

**COLLAGEN GLOW MINERAL SUNSCREEN SPF 50- zinc oxide lotion**  
**NAKED SUNDAYS PTY LTD**

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**NAKED SUNDAYS COLLAGEN GLOW MINERAL SUNSCREEN**

**ACTIVE INGREDIENT**

ZINC OXIDE 22.75%

**Purpose**

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 ageing 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.
- CONTAINS SODIUM BENZOATE.

Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

**DIRECTIONS**

- APPLY GENEROUSLY 15 MINUTES BEFORE SUN EXPOSURE.
- Reapply after 80 minutes of swimming or sweating.
- Reapply immediately after towel drying.
- Reapply at least every 2 hours.
- Children under 3 years of age: Ask a doctor.

**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:

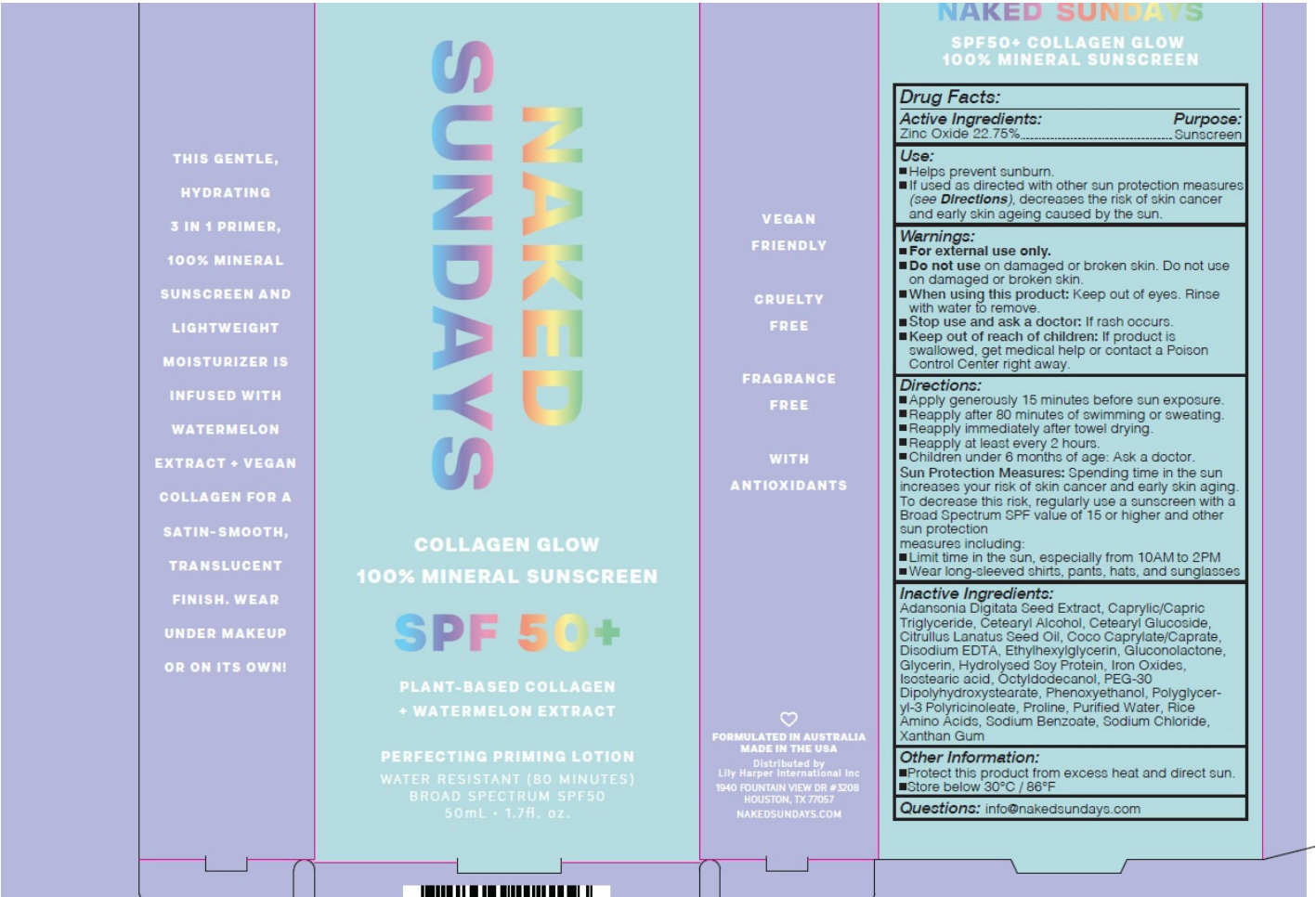
- Limit time in the sun, especially from 10.00AM - 2.00PM
- Wear long-sleeved shirts, pants, hats, and sunglasses

**INACTIVE INGREDIENTS**

CETEARYL GLUCOSIDE, CETOSTEARYL ALCOHOL, COCO-CAPRYLATE, DISODIUM EDETATE, GLUCONOLACTONE, HYDROLYZED ADANSONIA DIGITATA SEED EXTRACT, HYDROLYZED RICE PROTEIN, HYDROLYZED SOY PROTEIN, ISOSTEARIC ACID, CAPRIC TRIGLYCERIDES, OCTYLDODECANOL, PEG-30 DIPOLYHYDROXYSTEARATE, POLYGLYCERYL-3 POLYRICINOLEATE, PROLINE, WATER (AQUA), SODIUM BENZOATE, SODIUM CHLORIDE.

OTHER INFORMATION

Protect the product in this container from excessive heat and direct sun.



COLLAGEN GLOW MINERAL SUNSCREEN SPF 50			
zinc oxide lotion			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81104-100
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)		ZINC CATION	22.75 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>CETEARYL GLUCOSIDE</b> (UNII: 09FUA47KNA)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>COCO-CAPRYLATE</b> (UNII: 4828G836N6)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>GLUCONOLACTONE</b> (UNII: WQ29KQ9POT)	
<b>ADANSONIA DIGITATA SEED</b> (UNII: 2936P60TPX)	
<b>ORYZA SATIVA WHOLE</b> (UNII: 84IVV0906Z)	
<b>HYDROLYZED SOY PROTEIN (ENZYMATIC; 2000 MW)</b> (UNII: 1394NXB9L6)	
<b>ISOSTEARIC ACID</b> (UNII: X33R8U0062)	
<b>TRICAPRIN</b> (UNII: O1PB8EU98M)	
<b>OCTYLDODECANOL</b> (UNII: 461N1O614Y)	
<b>PEG-30 DIPOLYHYDROXYSTEARATE</b> (UNII: 9713Q0S7FO)	
<b>POLYGLYCERYL-3 RICINOLEATE</b> (UNII: MZQ63P0N0W)	
<b>PROLINE</b> (UNII: 9DLQ4CIU6V)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81104-100-01	50 mL in 1 TUBE; Type 0: Not a Combination Product	12/01/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2020	

**Labeler** - NAKED SUNDAYS PTY LTD (746918845)

Revised: 6/2025

NAKED SUNDAYS PTY LTD