

**APOSTROPHE SCREEN SPF43- sunscreen lotion**  
**Yoderm, Inc.**

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**Apostrophe Screen SPF43**

**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**

Zinc Oxide 7.5% Sunscreen

Octinoxate 7.5% Sunscreen

Octisalate 2.5% 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.

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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.

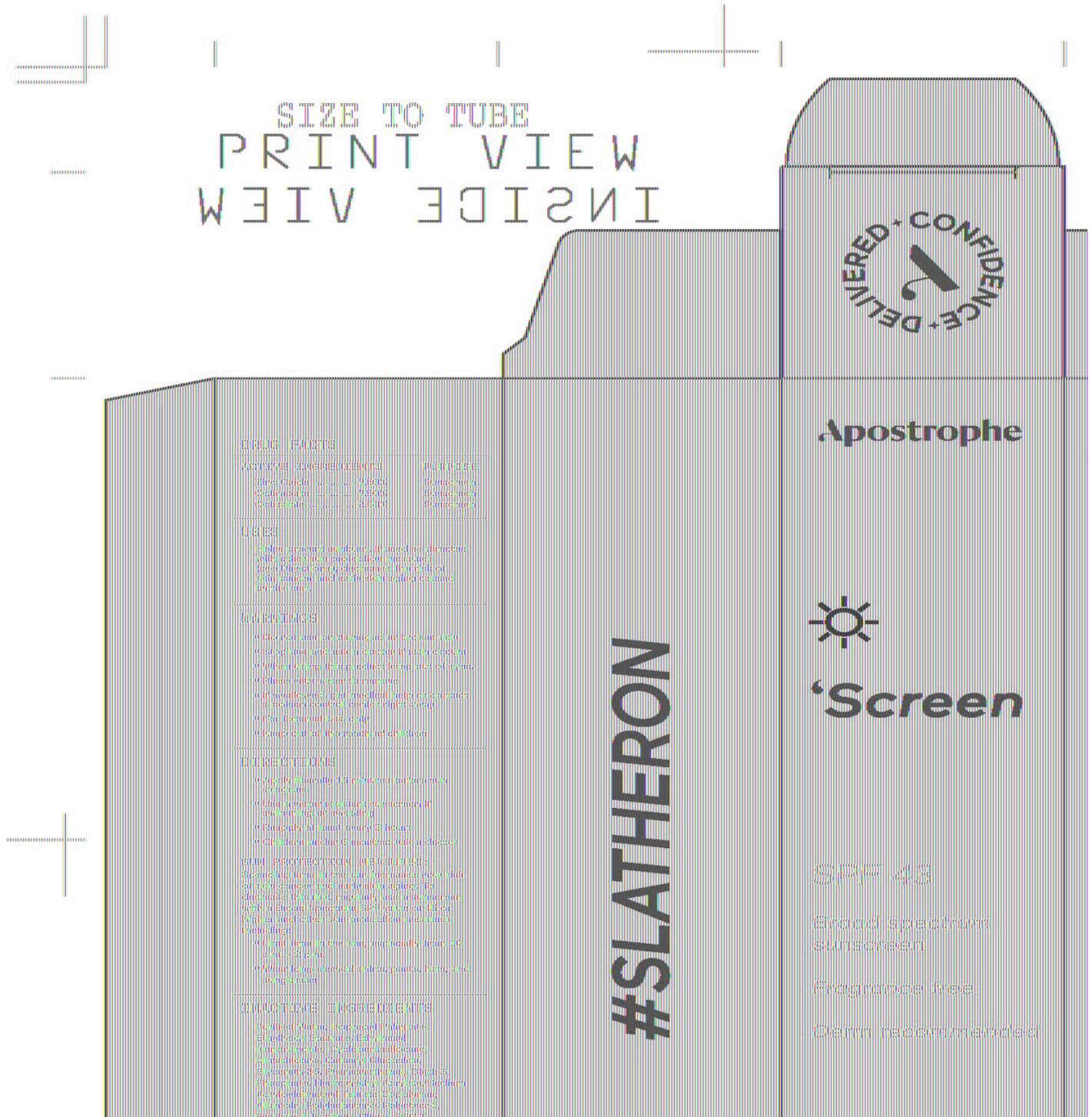
**Other Information**

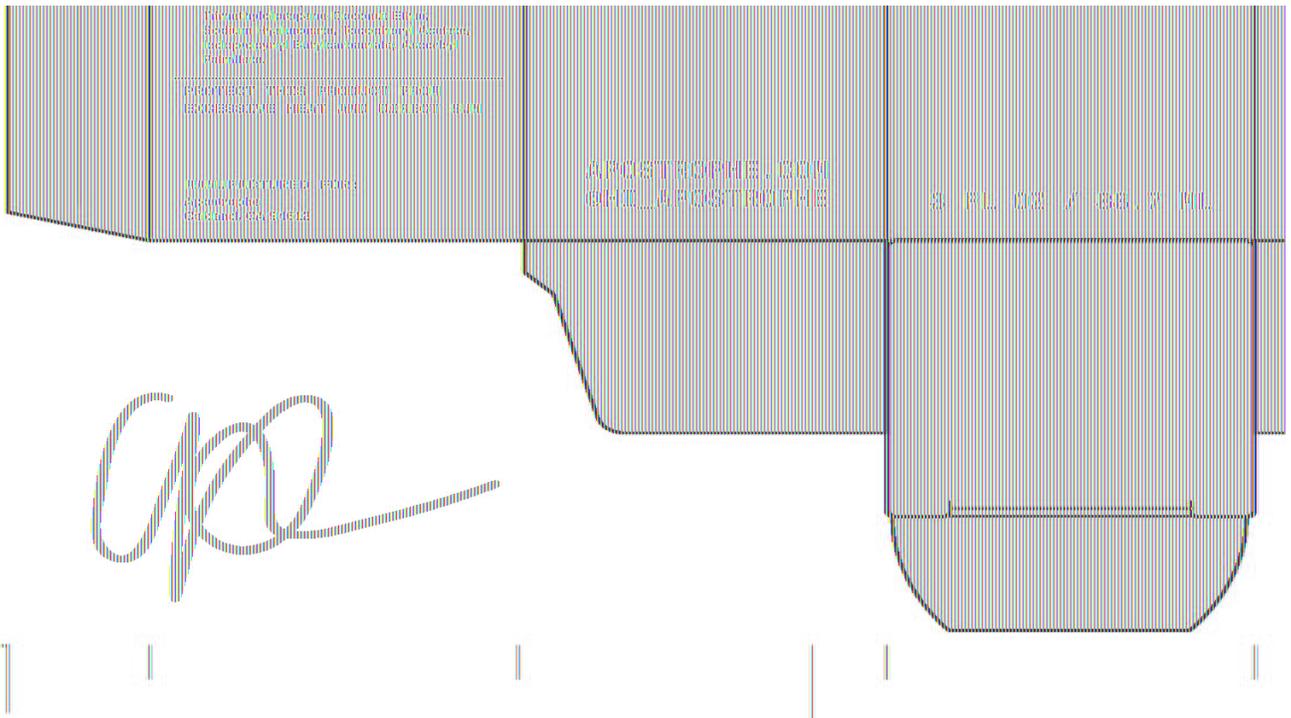
Protect this product from excessive heat and direct sun.

## Inactive Ingredients

purified water, isopropyl palmitate, ethylhexyl stearate, ethylhexyl isononanoate, cyclopentasiloxane, dimethicone, cetearyl glucoside, glycereth-26, phenoxyethanol, oleth-3 phosphate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, allantoin, polyisobutene, polyether-1, panthenol, butylene glycol, PEG-7 trimethylolpropane coconut ether, sodium hyaluronate, tocopheryl acetate, iodopropynyl butylcarbamate, ascorbyl palmitate, triethoxycaprylylsilane

## Labeling





## APOSTROPHE SCREEN SPF43

sunscreen lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:81780-007
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	25 g in 1000 g
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 g in 1000 g
