NATIVE MINERAL SUNCARE DAILY FACE BROAD SPECTRUM SPF 30 UNSCENTED- zinc oxide lotion The Procter & Gamble Manufacturing Company

Native Mineral Suncare Daily Face Broad Spectrum SPF 30 Unscented Drug Facts

ZINC OXIDE 20%

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 at least every 2 hours
- use 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:
 - o limit time in sun, especially from 10 a.m. 2 p.m.
 - wear long-sleeved shirts, pants, hats, and sunglasses
- · children under 6 months: ask a doctor

Other information

• protect this product from excessive heat and direct sun

Inactive ingredients

water, caprylic/capric triglyceride, coconut alkanes, glycerin, cetearyl alcohol, cocoglucoside, arachidyl alcohol, behenyl alcohol, arachidyl glucoside, polyhydroxystearic acid, benzyl alcohol, xanthan gum, coco-caprylate/caprate, tocopherol, persea gratissima (avocado) oil, helianthus annuus (sunflower) seed oil

Distr. by Native San Francisco, CA 94111

Questions or comments?

Please contact support#\@nativecos.com

PRINCIPAL DISPLAY PANEL - 50 mL Tube

BROAD SPECTRUM SPF 30

NATIVE

MINERAL SUNCARE

ZINC OXIDE SUNSCREEN

DAILY FACE

UNSCENTED

NO OXYBENZONE

NO OCTINOXATE

1.7 FL OZ (50 mL)

Drug Facts

Active ingredient Zinc oxide 20%.

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DISTR. BY NATIVE SAN FRANCISCO, CA 94111 CRUELTY FREE

21261570





100% MINERAL UV ACTIVE SUN PROTECTION



ABSORBS QUICKLY & NO RESIDUE



LIGHTWEIGHT & NON-GREASY FORMULA



BROAD SPECTRUM SPF 30



MINERAL SUNCARE ZINC OXIDE SUNSCREEN



DAILY FACE

UNSCENTED

NO OXYBENZONE NO OCTINOXATE

1.7 FL OZ (50 mL)

MADE WITH ♥ CLEAN . SIMPLE . EFFECTIVE

WE'VE GOT YOU COVERED

Say hello to 100% Mineral UV Active sun protection. Thoughtfully made with naturally derived zinc oxide and without oxybenzone or octinoxate. All sunshine, no worries.

DIRECTIONS

Apply liberally 15 minutes before sun exposure. Reapply at least every 2 hours.

NATIVE MINERAL SUNCARE DAILY FACE BROAD SPECTRUM SPF 30 UNSCENTED

zinc oxide lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:84126-157

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) 20 g in 100 mL ZINC OXIDE

Inactive Ingredients			
Ingredient Name	Strength		
COCONUT ALKANES (UNII: 1E5KJY107T)			
COCO GLUCOSIDE (UNII: ICS790225B)			
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)			
PERSEA GRATISSIMA (AVOCADO) OIL (UNII: 6VNO72PFC1)			
BEHENYL ALCOHOL (UNII: 9G10E216XY)			
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			
TOCOPHEROL (UNII: R0ZB2556P8)			
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)			
CETEARYL ALCOHOL (UNII: 2DMT128M1S)			
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)			
XANTHAN GUM (UNII: TTV12P4NEE)			
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)			
ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)			
HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL (UNII: 3W1JG795YI)			
BENZYL ALCOHOL (UNII: LKG8494WBH)			

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:84126-157- 50	1 in 1 CARTON	01/01/2025				
1		50 mL in 1 TUBE; 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	01/01/2025				

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 7/2025 The Procter & Gamble Manufacturing Company