REPLENIX SUNSCREEN SPF 50- zinc oxide lotion Topiderm, Inc.

Replenix® Sunscreen SPF 50

Drug Facts

| Active ingredient | Purpose | |
|-------------------|-----------|--|
| Zinc Oxide 14.5% | Sunscreen | |
| Octinoxate 7.5% | 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 10 a.m.-2 p.m.
- wear long-sleeve shirts, pants, hats, and sunglasses

Directions

- Apply liberally to face and neck and spread evenly 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

Purified Water, Niacinamide, Oleth-3 Phosphate, Neopentyl Glycol Diheptanoate,

Polyisobutene, Octyldodecyl Neopentanoate, Butylene Glycol, Tocopheryl Acetate, Sodium Hyaluronate, Caffeine, Panthenol, Dimethicone, Sucrose, Lactic Acid, Glycerin, Ethylhexyl Stearate, Triethoxycaprylylsilane, Xanthan Gum, Lauryl PEG Polydimethylsiloxyethyl Dimethicone, PEG-7 Trimethylolpropane Coconut Ether, Hydroxyethyl Acrylate/Sodium Acryloydimethyl Taurate Copolymer, Phenoxyethanol, Disodium EDTA.

PRINCIPAL DISPLAY PANEL - 63 g Bottle Label

REPLENIX® SUNSCREEN

ESSENTIAL OIL FREE SHEER SPF 50

14.5% Micronized Zinc Oxide Multi-tasking sheer application Sensitive skin friendly

BROAD SPECTRUM UVA/UVB SPF 50

Net wt. 2.22 oz. (63 g)

TOPIX PHARMACEUTICALS, INC. N. AMITYVILLE, NY 11701



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Drug Facts (continued)

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1200MB

swimming or sweating Use on children under 6 months of age: consult a doctor.

Inactive ingredients

Purified Water, Niacinamide, Oleth-3
Phosphate, Neopentyl Glycol Diheptanoate,
Polyisobutene, Octyldodecyl Neopentanoate,
Butylene Glycol, Tocopheryl Acetate, Sodium
Hyaluronate, Caffeine, Panthenol,
Dimethicone, Sucrose, Lactic Acid, Glycerin,
Ethylhexyl Stearate, Triethoxycaprylylsilane,
Xanthan Gum, Lauryl PEG
Polydimethylsiloxyethyl Dimethicone, PEG-7
Trimethylolpropane Coconut Ether,
Hydroxyethyl Acrylate/Sodium
Acryloydimethyl Taurate Copolymer,
Phenoxyethanol, Disodium EDTA.

REPLENIX SUNSCREEN SPF 50

zinc oxide lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:51326-220

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|--------------------------|---------------|
| ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) | ZINC OXIDE | 145 mg in 1 g |
| OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51) | OCTINOXATE | 75 mg in 1 g |

| Inactive | Ingredi | ents |
|----------|---------|------|
| | | |

Ingredient Name Strength
WATER (UNII: 059QF0KO0R)

NIACINAMIDE (UNII: 25X51I8RD4)

| OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL) | |
|---|--|
| NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X) | |
| OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T) | |
| BUTYLENE GLYCOL (UNII: 3XUS85K0RA) | |
| .ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| HYALURONATE SODIUM (UNII: YSE9PPT4TH) | |
| CAFFEINE (UNII: 3G6A5W338E) | |
| PANTHENOL (UNII: W/9CM0067Z) | |
| DIMETHICONE (UNII: 92RU3N3Y1O) | |
| SUCROSE (UNII: C151H8M554) | |
| LACTIC ACID, DL- (UNII: 3B8D35Y7S4) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5) | |
| TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E) | |
| XANTHAN GUM (UNII: TTV12P4NEE) | |
| PEG-7 TRIMETHYLOLPROPANE COCONUT ETHER (UNII: MVJ3AD73GG) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | |
| | |

| l | Packaging | | | | |
|---|---------------------------------|----------------------|---|-----------------------|--|
| | # Item Code Package Description | | Marketing Start Date | Marketing End Date | |
| | 1 | NDC:51326-220- 01 | 63 g in 1 BOTTLE; Type 0: Not a Combination Product | 10/16/2021 | |

| Marketing Information | | | |
|--|------|-------------------------|-----------------------|
| Marketing Application Number or Monogra Category Citation | | Marketing Start Date | Marketing End Date |
| OTC monograph drug | M020 | 10/16/2021 | |
| | | | |

Labeler - Topiderm, Inc. (049121643)

Registrant - Topiderm, Inc. (049121643)

| Establishment | | | | | |
|----------------|---------|-----------|------------------------|--|--|
| Name | Address | ID/FEI | Business Operations | | |
| Topiderm, Inc. | | 049121643 | MANUFACTURE(51326-220) | | |

Revised: 2/2023 Topiderm, Inc.