

LIPID SHIELD SPF 30- zinc oxide cream

Allure Labs

Drug Facts

Active Ingredient: Zinc Oxide 21%

Purpose: Sunscreen

Uses • Helps prevent sunburn and damage caused by UVA/UVB exposure • 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.

Do not use on damaged or broken skin

When using this product In the event of contact with eyes, rinse thoroughly with water

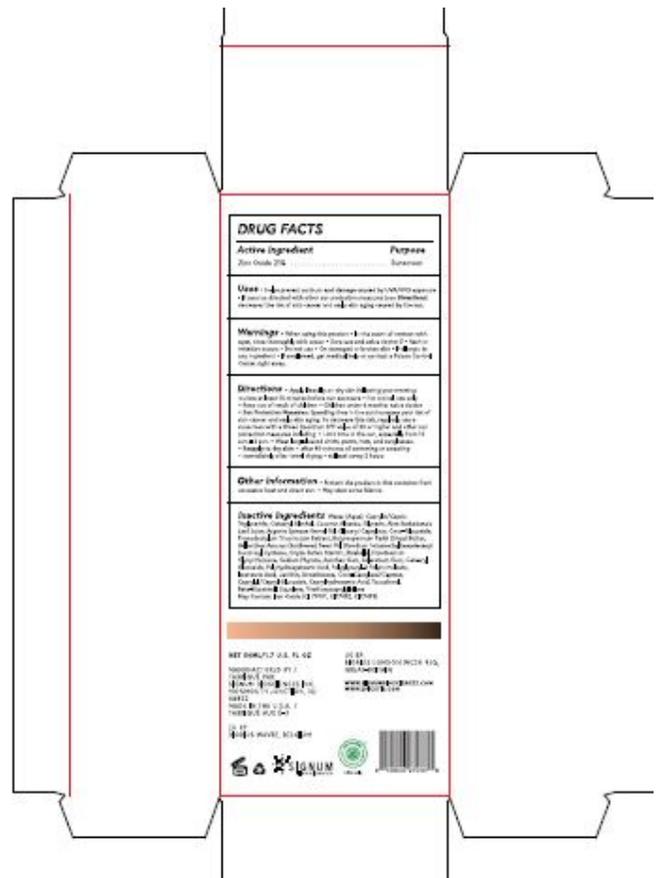
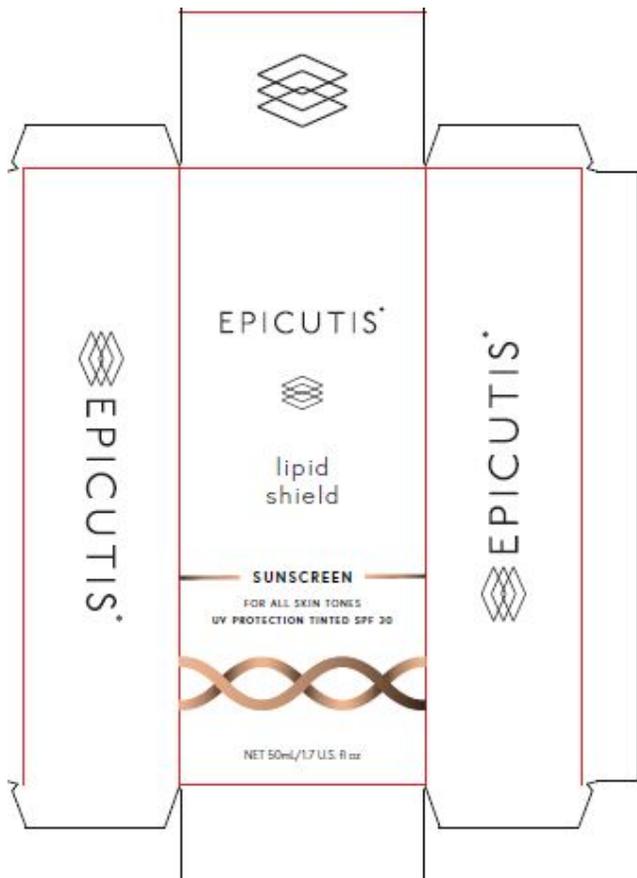
Stop use and ask a doctor if Rash or irritation occurs

