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

Native Mineral Sunscreen Broad Spectrum SPF 30 Tropical Breeze

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, coco-glucoside, arachidyl alcohol, behenyl alcohol, arachidyl glucoside, polyhydroxystearic acid, benzyl alcohol, xanthan gum, coco-caprylate/caprate, fragrance, 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

BROAD SPECTRUM SPF 30

MINERAL SUNCARE

ZINC OXIDE SUNSCREEN

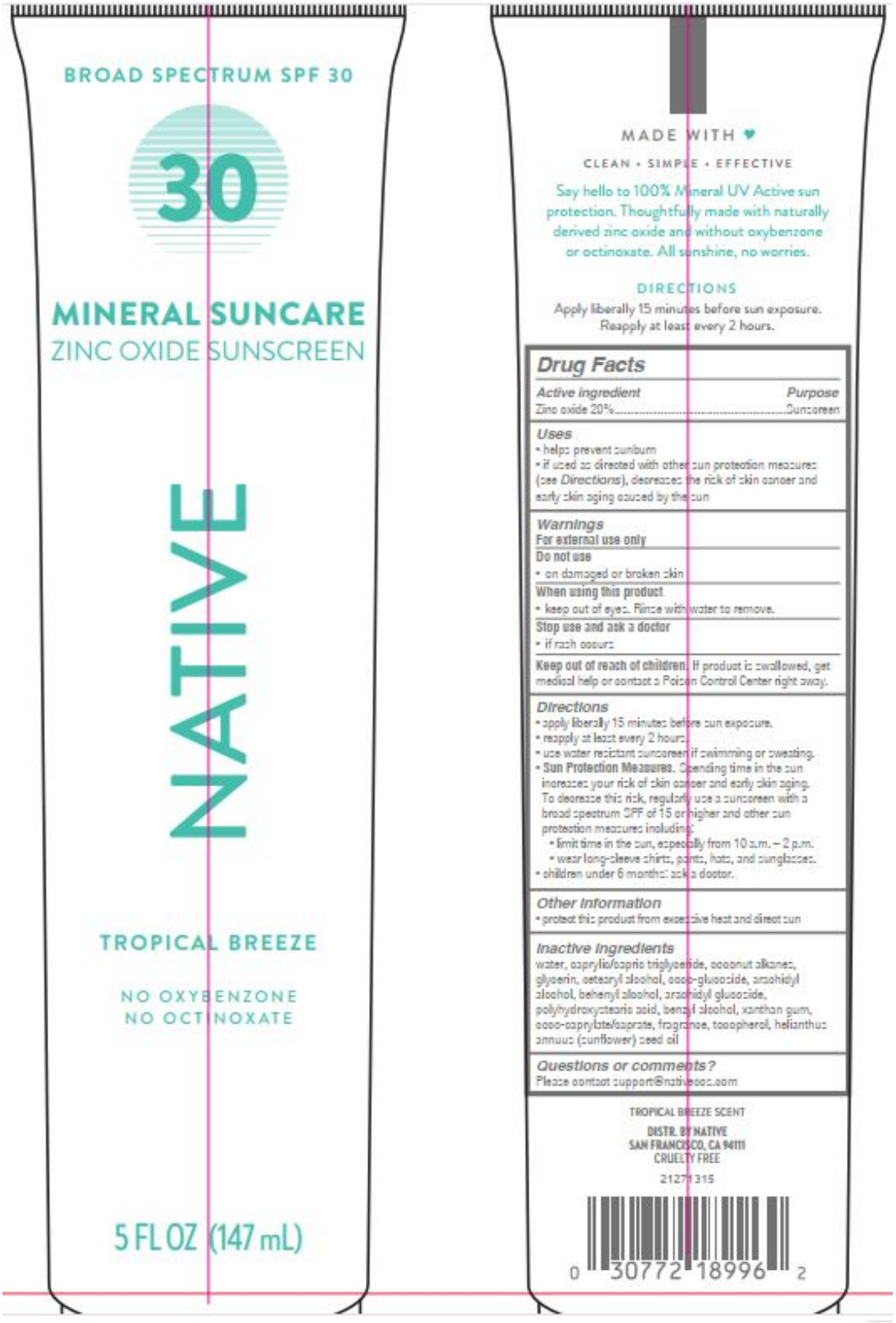
NATIVE

TROPICAL BREEZE

NO OXYBENZONE

NO OCTINOXATE

5 FL OZ (147 mL)



NATIVE MINERAL SUNSCREEN BROAD SPECTRUM SPF 30 TROPICAL BREEZE

zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84126-605

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	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: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
TOCOPHEROL (UNII: R0ZB2556P8)	
DOCOSANOL (UNII: 9G1OE216XY)	
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

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84126-605-14	147 mL in 1 TUBE; Type 0: Not a Combination Product	10/28/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/28/2024	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2025 The Procter & Gamble Manufacturing Company