ELTAMD UV SHEER- zinc oxide and octocrylene sunscreen cream CP Skin Health Group, Inc.

EltaMD UV Sheer

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

Octocrylene 10% Sunscreen Zinc Oxide 15% Sunscreen

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.

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.

Keep out of reach of children

Keep out of reach of children

Directions

Apply liberally 15 minutes before sun exposure. Reapply after 80 minutes of swimming or sweating, immediately after towel drying, 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.

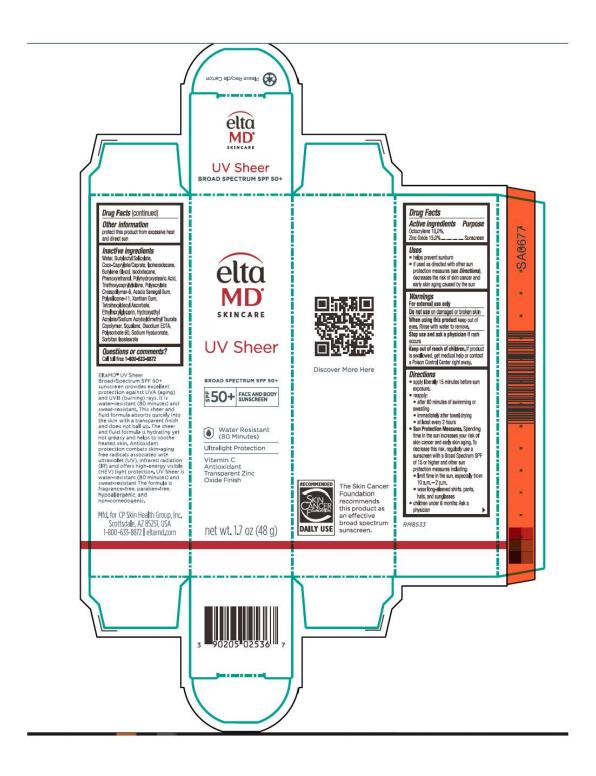
Inactive ingredients

water, butyloctyl salicylate, coco-caprylate/caprate, isohexadecane, butylene glycol, isododecane, phenoxyethanol, polyhydroxystearic acid, triethoxycaprylylsilane, polyacrylate crosspolymer-6, acacia Senegal gum, polysilicone-11, xanthan gum, tetrahexyldecyl ascorbate, ethylhexylglycerin, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, squalene, disodium EDTA, polysorbate 60, sodium hydroxide

Questions?

Call toll free 1-800-633-8872

Labeling



zinc oxide and octocrylene sunscreen cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72043-2563

Route of Administration TOPICAL

(45000 MPA.S) (UNII: Q7UI015FF9)

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	150 g in 1000 g		
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 a in 1000 a		

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYALURONIC ACID (UNII: S270N0TRQY)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
ISODODECANE (UNII: A8289P68Y2)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
ACACIA SENEGAL FLOWER (UNII: 72P931MTC2)	
XANTHAN GUM (UNII: TTV12P4NEE)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
SQUALANE (UNII: GW89575KF9)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED	

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- 2563-1	30 g in 1 TUBE; Type 0: Not a Combination Product	11/03/2020	11/09/2023
2	NDC:72043- 2563-7	50 g in 1 TUBE; Type 0: Not a Combination Product	11/03/2020	
3	NDC:72043- 2563-0	10 g in 1 TUBE; Type 0: Not a Combination Product	11/03/2020	
4	NDC:72043- 2563-2	2 g in 1 PACKET; Type 0: Not a Combination Product	11/03/2020	11/09/2023
5	NDC:72043- 2563-3	85 g in 1 TUBE; Type 0: Not a Combination Product	07/07/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/30/2020	

Labeler - CP Skin Health Group, Inc. (611921669)

Registrant - Swiss-American CDMO, LLC (080170933)

Establishment				
Na me	Address	ID/FEI	Business Operations	
Swiss-American CDMO, LLC		080170933	manufacture(72043-2563)	

Revised: 12/2023 CP Skin Health Group, Inc.