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

EltaMD UV Daily 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

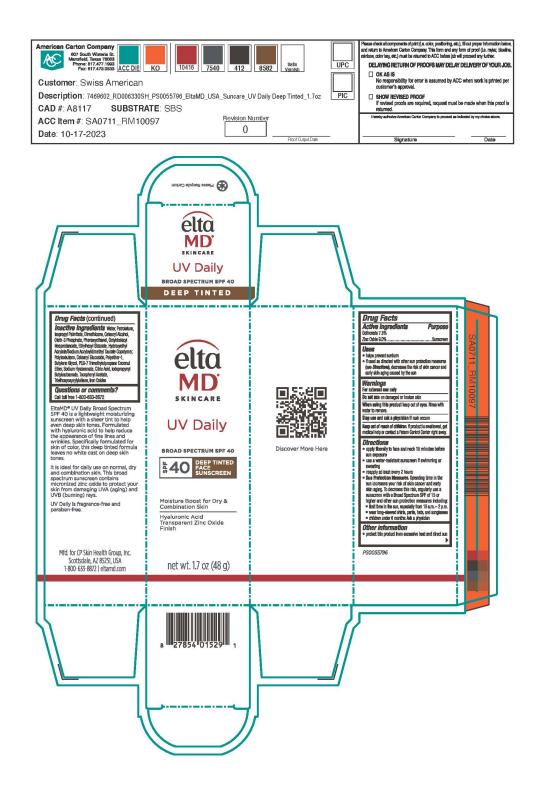
water, petrolatum, isopropyl palmitate, dimethicone, cetearyl alcohol, oleth-3 phosphate, phenoxyethanol, octyldodecyl neopentanoate, ethylhexyl stearate, hydroxyethyl

acrylate/sodium acryloyldimethyl taurate copolymer, polyisobutene, cetearyl glucoside, polyether-1, butylene glycol, PEG-7 trimethylolpropane cocnut ether, sodium hyaluronate, citric acid, iodopropynyl butylcarbamate, tocopheryl acetate, triethoxycaprylylsilane, iorn oxides

Questions

Call toll free 1-800-633-8872

Labeling



UV DAILY DEEP TINT

octinoxate, zinc oxide sunscreen lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72043-2920

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)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PETROLATUM (UNII: 4T6H12BN9U)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
HYALURONIC ACID (UNII: S270N0TRQY)	
PEG-7 TRIMETHYLOLPROPANE COCONUT ETHER (UNII: MVJ3AD73GG)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	

Product Characteristics			
Color	brown	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72043-	50 g in 1 BOTTLE; Type 0: Not a Combination	05/00/2024	

-	2920-1	Product	05/09/2024	
2	NDC:72043- 2920-2	2 g in 1 PACKET; Type 0: Not a Combination Product	05/09/2024	
Marketing Information				
M	larketing	nformation		
M	larketing Marketing Category	nformation Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

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(72043-2920)	

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