

PROJECT REEF SPF-30 MINERAL SUNSCREEN- zinc oxide cream
Project Reef LLC

PROJECT REEF SPF-30 Mineral Sunscreen

Drug Facts

Active Ingredients

Zinc Oxide 14.00%

Purpose

Sunscreen

Uses:

- Helps prevent sunburn
- higher SPF gives more sunburn protection
- retains SPF after 80 minutes of swimming or sweating
- 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:

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 and as needed.
- Children under 6 months of age: ask a doctor

- Reapply:
- after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:: **Sun Protection Measures.**
- Limit time in the sun, especially from 10 a.m. - 2 p.m.
- Wear long- sleeved shirts, pants, hats, and sunglasses

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, Polyhydroxystearic Acid, *Punica Granatum (Pomegranate) Extract, Rubus Idaeus (Raspberry) Extract, Stearic Acid, Tridecyl Salicylate, Xanthan Gum, Zemea (Corn) Propanediol.

Questions?

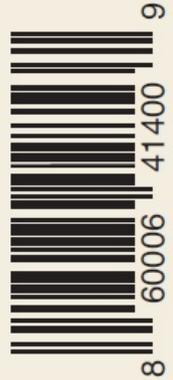
Call 800 914 0146

Package Labeling:

 <p>Project Reef MINERAL SUNSCREEN</p>	<p>Drug Facts</p>	<p style="writing-mode: vertical-rl; transform: rotate(180deg);">Distributed by: Project Reef LLC, 1215 S. Kihei Rd. Ste O #428, Kihei HI 96753</p>				
	<table border="1"> <thead> <tr> <th>Active Ingredients</th> <th>Purpose</th> </tr> </thead> <tbody> <tr> <td>Zinc Oxide 14.00 %</td> <td>Sunscreen</td> </tr> </tbody> </table>		Active Ingredients	Purpose	Zinc Oxide 14.00 %	Sunscreen
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	<p>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.</p>					
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Questions? Call 800 914 0146



This tube was made using recycled ocean bound plastics



MAUI, HAWAII

Plant-Based & Sustainable
Broad Spectrum SPF 30
Water Resistant for 80 min
4.2 oz / 125 ml

Each purchase removes 1lb of plastic
from the world's oceans.



PROJECT REEF SPF-30 MINERAL SUNSCREEN

zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81637-139
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

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 (UNII: AKN3KSD11B)	
COCONUT OIL (UNII: Q9L0O73W7L)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
ETHYL VANILLIN (UNII: YC9ST449YJ)	
NORI (UNII: 477TV3P5UX)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
HIPPOPHAE RHAMNOIDES SEED OIL (UNII: T53SBG6741)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
POMEGRANATE (UNII: 56687D1Z4D)	
RUBUS IDAEUS SEED (UNII: M3CL7US2ZG)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIDECYL SALICYLATE (UNII: AZQ08K38Z1)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CORN (UNII: 0N8672707O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81637-139-00	125 mL in 1 TUBE; Type 0: Not a Combination Product	03/08/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/08/2021	

