TRUSKIN SPF 30 MINERAL SUNSCREEN FACIAL SERUM- zinc oxide cream
Allure Labs

Drug Facts

Active ingredients: Zinc Oxide 16.1%

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.

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 ■ Shake well before use ■ Apply generously 15 minutes before sun soposure ■ 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 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.

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

Inactive Ingredients:
Water(Aqua),Caprylic/capric Triglyceride, Isopropyl Isostearate, Polyglyceryl -4 Diisostearate/Polyhydroxy stearate/sebacate, Coconut Alkanes, C13-15 Alkane, Glycerin, Sodium Ascorbyl Phosphate (Vitamin C), Sodium Chloride, Niacinamide, Sodium Hyaluronate, Camellia Sinensis (Green Tea) Leaf Extract, Opuntia Ficus-Indica Stem (Prickly Pear)Extract, Caesalpinia Spinosa (Spiny Holdback) Fruit Pod Extract, Helianthus Annuus (Sunflower) Sprout Extract, Coco-Caprylate/-Caprate, Polyhydroxystearic acid, Polyglyceryl – 3 Polyricinoleate, Isosteric acid, Lecithin, Ceramide NP, Ceramide AP, Ceramide EOP, Xanthan Gum, Maltodextrin, Ethylhexylglycerin, Caprylyl Glycol, Hexylene Glycol, Phenoxyethanol, Sodium Benzoate, Potassium Sorbate, Citric Acid, Phytosphingosine, Cholesterol, Sodium lauroyl lactylate, Carbomer. May Contain +/_ Iron oxides (CI77491, CI77492, CI 77499)

TRUSKIN SPF 30 MINERAL SUNSCREEN FACIAL SERUM

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4274
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	16.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
CAESALPINIA SPINOSA FRUIT POD (UNII: EXY4496LWD)	
HELIANTHUS ANNUUS SPROUT (UNII: 4P26HG1S5W)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q0OK5DOT4)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	

CERAMIDE AP (UNII: F1X8L2B00J)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
CARBOMER (UNII: 0A5MM307FC)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CHOLESTEROL (UNII: 97C5T2UQ7J)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ISOPROPYL ISOSTEARATE (UNII: C67IXB9Y7T)	
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)	
OPUNTIA FICUS-INDICA STEM (UNII: MUD8892KHL)	
CI 77491 (UNII: 1K09F3G675)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CI 77492 (UNII: EX438O2MRT)	
SODIUM LAUROYL LACTYLATE (UNII: 7243K85WFO)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
CITRIC ACID (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
CERAMIDE NP (UNII: 4370DF050B)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
NIACINAMIDE (UNII: 25X51I8RD4)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
CI 77499 (UNII: XM0M87F357)	
PHYTOSPHINGOSINE (UNII: GIN46U9Q2Q)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
C13-15 ALKANE (UNII: 114P5I43UJ)	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4274-2	1 in 1 CARTON	09/05/2025	
1	NDC:62742-4274-1	10 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	09/05/2025	

Labeler - Allure Labs (926831603)

Registrant - Allure Labs (926831603)

Establishment

Name	Address	ID/FEI	Business Operations
Allure Labs		926831603	manufacture(62742-4274)

Revised: 9/2025

Allure Labs