

**SHEER MINERAL ANTIOXIDANT SUNSCREEN SPF50- zinc oxide lotion**  
Topiderm, Inc.

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**Sheer Mineral Antioxidant Sunscreen**  
**SPF50**

***Drug Facts***

**Active ingredient**

Zinc Oxide 20.5%

**Purpose**

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.

**Warnings**

**For external use only.**

- **When using this product** keep out of eyes, rinse with water to remove.
- **Stop use if** signs of irritation or rash appear. If irritation or rash persists consult a doctor.
- **Keep out of reach of children.**
- In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.
- **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 sunscreen measures including:

- limit time in the sun, especially from 10a.m.-2 p.m.
- wear long-sleeve shirts, pants, hats, and sunglasses

**Directions**

- Apply liberally and spread evenly by hand 15 minutes before sun exposure
- Re-apply after swimming, excessive perspiring, or anytime after towel drying
- Use a water resistant sunscreen if swimming or sweating
- Use on children under 6 months of age: consult a doctor.

## **Inactive ingredients**

Acrylates Crosspolymer-4, Ascorbic Acid, Ascorbyl Palmitate, Biotin, C12-15 Alkyl Benzoate, C13-15 Alkane, Camellia Sinensis (Green Tea) Polyphenols, Cetearyl Alcohol, Ceteth-10 Phosphate, Dicapryl Phosphate, Disodium EDTA, Isostearic Acid, Lauryl PEG/PPG-18/18 Methicone, Phenoxyethanol, Phospholipids, Phyllanthus Emblica Fruit Extract, Polyhydroxystearic Acid, Potassium Cetyl Phosphate, Potassium Hydroxide, Purified Water, Resveratrol, Retinyl Palmitate, Tocopheryl Acetate, Triethoxycaprylylsilane, Ubiquinone, Xanthan Gum.

## **PRINCIPAL DISPLAY PANEL - 57 g Container Label**

COMPLIMENTS OF

TOPIX  
PHARMACEUTICALS, INC

DAILY

Sheer Mineral  
Antioxidant  
Sunscreen SPF 50

Broad Spectrum SPF 50

Net wt. 2 oz. (57 g)

Available  
Custom Branded  
800.445.2595  
c.service@topixpharm.com

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R0424

Made in U.S.A.

1440MB

### Drug Facts (continued)

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# SHEER MINERAL ANTIOXIDANT SUNSCREEN SPF50

zinc oxide lotion

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51326-144
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	205 mg in 1 g

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>ACRYLATES CROSSPOLYMER-4</b> (UNII: GEV2EL4D9G)	
<b>ASCORBIC ACID</b> (UNII: PQ6CK8PD0R)	
<b>ASCORBYL PALMITATE</b> (UNII: QN83US2B0N)	
<b>BIOTIN</b> (UNII: 6SO6U10H04)	
<b>ALKYL (C12-15) BENZOATE</b> (UNII: A9EJ3J61HQ)	
<b>C13-15 ALKANE</b> (UNII: 114P5I43UJ)	
<b>GREEN TEA LEAF</b> (UNII: W2ZU1RY8B0)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>CETETH-10 PHOSPHATE</b> (UNII: 4E05O5N49G)	
<b>DIHEXADECYL PHOSPHATE</b> (UNII: 2V6E5WN99N)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>ISOSTEARIC ACID</b> (UNII: X33R8U0062)	
<b>LAURYL PEG/PPG-18/18 METHICONE</b> (UNII: ZJ5S27D9NX)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>HYDROGENATED SOYBEAN LECITHIN</b> (UNII: H1109Z9J4N)	
<b>PHYLLANTHUS EMBLICA FRUIT</b> (UNII: YLX4CW2576)	
<b>POLYHYDROXYSTEARIC ACID (2300 MW)</b> (UNII: YXH47AOU0F)	
<b>POTASSIUM CETYL PHOSPHATE</b> (UNII: 03KCY6P7UT)	
<b>POTASSIUM HYDROXIDE</b> (UNII: WZH3C48M4T)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>RESVERATROL</b> (UNII: Q369O8926L)	
<b>VITAMIN A PALMITATE</b> (UNII: 1D1K0N0VVC)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)	
<b>UBIDECARENONE</b> (UNII: EJ27X76M46)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51326-144-01	57 g in 1 CONTAINER; Type 0: Not a Combination Product	04/25/2024	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M020	04/25/2024	

**Labeler** - Topiderm, Inc. (049121643)**Registrant** - Topiderm, Inc. (049121643)**Establishment**

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-144)

**Establishment**

Name	Address	ID/FEI	Business Operations
Topix Pharmaceuticals, Inc.		117745066	PACK(51326-144)

Revised: 3/2025

Topiderm, Inc.