DERMA SWISS SPF 50 MINERAL- titanium dioxide, zinc oxide lotion Prime Enterprises Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Derma Swiss SPF 50 Mineral Lotion

Active ingredients

Titanium Dioxide 4 %

Zinc Oxide 4 %

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

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

Directions

- apply liberally 15 minutes before sun exposure
- reapply:
- after 80 minutes of swimming or sweating
- immediately after towel drying
- 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 broad spectrum SPF 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-sleeve shirts, pants, hats, and sunglasses
- children under 6 months: Ask a doctor

Inactive ingredients:

Allantoin, Aloe Barbadensis Leaf Extract, Alumina, Bisabolol, Butyrospermum Parkii (Shea) Butter, Camellia Oleifera (Green Tea) Leaf Extract, Cetyl PEG/PPG-10/1 Dimethicone, Cyclopentasiloxane,

Dimethicone Crosspolymer, Disodium EDTA, Disteardimonium Hectorite, Ethylhexylglycerin, Glycerin, Helianthus Annuus (Sunflower) Seed Oil, Hexyl Laurate, Panthenol, PEG-10 Dimethicone, Phenoxyethanol, Polyglyceryl-4 Isostearate, Polymethylsilsesquioxane, Silica, Stearic Acid, Tocopheryl Acetate, Water

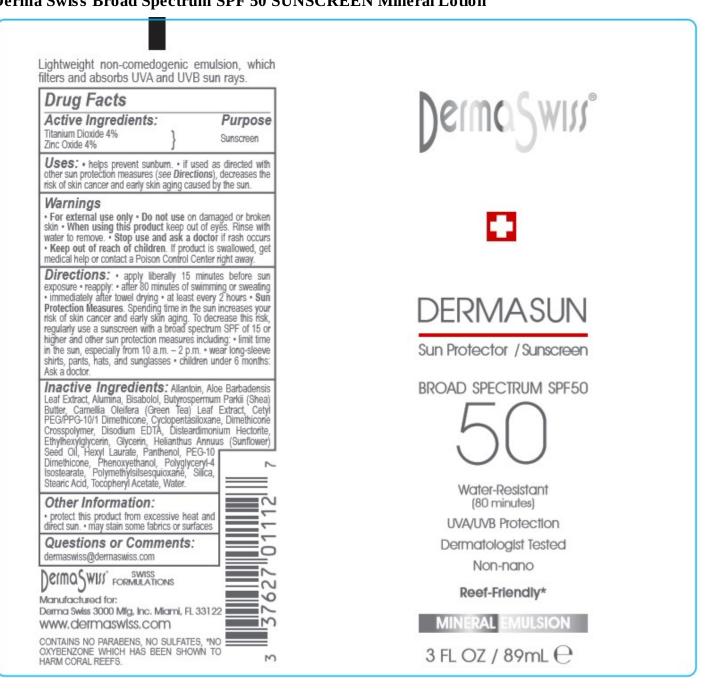
Other information

- protect this product from excessive heat and direct sun
- may stain some fabrics or surfaces

Questions or comments?

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Derma Swiss Broad Spectrum SPF 50 SUNSCREEN Mineral Lotion



DERMA SWISS SPF 50 MINERAL

titanium dioxide, zinc oxide lotion

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58443-0362

Route of Administration TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	42.4 mg in 1 mL			
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	42.4 mg in 1 mL			

Inactive Ingredients	
Ingredient Name	Strength
EDETATE DISO DIUM ANHYDRO US (UNII: 8 NLQ36 F6 MM)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)	
CAMELLIA OLEIFERA LEAF (UNII: 5077EL0C60)	
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)	
.ALPHATO CO PHERO L ACETATE (UNII: 9E8 X80 D2L0)	
ALLANTO IN (UNII: 344S277G0Z)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8 PR7V1SVM0)	
SHEA BUTTER (UNII: K49 155WL9 Y)	
WATER (UNII: 059QF0KO0R)	
CETYL PEG/PPG-10/1 DIMETHICO NE (HLB 2) (UNII: V2W71V8T0X)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	
DISTEARDIMO NIUM HECTO RITE (UNII: X687XDK09L)	
GLYCERIN (UNII: PDC6A3C0OX)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820 DPX33S7)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
LEVOMENOL (UNII: 24WE03BX2T)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
PANTHENOL (UNII: WV9CM0O67Z)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
ALUMINUM OXIDE (UNII: LMI26O6933)	

Product Characteristics				
Color	white (White to Off-White)	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:58443-0362-3	89 mL in 1 TUBE; Type 0: Not a Combination Product	11/20/2019			
Marketing Info	ormation				
Marketing Info		Marketing Start Date	Marketing End Date		
Marketing Info Marketing Category OTC monograph final		Marketing Start Date 11/20/2019	Marketing End Date		

Labeler - Prime Enterprises Inc. (101946028)

Registrant - Prime Enterprises Inc. (101946028)

Establishment			
Name	Address		Business Operations
Prime Enterprises Inc.		10 19 46 0 28	pack(58443-0362), manufacture(58443-0362), label(58443-0362), analysis(58443-0362)

Revised: 9/2020 Prime Enterprises Inc.