SPF 50 SUNSCREEN STICK- zinc oxide stick
Australian Gold

Drug Facts

Active ingredient

Zinc Oxide....20.0%

Purpose

Sunscreen

Keep Out of Reach of Children

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

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.

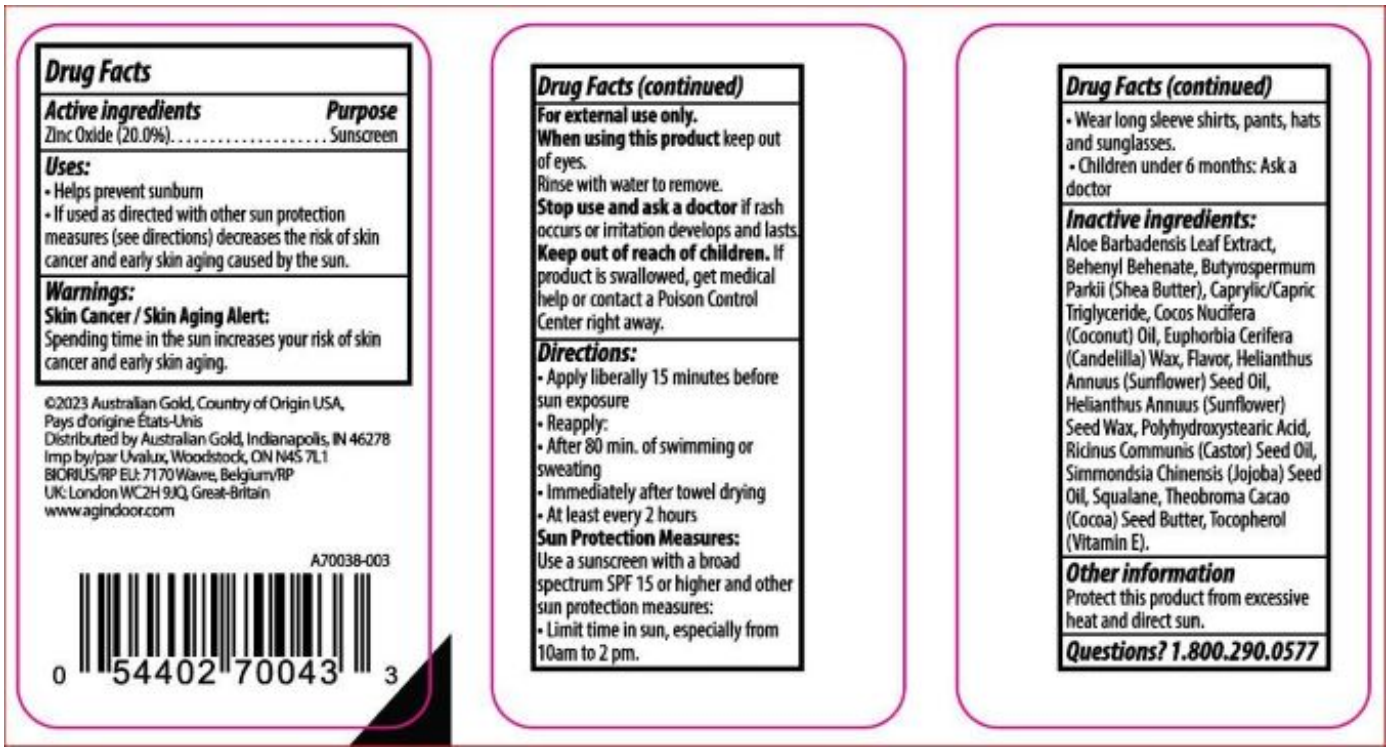
Directions

Apply liberally 15 minutes before sun exposure. Reapply after 80 minutes of swimming or sweating. Immediately after towel drying. At least every 2 hours. 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 Broad Spectrum SPF value 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-sleeve shirts, pants, hats and sunglasses. Children under 6 months: Ask a doctor.

Inactive Ingredients

Aloe Extract, Behenyl Behenate, Shea Butter, Caprylic/Capric Triglyceride, Coconut Oil, Candelilla Wax, Sunflower Oil, Sunflower Wax, Polyhydroxystearic Acid, Castor Oil, Jojoba Oil, Squalane, Cocoa Butter, Tocopherol (Vitamin E).

Package/Label Principal Display Panel



Drug Facts

Active ingredients — **Purpose**
Zinc Oxide (20.0%) ..... 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:
Skin Cancer / Skin Aging Alert:
Spending time in the sun increases your risk of skin cancer and early skin aging.

Distributed by Australian Gold, Indianapolis, IN 46278
Imp by/par Uvalux, Woodstock, ON N4S 7L1
BIORIUS/RP EU: 7170 Wavre, Belgium/RP
UK: London WC2H 9JQ, Great-Britain
www.agindoor.com

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0 54402 70043 3

Drug Facts (continued)

For external use only.
When using this product keep out of eyes.
Rinse with water to remove.
Stop use and ask a doctor if rash occurs or irritation develops and lasts.
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:
- After 80 min. of swimming or sweating
- Immediately after towel drying
- At least every 2 hours

Sun Protection Measures:
Use a sunscreen with a broad spectrum SPF 15 or higher and other sun protection measures:
- Limit time in sun, especially from 10am to 2 pm.

Drug Facts (continued)

- Wear long sleeve shirts, pants, hats and sunglasses.
- Children under 6 months: Ask a doctor

Inactive ingredients:
Aloe Barbadensis Leaf Extract, Behenyl Behenate, Butyrospermum Parkii (Shea Butter), Caprylic/Capric Triglyceride, Cocos Nucifera (Coconut) Oil, Euphorbia Cerifera (Candelilla) Wax, Flavor, Helianthus Annuus (Sunflower) Seed Oil, Helianthus Annuus (Sunflower) Seed Wax, Polyhydroxystearic Acid, Ricinus Communis (Castor) Seed Oil, Simmondsia Chinensis (Jojoba) Seed Oil, Squalane, Theobroma Cacao (Cocoa) Seed Butter, Tocopherol (Vitamin E).

Other information
Protect this product from excessive heat and direct sun.

Questions? 1.800.290.0577

SPF 50 SUNSCREEN STICK

zinc oxide stick

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84061-129
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	200 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
COCONUT OIL (UNII: Q9L0O73W7L)	395 mg in 1 g
WHITE WAX (UNII: 7G1J5DA97F)	162 mg in 1 g
COCOA BUTTER (UNII: 512OYT1CRR)	57 mg in 1 g
SUNFLOWER OIL (UNII: 3W1JG795YI)	64 mg in 1 g

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84061-129-01	14 g in 1 CONTAINER; Type 0: Not a Combination Product	03/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/01/2024	

Labeler - Australian Gold (073212842)

Registrant - OraLabs (801824756)

Establishment

Name	Address	ID/FEI	Business Operations
OraLabs		801824756	manufacture(84061-129) , label(84061-129) , analysis(84061-129)

Revised: 1/2026 Australian Gold