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

Native Mineral Sunscreen 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:
 - 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

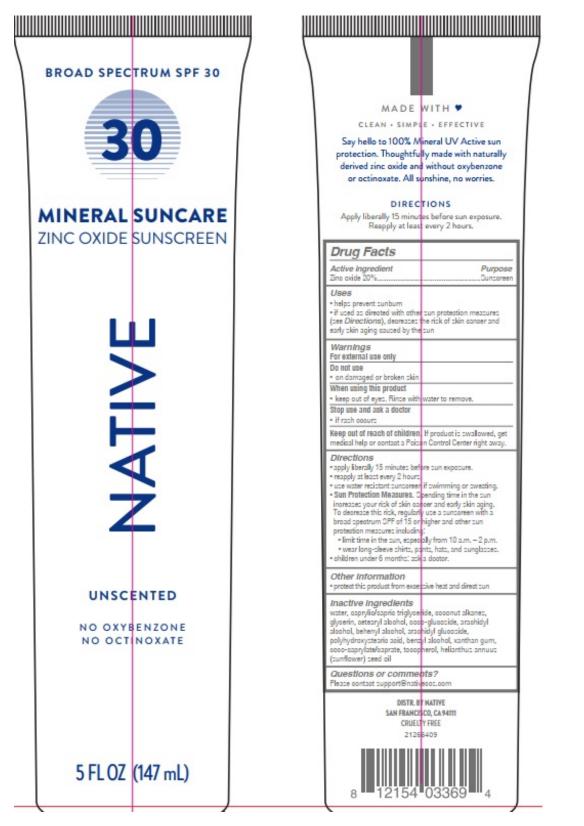
water, caprylic/capric triglyceride, coconut alkanes, glycerin, cetearyl alcohol, cocoglucoside, arachidyl alcohol, behenyl alcohol, arachidyl glucoside, polyhydroxystearic acid, benzyl alcohol, xanthan gum, coco-caprylate/caprate, tocopherol, helianthus annuus (sunflower) seed oil

Distr. by Native San Francisco, CA 94111

Questions or comments?

Please contact support@nativecos.com

PRINCIPAL DISPLAY PANEL - 147 mL Tube



- UNSCENTED
- NO OXYBENZONE
- NO OCTINOXATE
- NATIVE
- .
- MINERAL SUNSCREEN
- SUNSCREEN WITH

ZINC OXIDE BROAD SPECTRUM SPF 30 5 FL OZ (147 mL)

NATIVE MINERAL SUNSCREEN BROAD SPECTRUM SPF 30 UNSCENTED zinc oxide lotion **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:69423-550 **Route of Administration** TOPICAL **Active Ingredient/Active Moiety Basis of Strength Ingredient Name** Strength **ZINC OXIDE** (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) ZINC OXIDE 20 g in 100 mL **Inactive Ingredients Ingredient Name** Strength COCONUT ALKANES (UNII: 1E5KJY107T) COCO GLUCOSIDE (UNII: ICS790225B) COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H) WATER (UNII: 059QF0K00R) GLYCERIN (UNII: PDC6A3C0OX) TOCOPHEROL (UNII: R0ZB2556P8) DOCOSANOL (UNII: 9G10E216XY) CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S) POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F) MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU) ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)

SUNFLOWER OIL (UNII: 3W1JG795YI) BENZYL ALCOHOL (UNII: LKG8494WBH) XANTHAN GUM (UNII: TTV12P4NEE)

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-550- 14	147 mL in 1 TUBE; Type 0: Not a Combination Product	04/01/2021	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M020	04/01/2021				

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 9/2024

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