# ELTAMD UV LUMINOUS- titanium dioxide, zinc oxide sunscreen lotion CP Skin Health Group, Inc

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#### **EltaMD UV Luminous**

## **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.

#### 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.

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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.

# Keep out of reach of children

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## **Active Ingredients**

Titanium Dioxide 7.2% Sunscreen Zinc Oxide 9.0% Sunscreen

### Other information:

Protect this product from excessive heat and direct sun.

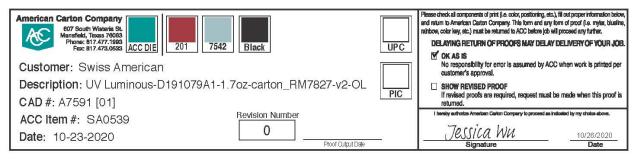
## **Inactive Ingredients**

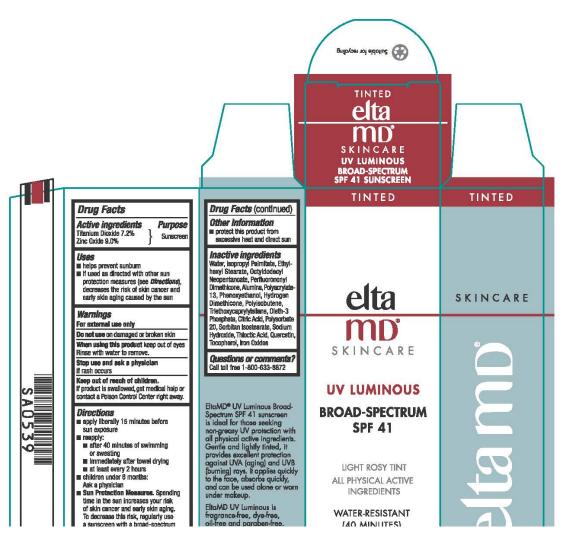
water, isopropyl palmitate, ethylhexyl stearate, octyldodecyl neopentanoate, perfluorononly dimethicone, alumina, polyacrylate-13, phenoxyethanol, hydrogen dimethicone, polyisobutene, triethoxycaprylylsilane, oleth-3 phosphate, citric acid, polysorbate 20, sorbitan isostearate, sodium hydroxide, thioctic acid, quercetin, tocopherol, iron oxides

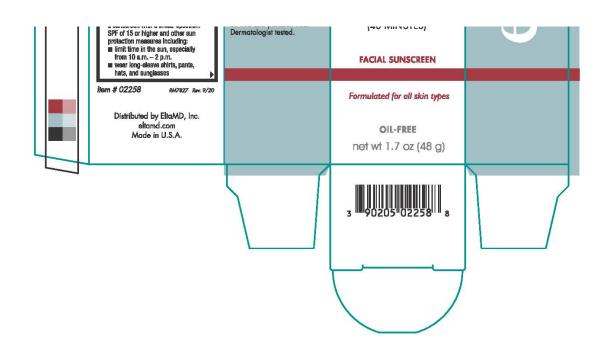
#### **Questions?**

Call toll free 1-800-633-8872

## Labeling







# **ELTAMD UV LUMINOUS**

titanium dioxide, zinc oxide sunscreen lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72043-2256
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	72 g in 1000 g		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	90 g in 1000 g		

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)			

ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M) OCTYL STEARATE (UNII: 772Y4UFC8B) OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T) OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL) FERRIC OXIDE YELLOW (UNII: EX43802MRT) FERRIC OXIDE RED (UNII: 1K09F3G675) FERROSOFERRIC OXIDE (UNII: XM0M87F357) TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E) **ALUMINUM OXIDE** (UNII: LMI26O6933) HYDROGEN DIMETHICONE (13 CST) (UNII: 4QGR4P2YOI) POLYSORBATE 20 (UNII: 7T1F30V5YH) POLYISOBUTYLENE (1000 MW) (UNII: 5XB3A63Y52) **ALPHA LIPOIC ACID** (UNII: 73Y7P0K73Y) QUERCETIN (UNII: 9IKM0I5T1E) **BUTYLENE GLYCOL** (UNII: 3XUS85K0RA) **SORBITAN ISOSTEARATE** (UNII: 01S2G2C1E4) TOCOPHEROL (UNII: R0ZB2556P8)

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- 2256-1	10 g in 1 TUBE; Type 0: Not a Combination Product	12/09/2020	11/14/2023	
2	NDC:72043- 2256-7	48 g in 1 TUBE; Type 0: Not a Combination Product	12/09/2020		
3	NDC:72043- 2256-3	30 g in 1 TUBE; Type 0: Not a Combination Product	12/09/2020	11/14/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/09/2020	

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

# Registrant - Swiss-American CDMO, LLC (080170933)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>

CP Skin Health Group, Inc Revised: 11/2023