

CLARITY RX PHYSICAL SKIN DEFENSE SPF 50- zinc oxide and titanium dioxide lotion
Topiderm, Inc.

Clarity Rx® Physical Skin Defense
SPF 50

Drug Facts

<i>Active ingredient</i>	<i>Purpose</i>
Zinc Oxide 11%	Sunscreen
Titanium Dioxide 7%	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, Dimethicone, Methyl Trimethicone, Isononyl Isononanoate, Oleth-3 Phosphate, PEG-9 Polydimethylsiloxyethyl Dimethicone, Alumina, Butylene Glycol, Octyldodecyl Neopentanoate, VP/Eicosene Copolymer, Sodium Hyaluronate, Tocopheryl Acetate, Caffeine, Polyisobutene, Iron Oxides, Sucrose, Glycerin, Ethylhexyl Stearate, Isopropyl Titanium Triisostearate, Polysorbate 20, Polyacrylate-13, Panthenol, Cetearyl Alcohol, Dicetyl Phosphate, Ceteth-20 Phosphate, PEG-8 Methyl Ether Triethoxysilane, Hydrogen Dimethicone, Polyglyceryl-6 Polyricinoleate, Xanthan Gum, Sodium Hydroxide, Phenoxyethanol, Disodium EDTA.

PRINCIPAL DISPLAY PANEL - 99 g Bottle Label

CLARITY^{RX}®

5

PHYSICAL SKIN
DEFENSE

Mineral SPF 50 with
Antioxidants

Broad-Spectrum,
Perfecting Sunscreen
for Face and Body

Net 3.5 fl. oz. / 99 g

CLARITY^{RX}

PHYSICAL SKIN
DEFENSE

Mineral SPF 50 with
Antioxidants

Broad-Spectrum,
Perfecting Sunscreen
for Face and Body

Net 3.5 fl. oz. / 99 g

Drug Facts

Active ingredient Purpose

Zinc Oxide 11%.....Sunscreen

Titanium Dioxide 7%.....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

R0225 MADE IN U.S.A. 1203MB

Continued on back of peel panel

Drug Facts (continued)

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, Dimethicone, Methyl Trimethicone, Isononyl Isononanoate, Oleth-3 Phosphate, PEG-9 Polydimethylsiloxylethyl Dimethicone, Alumina, Butylene Glycol, Octyldodecyl Neopentanoate, VP/Eicosene Copolymer, Sodium Hyaluronate, Tocopheryl Acetate, Caffeine, Polyisobutene, Iron Oxides, Sucrose, Glycerin, Ethylhexyl Stearate, Isopropyl Titanium Triisostearate, Polysorbate 20, Polyacrylate-13, Panthenol, Cetearyl Alcohol, Dicetyl Phosphate, Ceteth-20 Phosphate, PEG-8 Methyl Ether Triethoxysilane, Hydrogen Dimethicone, Polyglyceryl-6 Polyricinoleate, Xanthan Gum, Sodium Hydroxide, Phenoxyethanol, Disodium EDTA.

CLARITY RX PHYSICAL SKIN DEFENSE SPF 50

zinc oxide and titanium dioxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51326-035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	110 mg in 1 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	75 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
METHYL TRIMETHICONE (UNII: S73ZQI0GXM)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)	
PEG-9 POLYDIMETHYLSILOXYETHYL DIMETHICONE (UNII: TYP81E471F)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CAFFEINE (UNII: 3G6A5W338E)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM ACRYLATE/SODIUM ACRYLOYLDIMETHYLTaurate COPOLYMER (4000000 MW) (UNII: 1DXE3F3OZX)	
DEXPANTHENOL (UNII: 1O6C93RI7Z)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
TRIETHOXYSILANE (UNII: 8T460WDH89)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51326-035-01	99 g in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M020	01/01/2022	

Labeler - Topiderm, Inc. (049121643)

Establishment

Name	Address	ID/FEI	Business Operations
Topiderm, Inc.		049121643	MANUFACTURE(51326-035)

Revised: 4/2025

Topiderm, Inc.