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	75 g in 1000 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)	
<b>ETHYLHEXYL STEARATE</b> (UNII: EG3PA2K3K5)	
<b>ETHYLHEXYL ISONONANOATE</b> (UNII: I6KB4GE3K4)	
<b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>CETEARYL GLUCOSIDE</b> (UNII: 09FUA47KNA)	
<b>GLYCERETH-26</b> (UNII: NNE56F2N14)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>OLETH-3 PHOSPHATE</b> (UNII: 8Q0Z18J1VL)	
<b>HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%)</b> (UNII: 86FQE96TZ4)	

<b>ALLANTOIN</b> (UNII: 344S277G0Z)	
<b>POLYISOBUTYLENE (1000 MW)</b> (UNII: 5XB3A63Y52)	
<b>PANTHENOL</b> (UNII: WW9CM0067Z)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>HYALURONIC ACID</b> (UNII: S270N0TRQY)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>IODOPROPYNYL BUTYLCARBAMATE</b> (UNII: 603P14DHEB)	
<b>ASCORBYL PALMITATE</b> (UNII: QN83US2B0N)	
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81780-007-03	1 in 1 CARTON	04/09/2021	
1		87 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:81780-007-02	2 g in 1 PACKET; Type 0: Not a Combination Product	04/09/2021	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/09/2021	

**Labeler** - Yoderm, Inc. (079220017)

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

### Establishment

Name	Address	ID/FEI	Business Operations
Swiss-American CDMO, LLC		080170933	manufacture(81780-007)