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

Directions • Apply liberally on dry skin following your morning routine at least 15 minutes before sun exposure • For topical use only • Keep out of reach of children • Children under 6 months: ask a doctor • 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 30 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. • Reapply to dry skin: • after 80 minutes of swimming or sweating • immediately after towel drying • at least every 2 hours

Other Information • Protect the product in this container from excessive heat and direct sun. • May stain some fabrics.

Inactive Ingredients: Water (Aqua), Caprylic/Capric Triglyceride, Cetearyl Alcohol, Coconut Alkanes, Glycerin, Aloe Barbadensis Leaf Juice, Argania Spinosa Kernel Oil, Glyceryl Caprylate, Coco-Glucoside, Phaeodactylum Tricornutum Extract, Butyrospermum Parkii (Shea) Butter, Helianthus Annuus (Sunflower) Seed Oil, Disodium Tetramethylhexadecenyl Succinoyl Cysteine, Oryza Sativa Starch, Bisabolol, Dipotassium Glycyrrhizinate, Sodium Phytate, Xanthan Gum, Sclerotium Gum, Cetearyl Glucoside, Polyhydroxystearic Acid, Polyglyceryl-3 Polyricinoleate, Isostearic Acid, Lecithin, Dimethicone, Coco-Caprylate/Caprinate, Caprylyl/Capryl Glucoside, Caprylhydroxamic Acid, Tocopherol, Beta-Sitosterol, Squalene, Triethoxycaprylylsilane
May Contain: Iron Oxide (CI 77491, CI77492, CI77499)



LIPID SHIELD SPF 30

zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4273
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	21 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERYL CAPRYLATE (UNII: TM2TZD4G4A)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHAEODACTYLUM TRICORNUTUM (UNII: Y5W63E7HLV)	
SHEA BUTTER (UNII: K49155WL9Y)	
DISODIUM TETRAMETHYLHEXADECENYL SUCCINOYL CYSTEINE (UNII: CZA8L664YS)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
DIPOTASSIUM GLYCYRRHIZATE (UNII: CA2Y0FE3FX)	
CI 77499 (UNII: XM0M87F357)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	

COCO-GLUCOSIDE (UNII: ICS790225B)
HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL (UNII: 3W1JG795YI)
BISABOLOL (UNII: 24WE03BX2T)
XANTHAN GUM (UNII: TTV12P4NEE)
SCLEROTIUM GUM (UNII: 2X51AD1X3T)
ISOSTEARIC ACID (UNII: X33R8U0062)
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)
CAPRYLYL/CAPRYL GLUCOSIDE (UNII: E00JL9G9K0)
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)
GLYCERIN (UNII: PDC6A3C0OX)
WATER (UNII: 059QF0KO0R)
CETEARYL ALCOHOL (UNII: 2DMT128M1S)
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)
.BETA.-SITOSTEROL (UNII: S347WMO6M4)
CI 77492 (UNII: EX438O2MRT)
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q0OK5DOT4)
ORYZA SATIVA (RICE) STARCH (UNII: 4DGK8B7I3S)
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)
ARGANIA SPINOSA KERNEL OIL (UNII: 4V59G5UW9X)
SODIUM PHYTATE (UNII: 88496G1ERL)
DIMETHICONE (UNII: 92RU3N3Y1O)
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)
CI 77491 (UNII: 1K09F3G675)
SQUALENE (UNII: 7QVM220FJH)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4273-2	1 in 1 CARTON	08/07/2025	
1	NDC:62742-4273-1	50 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/07/2025	

Labeler - Allure Labs (926831603)

Registrant - Allure Labs (926831603)

Establishment

Name	Address	ID/FEI	Business Operations
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Allure Labs		926831603	manufacture(62742-4273)
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Revised: 8/2025

Allure Labs