UV CLEAR DEEP TINT- octinoxate, zinc oxide sunscreen lotion
CP Skin Health Group, Inc.

EltaMD UV Clear Deep Tint

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

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Uses

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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.

Active Ingredients

Octinoxate 7.5%

Zinc Oxide 9.0%

Inactive Ingredients

water, cyclopentasiloxane, niacinamide, octyldodecyln eopentanoate, hydroxyethyl acrylate/sodium acryloldimethyl taurate copolymer, butylee glycol, isopropyl palmitate,

phenoxyethanol, polyisobutene, ethylheyl stearate, triethoxycaprylylsilane, tocopheryl acetate, PEG-7 trimethylolpropane coconut ether, oleth-3 phosphate, iodopropynyl butylcarbamate, latic acid, sodiym hyaluroate, iron oxides

Questions

Call toll free 1-800-633-8872

Labeling

UV CLEAR DEEP TINT

octinoxate, zinc oxide sunscreen lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72043-2530

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	750 g in 1000 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	900 g in 1000 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
LACTIC ACID (UNII: 33X04XA5AT)	
HYALURONIC ACID (UNII: S270N0TRQY)	
NIACINAMIDE (UNII: 25X51I8RD4)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)	
PEG-7 TRIMETHYLOLPROPANE COCONUT ETHER (UNII: MVJ3AD73GG)	

Product Characteristics

Color	brown	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72043-2530-1	50 g in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2024	
2	NDC:72043-2530-2	2 g in 1 PACKET; Type 0: Not a Combination Product	05/08/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	05/08/2024	

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(72043-2530)

Revised: 5/2024 CP Skin Health Group, Inc.