CLEAR CLEAR ZINC SOLAR PROTECTION SPF 30- zinc oxide cream Allure Labs, LLC

Drug Facts

Zinc Oxide: 17%

Purpose: Suncreen

Uses: Help prevent sunburn

Warnings: For external use only

Do not use on damged or broken skin.

Keep out of reach of children. if swalloed, get medical help or contact a Poison Control Center right away.

Directions: Massage into skin with fingertips and reapply as needed. Avoid eye area while applying.

Other Ingredients: Water (Aqua), Cyclopentasiloxane, Dimethicone/PEG10/15 Crosspolymer, Dimethicone/Vinyl Dimethicone Crossploymer, Polyglyceryl-3 polydimethylsiloxyethyl Dimethicone, Cyclomethicone, PEG/PPG-18/18 Dimethicone, Hexyl Triethoxysilylethyl Polydimethylsiloxyethyl Dimethicone, C12-15 Alkyl Benzoate, Polyisobutene, Isododecane, Phenoxyethanol, Sodium Hyaluronate, Caprylyl Glycol, Tocopheryl Acetate, Ethylhexylglycerin, Hexylene Glycol, Mica, Aloe Barbadensis Leaf extract, Camellia Sinensis Leaf Extract, Olea Europaea (Olive) Leaf Extract, Silica

Physicians' Aesthetic Research, Inc. Distr.

San Clemente, CA 92672. Made in U.S.A. www.lucrece.com

A very fine and high quality sun defender, this silicon based formulation is derived from pure and high quality natural resources that provide full spectrum solar protection. Ideal for day to day use, this product is made using the finest cosmetic quality Zinc Oxide, which also acts as a natural anti-irritant in addition to providing outstanding UVA & UVB blocking ability.

Drug Facts	
Active ingredient	Purpose
Zinc Oxide 17%	Sunscreen

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Innovative Skin Technology

CLEAR

CLEAR ZINC SOLAR PROTECTION SPF 30



Net Wt. 2.5 oz. (71 g)

CLEAR CLEAR ZINC SOLAR PROTECTION SPF 30

zinc oxide cream

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:62742-4109 Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	170 mg in 1 g	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)			
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)			
POLYGLYCERYL-3 POLYDIMETHYLSILOXYETHYL DIMETHICONE (4000 MPA.S) (UNII: RLA2U05Z4Q)			
CYCLOMETHICONE (UNII: NMQ347994Z)			

PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)
ISODODECANE (UNII: A8289P68Y2)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
CAPRYLYL GLYCOL (UNII: 00YIU5438U)
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
HEXYLENE GLYCOL (UNII: KEH0A3F75J)
MICA (UNII: V8A1AW0880)
ALOE VERA LEAF (UNII: ZY81Z83H0X)
CAMELLIA SINENSIS FLOWER (UNII: 912BJY2J17)
OLEA EUROPAEA LEAF (UNII: MJ95C3OH47)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742- 4109-1	71 g in 1 TUBE; Type 0: Not a Combination Product	12/04/2017	

Marketing Information				
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M020	12/04/2017		

Labeler - Allure Labs, LLC (926831603)

Registrant - Allure Labs, LLC (926831603)

Establishment				
Name	Address	ID/FEI	Business Operations	
Allure Labs, LLC		926831603	manufacture(62742-4109)	

Revised: 12/2024 Allure Labs, LLC