ELTAMD UV PURE SPF47- zinc oxide and titanium dioxide sunscreen cream CP Skin Health Group, Inc.

EltaMD UV Pure SPF47

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.

Active Ingredients

Zinc Oxide 10.0% Sunscreen
Titanium Dioxide 5.5% Sunscreen

Uses

helps prevents 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

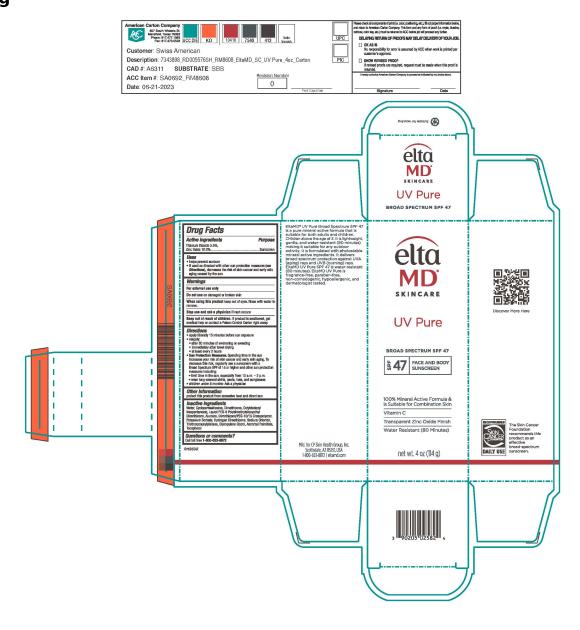
Apply liberally 15 minutes before sun exposure Reapply after 80 minutes of swimming or sweating immediately after towel drying at least every 2 hours Sun Protection Measures. Spending time int he sun increases your risk of skin cancer and early skin aging. To decrease this risk, egularly use a sunscreen 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, Cyclopentasiloxane, Dimethicone, Octyldodecyl neopentanoate, laury PEG-9 Polydimethylsiloxyethyl Dimethicone, Dimethicone/PEG-10/15 Crosspolymer, Ascorbyl Palmitate, Tocopherol, Sodium Chloride, Triethoxycaprylylsilane, Alumina, Hydrogen Dimethicone, Potassium Sorbate, Citric Acid

KEEP OUT OF REACH OF CHILDREN

Labeling



ELTAMD UV PURE SPF47 zinc oxide and titanium dioxide sunscreen cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:72043-2582 Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	100 g in 1000 g	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	55 g in 1000 g	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
DIMETHICONE (UNII: 92RU3N3Y10)			
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)			
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)			
ASCORBYL PALMITATE (UNII: QN83US2B0N)			
LAURYL PEG-9 POLYDIMETHYLSILOXYETHYL DIMETHICONE (UNII: 25G622K2RA)			
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)			
HYDROGEN DIMETHICONE (13 CST) (UNII: 4QGR4P2YOI)			
ALUMINUM OXIDE (UNII: LMI26O6933)			

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:72043- 2582-4	114 g in 1 BOTTLE; Type 0: Not a Combination Product	01/10/2018		
2	NDC:72043- 2582-2	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(72043-2582)		

Revised: 3/2024 CP Skin Health Group, Inc.