# NAKED SUNDAYS CABANA GLOW MINERAL GLOW SERUM DROPS BLUSH 15ML SPF 50- zinc oxide liquid Wild Child Laboratories PTY LTD

-----

## NAKED SUNDAYS CABANA GLOW MINERAL GLOW SERUM DROPS Blush 15mL

## **ACTIVE INGREDIENTS**

**Active Ingredients:** 

Zinc Oxide 14.9%

# **Purpose**

Sunscreen

#### **USES**

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

Warnings: For external use only. Do not use on damaged or broken skin Avoid prolonged sun exposure. If it gets in your eyes, wash well with water. Keep out of reach of children. Stop use and ask a doctor if rashes occur. If product is swallowed, get medical help or contact a Poison Control Center right away.

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

#### **DIRECTIONS**

- SPRAY 7 TIMES WITH MOUTH AND LIPS CLOSED 8-10" AWAY FROM FACE 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**

Inactive Ingredients: Aluminium Hydroxide, Butyloctyl Salicylate, C12-15 Alkyl Benzoate, Coco-Caprylate/Caprate, Dimethicone, Ethylhexyl Methoxycrylene, Ethylhexylglycerin, Hibiscus sabdariffa Flower Extract, Hydroxyacetophenone, Iron Oxides/(CI 77499, CI 77491, CI 77492), Isododecane, Isostearic Acid, Lauryl PEG-8 Dimethicone, Magnesium Sulfate, Mica/CI 77019, Phenoxyethanol, Polyglyceryl-3 Polyricinoleate, Propanediol, Purified Water, Silica, Sodium Hyaluronate, Stearic Acid, Tin Oxide, Titanium dioxide/CI 77891

# OTHER INFORMATION

Other Information: Protect this product from excess heat and direct sun. Store below 30°C / 86°F.

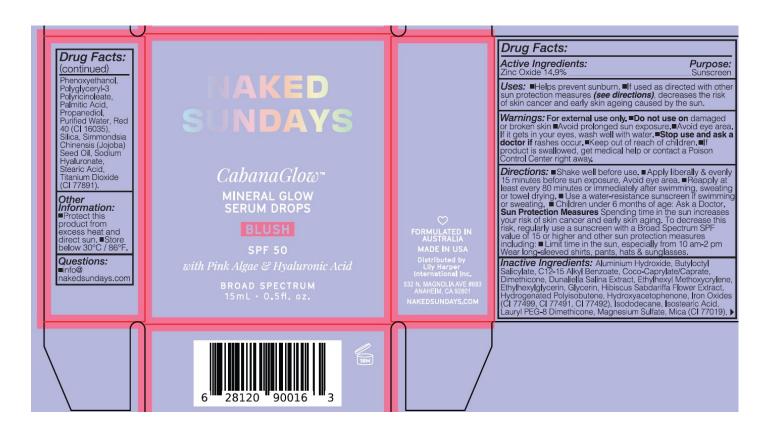
Directions: Shake well before use.

Apply liberally and reapply every 80 minutes, especially after swimming or towel drying.

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 lime in the sun, especially from 10 am-2 pm Wear longsleeved shirts, pants, hats, and sunglasses For Children under 6 months of age ask a doctor

Sunscreen

#### Label



# NAKED SUNDAYS CABANA GLOW MINERAL GLOW SERUM DROPS BLUSH 15ML SPF 50

zinc oxide liquid

_			
Drad	IICT.	Intorr	mation
FIUU	uct		Hativii

Product Type HUMAN OTC DRUG Item Code (Source) NDC:30807-2570

Route of Administration TOPICAL

# **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)

ZINC OXIDE

14.9 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
STANNIC OXIDE (UNII: KM7N50LOS6)		
DIMETHICONE (UNII: 92RU3N3Y1O)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)		
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)		
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
ISOSTEARIC ACID (UNII: X33R8U0062)		
WATER (UNII: 059QF0KO0R)		
PROPANEDIOL (UNII: 5965N8W85T)		
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)		

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
HIBISCUS SABDARIFFA FLOWER (UNII: 45TGG6IU6M)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
LAURYL PEG-8 DIMETHICONE (300 CPS) (UNII: ELL2U7K8T8)	
MICA (UNII: V8A1AW0880)	
MAGNESIUM SULFATE ANHYDROUS (UNII: ML30MJ2U7I)	
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q00K5D0T4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
ISODODECANE (UNII: A8289P68Y2)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:30807- 2570-1	15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	12/20/2023	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M020	06/30/2023		

# Labeler - Wild Child Laboratories PTY LTD (890661643)

Revised: 10/2024 Wild Child Laboratories PTY LTD