

ELTAMD UV DAILY TINTED SPF40- zinc oxide and octinoxate sunscreen lotion
CP Skin Health Group, Inc.

EltAMD UV Daily Tinted SPF40

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 physician if rash occurs Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Active Ingredients

Zinc Oxide 9.0% Sunscreen

Octinoxate 7.5% 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 to face, neck and backs of hands 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 a.m. - 2 p.m. wear long-sleeve shirts, pants, hats and sunglasses children under 6 months: Ask a physician

Inactive Ingredients

purified water, petrolatum, isopropyl palmitate, cetearyl glucoside, dimethicone hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, polyisobutene, PEG-7 trimethylolpropane coconut ether, sodium hyaluronate, tocopheryl acetate, polyether-1, citric acid, oleth-3 phosphate, phenoxyethanol, butylene glycol, idodpropynyl butylcarbamate, triethoxycaprylylsilane, iron oxides, ethylhexyl stearate, octyldodecyl neopentanoate

KEEP OUT OF REACH OF CHILDREN

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Labeling

CAD #: A8117 SUBSTRATE: SBS

ACC Item #: SA0644

Date: 10-10-2022

Revision Number

0

Proof Output Date

I hereby authorize American Carton Company to proceed as indicated by my choice above.

Signature

Date



Drug Facts (continued)

Inactive ingredients: Water, Petroleum, Isopropyl Palmitate, Dimethicone, Cetearyl Glucoside, Oleth-3 Phosphate, Phenoxethanol, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Iron Oxides, Polyisobutene, Octyldodecyl Neopentanoate, Ethylhexyl Stearate, Polyether-1, Butylene Glycol, Cetyl Alcohol, Stearyl Alcohol, PEG-7 Trimethylpropylsorbate, Coconut Ethyl Phosphate, Sodium Hyaluronate, Citric Acid, Isododecyl Dimethylcarbamate, Tocopheryl Acetate, Triethoxycaprylylsilane

Questions or comments?
Call toll free 1-800-633-8872

EltaMD® UV Daily Broad Spectrum SPF 40 is a lightweight moisturizing sunscreen with a sheer tint to help even skin tone. Formulated with hyaluronic acid to help reduce the appearance of fine lines and wrinkles.

It is ideal for daily use on normal, dry and combination skin. This broad spectrum sunscreen contains micronized zinc oxide to protect your skin from damaging UVA (aging) and UVB (burning) rays.

UV Daily is fragrance-free, paraben-free, and non-comedogenic.

Mfd. for CP Skin Health Group, Inc.
Scottsdale, AZ 85251, USA
1-800-633-8872 | eltamd.com

BROAD SPECTRUM SPF 40
SPL 40 TINTED FACE SUNSCREEN

Moisture Boost for Dry & Combination Skin
Hyaluronic Acid
Transparent Zinc Oxide Finish

net wt. 1.7 oz (48 g)



Please Recycle Carton

TINTED
BROAD SPECTRUM SPF 40

UV Daily
SKINCARE
elta MD

elta MD
SKINCARE

UV Daily



Discover More Here



The Skin Cancer Foundation recommends this product as an effective broad spectrum sunscreen.

Drug Facts

Active ingredients Purpose
Octinoxate 7.5%, Zinc Oxide 9.0%.....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

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

Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally to face and neck 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 a.m. – 2 p.m.
 - wear long-sleeved shirts, pants, hats, and sunglasses
 - children under 6 months: Ask a physician

Other Information

protect this product from excessive heat and direct sun

RM8510

SA0644

ELTAMD UV DAILY TINTED SPF40

zinc oxide and octinoxate sunscreen lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72043-2269
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)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
OCTYL STEARATE (UNII: 772Y4UFC8B)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PETROLATUM (UNII: 4T6H12BN9U)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
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)	

Product Characteristics

Color	brown	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72043-2269-1	48 g in 1 BOTTLE; Type 0: Not a Combination Product	01/10/2018	

2	NDC:72043-2269-2	2 g in 1 PACKET; Type 0: Not a Combination Product	07/06/2022
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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(72043-2269)

Revised: 11/2023

CP Skin Health Group, Inc.