUNSCREEN zinc oxide cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:62742-4200 Route of Administration TOPICAL Active Ingredient/Active Moiety

**Basis of Strength** 

Strength

**Ingredient Name** 

Inactive Ingredients  Ingredient Name	Strength				
WATER (UNII: 059QF0KO0R)	Strength				
GLYCERIN (UNII: PDC6A3C0OX)					
POLYSORBATE 60 (UNII: CAL22UVI4M)					
SO RBITAN MONO STEARATE (UNII: NVZ4I0 H58 X)  MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9 H2L21V7U)					
FERULIC ACID (UNII: AVM951ZWST)					
PHORMIUM TENAX SEED OIL (UNII: OZ80CEA779)					
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)					
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q0OK5DOT4)					
PUNICA GRANATUM ROOT BARK (UNII: CLV24I3T1D)					
ALOE VERA LEAF (UNII: ZY81Z83H0X)					
CETEARYL GLUCO SIDE (UNII: 09 FUA47KNA)					
HO NEY (UNII: Y9 H1V576 FH)					
GARDENIA JASMINO IDES WHO LE (UNII: 0 PK353KHF0)					
CARRAGEENAN (UNII: 5C69 YCD2YJ)					
STEARYL ALCOHOL (UNII: 2KR8914H1Y)					
CETYL ALCOHOL (UNII: 936JST6JCN)					
ASTAXANTHIN (UNII: 8 XPW32PR7I)					
NIACINAMIDE (UNII: 25X51I8RD4)					
ALPHATOCOPHEROL (UNII: H4N855PNZ1)					
TOTAROL (UNII: 67NH2854WW)					
ALMOND OIL (UNII: 66 YXD4DKO9)					
XANTHAN GUM (UNII: TTV12P4NEE)					
DEHYDRO ACETIC ACID (UNII: 2KAG279 R6 R)					
STEARIC ACID (UNII: 4ELV7Z65AP)					
ISOSTEARIC ACID (UNII: X33R8U0062)					
BENZYL ALCOHOL (UNII: LKG8494WBH)					
LECITHIN, SO YBEAN (UNII: 1DI56 QDM62)					
PHYTATE SO DIUM (UNII: 88496G1ERL)					

F	Packaging					
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>		
1	NDC:62742-4200-2	1 in 1 CARTON	02/08/2021			
1	NDC:62742-4200-1	59 g 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	02/08/2021				

## Registrant - Allure Labs Inc (926831603)

Establishment				
Name	Address	ID/FEI	Business Operations	
Allure Labs		926831603	manufacture(62742-4200)	

Revised: 2/2021 Allure Labs Inc