

MATTE SUNSCREEN SPF 50- titanium dioxide, zinc oxide lotion
CLINICAL SKIN LLC

Clinical Skin Matte Sunscreen SPF 50

Drug Facts

Active Ingredients

Titanium Dioxide 5.8%

Zinc Oxide 14.4%

Purpose

Sunscreen

Use

- 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

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 doctor

if rash occurs.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure
- reapply at least every 2 hours • use a water resistant sunscreen if swimming or

sweating

- 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 value of 15 or higher and other sun protection measures including:
- limit time in the sun, especially from 10 a.m. – 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months of age: Ask a doctor

Other Information

- store at Room Temperature (15°C-25°C (59°F-77°F))
- protect the product in this container from excessive heat and direct sun

Inactive ingredients

Aloe Barbadensis (Aloe Vera) Leaf Extract, Alumina, Aqua (Water), Ascorbic Acid, Bioflavonoids, Brassica Oleracea Italica (Broccoli) Extract, Caprylhydroxamic Acid, Caprylic/Capric Triglyceride, Caprylyl Glycol, Carbomer, Ceramide AP, Ceramide EOP, Ceramide NG, Ceramide NP, Cetearyl Alcohol, Cetearyl Glucoside, Cholesterol, Dicaprylyl Carbonate, Dimethicone, Dipotassium Glycyrrhizate, Disodium EDTA, Ethylhexylglycerin, Glycerin, Lauryl Glucoside, Methyl Methacrylate/Glycol Dimethacrylate Crosspolymer, Niacinamide, Phenoxyethanol, Phytosphingosine, Polyglyceryl-2 Dipolyhydroxystearate, Polyhydroxystearic Acid, Propanediol, Silica, Sodium Lauroyl Lactylate, Sodium Stearoyl Glutamate, Stearic Acid, Superoxide Dismutase, Tocopherol, Triethoxycaprylylsilane, Xanthan Gum

CLINICAL+ SKIN

MATTE SUNSCREEN

UV BROAD SPECTRUM

SPF 50

100% MINERAL

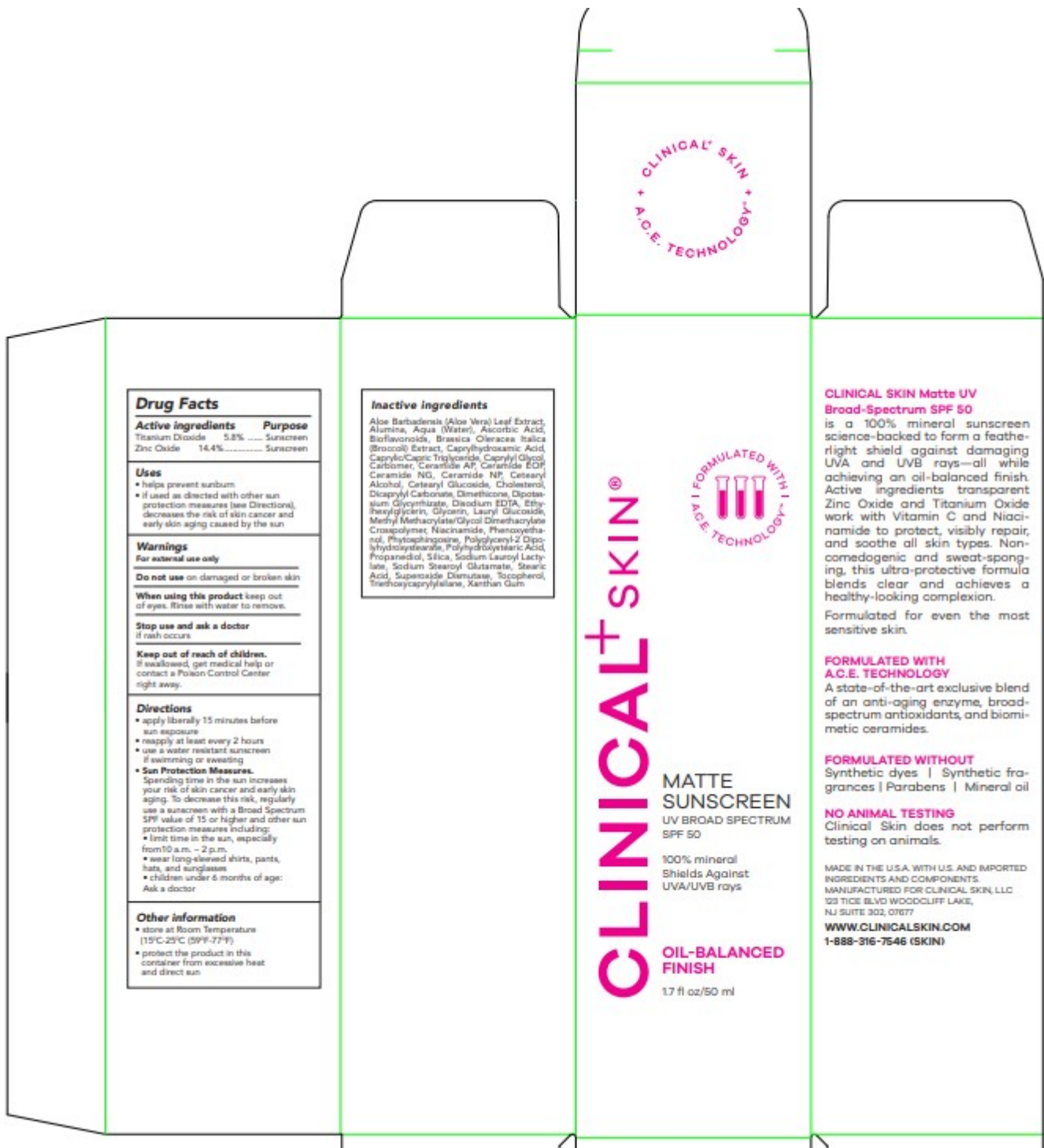
SHIELDS AGAINST

UVA/UVB RAYS

OIL-BALANCED

FINISH

1.7 FL OZ/50 ML



MATTE SUNSCREEN SPF 50

titanium dioxide, zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84248-025
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	14.4 mg in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	5.8 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BROCCOLI SEED OIL (UNII: SY01LVD4G4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
SODIUM LAUROYL LACTYLATE (UNII: 7243K85WFO)	
GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
METHYL METHACRYLATE/GLYCOL DIMETHACRYLATE CROSSPOLYMER (UNII: EG97988M5Q)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHYTOSPHINGOSINE (UNII: GIN46U9Q2Q)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CERAMIDE NG (UNII: C04977SRJ5)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CITRUS BIOFLAVONOIDS (UNII: BD70459I50)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CHOLESTEROL (UNII: 97C5T2UQ7J)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
MUSKMELON (UNII: ZV095H5633)	
NIACINAMIDE (UNII: 25X51I8RD4)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PROPANEDIOL (UNII: 5965N8W85T)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CERAMIDE AP (UNII: F1X8L2B00J)	
CERAMIDE NP (UNII: 4370DF050B)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC 64240-025			

1	NDC:84248-025-01	1 in 1 CARTON	06/17/2024	
1		50 mL in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020		06/17/2024	

Labeler - CLINICAL SKIN LLC (106954468)

Registrant - CLINICAL SKIN LLC (106954468)

Revised: 6/2024

CLINICAL SKIN LLC