ELTAMD UV DAILY- octinoxate, zinc oxide sunscreen lotion CP Skin Health Group, Inc

EltaMD UV Daily

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 physician if rash occurs. If product is swallowed get medical help or contact a Poison Control Center right away.

Active ingredients

Octinoxate 7.5% Sunscreen Zinc Oxide 9.0% Sunscreen

Indications & Uses:

Helps prevent sunburn. If used as directed with other sun protection measure (See Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Purpose

Helps prevent sunburn. If used as directed with other sun protection measure (See Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Keep out of reach of children

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Directions:

Apply liberally 15 minutes before sun exposure. Use a water-resistant sunscreen if swimming or sweating. Reapply 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 10 am to 2 pm. Wear long-sleeve shirts, pants, hats, and sunglasses. Children under 6 months: ask a physician.

Other information:

Protect this product from excessive heat and direct sun.

Inactive ingredients

water, petrolatum, isopropyl palmitate, dimethicone, cetearyl alcohol, oleth-3 phosphate, phenoxyethanol, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, polyisobutene, cetearyl glucoside, polyether-1, butylene glycol, PEG-7 trimethylolpropane coconut ether, sodium hyaluronate, iodopropynyl butylcarbamate, tocopheryl acetate, triethoxycaprylylsilane

Questions?

Call toll free 1-800-633-8872

Packaging



ELTAMD UV DAILY

octinoxate, zinc oxide sunscreen lotion

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:72043-6804

Route of Administration

TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	90 g in 1000 g		
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 g in 1000 g		

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PETROLATUM (UNII: 4T6H12BN9U)	
DIMETHICONE (UNII: 92RU3N3Y10)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HYALURONIC ACID (UNII: S270N0TRQY)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)	
PEG-7 TRIMETHYLOLPROPANE COCONUT ETHER (UNII: MVJ3AD73GG)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Product Characteristics					
Color	white	Score			
Shape		Size			
Flavor		Imprint Code			
Contains					

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:72043- 6804-2	2 g in 1 PACKET; Type 0: Not a Combination Product	05/31/2024		
2	NDC:72043- 6804-1	48 g in 1 BOTTLE; Type 0: Not a Combination Product	05/31/2024		

Marketing In	formation		
Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M020	05/31/2024	

Labeler - CP Skin Health Group, Inc (611921669)

Registrant - Swiss-American CDMO, LLC (080170933)

Establishment				
Name	Address	ID/FEI	Business Operations	
Swiss-American CDMO, LLC		080170933	manufacture(72043-6804)	

Revised: 5/2024 CP Skin Health Group, Inc