PROJECT REEF MINERAL SUNSCREEN SPF-50- zinc oxide, titanium dioxide lotion Project Reef LLC

PROJECT REEF Mineral Sunscreen SPF-50

Drug Facts

Active Ingredients

Zinc Oxide 14.00 % Titanium Dioxide 6.00 %

Purpose

Sunscreen

Uses:

• Helps prevent sunburn. • higher SPF gives more sunburn protection • If used as directed with other sun protection measures (see), decreases the risk of skin cancer and early skin aging caused by the sun

Directions

Warnings:

not intended for ingestion. 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 or irritation develops and lasts.

Keep out of reach of children.

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

If pregnant or breast-feeding, ask a health professional before use

Directions:

Apply liberally 15 minutes before sun exposure • Children under 6 months of age: ask a doctor.
Reapply at least every 2 hours • Reapply as needed or after towel drying, swimming, or sweating.
Use a water-resistant sunscreen if swimming or sweating • Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease the 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-sleeved shirts, pants, hats, and sunglasses

Sun Protection Measures.

Other Information:

• Protect the product in this container from excessive heat and direct sun.

Inactive Ingredients:

Algae Extract, *Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Benzyl Alcohol, *Butyrospermum Parkii (Shea) Butter, *Camellia Sinensis (Green Tea) Extract, Caprylic/Capric Triglyceride, Citric Acid, Citrus Aurantium Dulcis (Orange) Oil, *Cocos Nucifera (Coconut) Oil, Decyl Glucoside, Dehydroacetic Acid, Ethyl Vanillin, Fucus Vesiculosus (Seaweed) Extract, *Glycerin, Glyceryl Stearate, Helianthus Annuus, *Hippophae Rhamnoides (Sea Buckthorn) Oil, Hydroxyethylcellulose, Lavandula Angustifolia (Lavender) Oil, Polyhydroxystearic Acid, *Punica Granatum (Pomegranate) Extract, Rubus Idaeus (Raspberry) Extract, Stearic Acid, Tridecyl Salicylate, Xanthan Gum, Zemea (Corn) Propanediol. *Denotes Organic Ingredient

Questions?

Call 800 914 0146

Package Labeling:







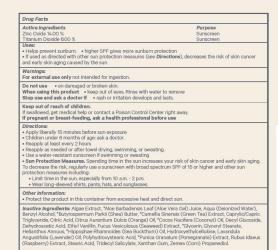












Questions? Call 800 914 0146 Distributed by: Project Reef LLC, 1215 S. Kihel Rd. Ste O #428, Kihel Hl 96753



MAUI, HAWAII

Plant Based & Sustainable Broad Spectrum SPF 50 Water Resistant for 80 Minutes

32FL OZ / 946.35ML

PROJECT REEF MINERAL SUNSCREEN SPF-50

zinc oxide, titanium dioxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81637-299
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	140 mg in 1 mL	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	60 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
WATER (UNII: 059QF0KO0R)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
SHEA BUTTER (UNII: K49155WL9Y)		
GREEN TEA LEAF (UNII: W2ZU1RY8B0)		
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
ORANGE OIL, COLD PRESSED (UNII: AKN3KSD11B)		
COCONUT OIL (UNII: Q9L0O73W7L)		

DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
ETHYL VANILLIN (UNII: YC9ST449YJ)	
FUCUS VESICULOSUS (UNII: 535G2ABX9M)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
PUNICA GRANATUM ROOT BARK (UNII: CLV24I3T1D)	
RASPBERRY (UNII: 4N14V5R27W)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIDECYL SALICYLATE (UNII: AZ Q08K38Z1)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CORN (UNII: 0N86727070)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81637- 299-00	946.35 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/21/2022	

Labeler - Project Reef LLC (096500461)

Revised: 11/2023 Project Reef LLC