SILK MINERAL SPF 50 SUNSCREEN FACE- zinc oxide lotion Baxter Laboratories

Standard Procedure Silk Mineral SPF 50 Sunscreen Face Lotion

Active Ingredient Purpose

Zinc Oxide20%....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.

Keep out of reach of children. If product is swallowed, get medical help or contact a poison control center right awa.

Stop use and ask a doctor if rash occurs.

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.

Directions

Apply liberally 15 minutes before sun exposure

Reapply at least every 2 hours

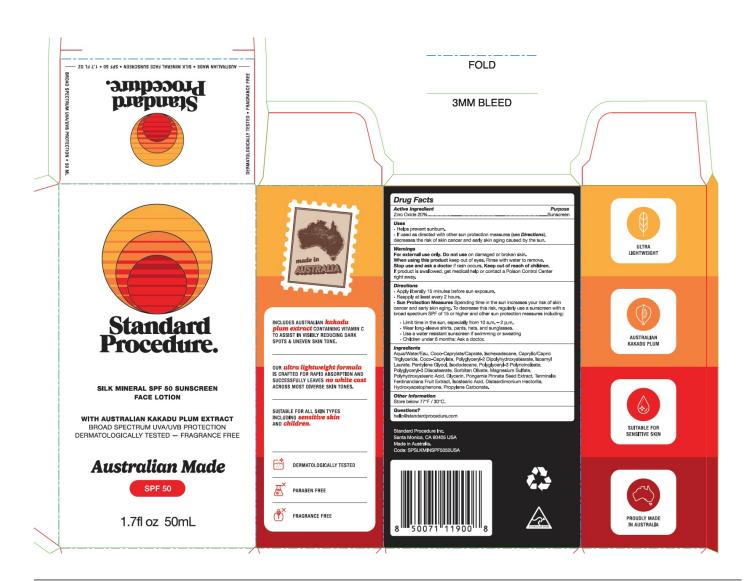
Ingredients

Aqua/Water/Eau, Coco-Caprylate/Caprate, Isohexadecane, Caprylic/Capric Triglyceride, Coco-Caprylate, Polyglyceryl-2 Dipolyhydroxystearate, Isoamyl Laurate, Pentylene Glycol, Isododecane, Polyglyceryl-3 Polyricinoleate, Polyglyceryl-3 Diisostearate, Sorbitan Olivate, Magnesium Sulfate, Polyhydroxystearic Acid, Glycerin, Pongamia Pinnata Seed Extract, Terminalia Ferdinandiana Fruit Extract, Isostearic Acid, Disteardimonium Hectorite, Hydroxyacetophenone, Propylene Carbonate

Standard Procedure

Silk Mineral SPF 50 Sunscreen Face Lotion

1.7 fl oz 50 mL



SILK MINERAL SPF 50 SUNSCREEN FACE

zinc oxide lotion

Product Information	oduct Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70157-026
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	
PENTYLENE GLYCOL (UNII: 50C1307PZG)		
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)		
GLYCERIN (UNII: PDC6A3C0OX)		
KAKADU PLUM (UNII: 0ZQ1D2FDLI)		
WATER (UNII: 059QF0KO0R)		

COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q00K5D0T4)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
ISODODECANE (UNII: A8289P68Y2)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
PONGAMIA PINNATA SEED (UNII: C2BRV53B1V)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
MAGNESIUM SULFATE (UNII: DE08037SAB)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:70157-026- 01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2025	

Marketing In	eting Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	05/09/2025	

Labeler - Baxter Laboratories (740537709)

Revised: 4/2025 Baxter Laboratories