

**ELTAMD UV SPORT SPF50- zinc oxide, octinoxate and octisalate
sunscreen lotion
CP Skin Health Group, Inc.**

EltAMD UV Sport SPF50

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 contact a Poison Control Center right away.

Active ingredients

Zinc Oxide 9.0% Sunscreen
Octinoxate 7.5% Sunscreen
Octisalate 5.0% 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.

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 10a.m.-2p.m. wear long-sleeve shirts, pants, hats and sunglasses Children under 6 months: Ask a physician

Doseage and administration

Apply liberally 15 minutes before sun exposure Reapply after 80 minutes of swimming or sweating immediately after towel drying at least every 2 hours








Inactive ingredients

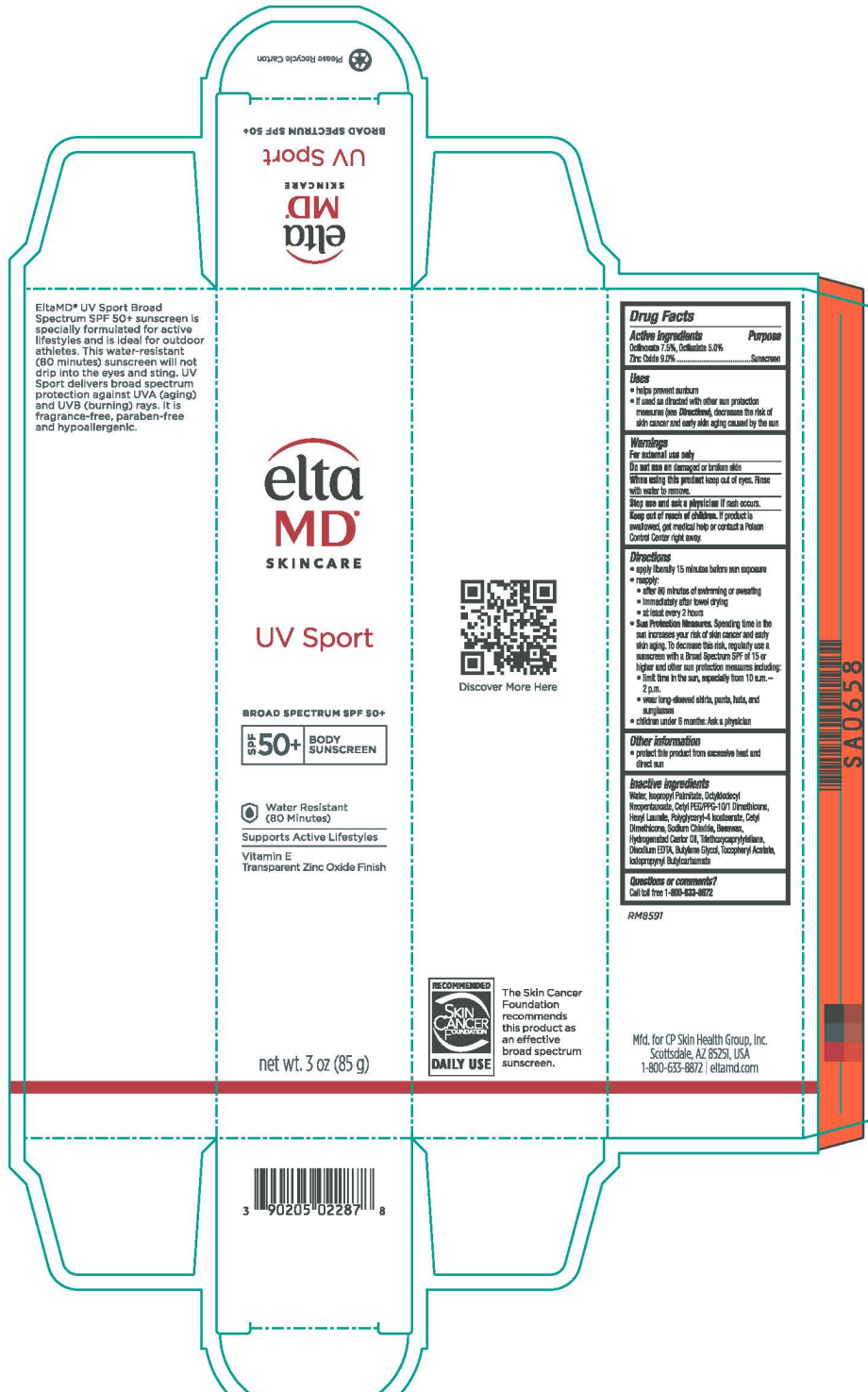
Beeswax, Butylene Glycol, Cetyl Dimethicone, Cetyl PEG/PPG-10/1 Dimethicone, Disodium EDTA, Hexyl Laurate, Hydrogenated Castor Oil, Iodopropynyl Butylcarbamate, Isopropyl Palmitate, Octyldoecyl Neopentanoate, Polyglyceryl-4 Isostearate, Purified Water, Sodium Chloride, Tocopheryl Acetate, Triethoxycaprylylsilane

Keep out of reach of children

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Labeling

American Carton Company  807 South Wisteria St. Mesquite, Texas 78063 Phone: 817.477.1982 Fax: 817.473.0533	 ACC DIE	 KO	 10416	 7540	 412	 Satin Varnish	<input type="checkbox"/> UPC <input type="checkbox"/> PIC	Please check all components of print (i.e. color, positioning, etc.). If out proper information below, and return to American Carton Company. This form and any form of proof (i.e. raster, halftone, inkjet, color key, etc.) must be returned to ACC before job will proceed any further. DELAYING RETURN OF PROOFS MAY DELAY DELIVERY OF YOUR JOB. <input type="checkbox"/> OK AS IS No responsibility for error is assumed by ACC when work is printed per customer's approval. <input type="checkbox"/> SHOW REVISED PROOF If revised proofs are required, request must be made when this proof is returned. I hereby authorize American Carton Company to proceed as indicated by my choice above.
	Customer: Swiss American Description: 7027632_RM8591 CAD #: A9658 SUBSTRATE: SBS ACC Item #: SA0658 Date: 11-15-2022	Revision Number 0	Proof Output Date	Signature _____ Date _____				





ELTAMD UV SPORT SPF50

zinc oxide, octinoxate and octisalate sunscreen lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72043-2287
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
octisalate (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	octisalate	50 g in 1000 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
CETYL DIMETHICONE 150 (UNII: 5L694Y0T22)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 1.5) (UNII: V2W71V8T0X)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72043-2287-3	84 g in 1 TUBE; Type 0: Not a Combination Product	01/10/2018	
2	NDC:72043-2287-9	226 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2024	
3	NDC:72043-2287-8	198 g in 1 BOTTLE; Type 0: Not a Combination Product	01/10/2018	05/30/2024
4	NDC:72043-2287-2	2 g in 1 PACKET; Type 0: Not a Combination Product	07/07/2022	
5	NDC:72043-2287-6	1815 g in 1 BOTTLE; Type 0: Not a Combination Product	07/07/2022	09/11/2023

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(72043-2287)

Revised: 5/2024

CP Skin Health Group, Inc.