ELTAMD UV SPF 30 PLUS- zinc oxide and octinoxate sunscreen lotion
CP Skin Health Group, Inc.

EltaMD UV Lotion

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. Keep out of reach of children.

Active Ingredients

Octinoxate 7.5% Sunscreen

Zinc Oxide 7.0% 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.

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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 another sun protection measures including: limit time in the sun, expecially from 10 am - 2 pm wear long-sleeve shirts, pants, hats and sunglasses Children under 6 months: Ask a physician

Inactive Ingredients

purified water, petrolatum, isopropyl palmitate, octyl stearate, glyceryl stearate, cetearyl glucoside, dimethicone, PEG-100 stearate, hydroxyethyl acrylate/sodium acryloyldimethyl tautrate copolymer, polyisobutene, PEG-7 trimethylolpropane coconut ether, sodium hyaluronate, tocopheryl acetate, ploye ther-1, citric acid, oleth-3 phosphate, phenoxyethanol, butylene glycol, iodopropynyl butylcarbamate, triethoxycaprylylsilane

KEEP OUT OF REACH OF CHILDREN

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Labeling

UVLotion7ozBox080510D.pdf 1 11/12/20 11:36 AM

7oz Pump Bottle Box
Straight Tuck
080510D Dieline

ELTAMD UV SPF 30 PLUS

zinc oxide and octinoxate sunscreen lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72043-2286
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	70 g in 1000 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 g in 1000 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
PETROLATUM (UNII: 4T6H12BN9U)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
OCTYL STEARATE (UNII: 772Y4UFC8B)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
PEG-100 STEARATE (UNII: YD01N1999R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72043-	198 g in 1 BOTTLE; Type 0: Not a Combination	01/10/2018	

1	2286-8	Product	01/10/2018	
2	NDC:72043-2286-2	2 g in 1 PACKET; Type 0: Not a Combination Product	07/06/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/10/2018	

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(72043-2286)

Revised: 11/2023 CP Skin Health Group, Inc.