# STANDARD PROCEDURE SPF 50 SUNSCREEN FACE AND BODY- avobenzone, homosalate, octisalate, octocrylene lotion Baxter Laboratories

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# Standard Procedure SPF 50+ Sunscreen Face and Body Lotion

Active Ingredients Purpose

Avobenzone3%, Homosalate 10%, Octisalate 5%, Octocrylene 8% Sunscreen

#### **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 aging caused by the sun
- Keep out of reach of children
- If swallowed, get medical help or contact a Poison Control Center right away

## 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 generously and evenly
- Reapply
- after 80 minutes of swimming
- Immediately after towel drying
- at least every 2 hours
- use a 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 broad spectrum SPF 15 or higher and other sun protection measures including:
- limit time in the sun, especially from 10 a.m. to 2 p.m.
- wear long-sleeve shirts, pants, hats, and sunglasses
- Children under 6 months: ask a doctor

# Inactive Ingredients

Water, Beeswax, Isopropyl Palmitate, Cetearyl Alcohol, Cyclopentasiloxane, Cyclohexasiloxane, Ceteareth-20, Hydroxyacetophenone, Carbomer, Benzyl alcohol, Phenoxyethanol, Sodium Stearoyl Glutamate, Triethanolamine, Tocopheryl Acetate, Glycerin, Maltodextrin, Aloe Barbadensis Leaf Juice, Mangifera Indica (Mango) Fruit Extract, Carica Papaya (Papaya) Fruit Extract, Ananas Sativus (Pineapple) Fruit Extract, Citric Acid.

#### Other information

• protect the product in this container from excessive heat and direct sun

## **Questions?**

1 213-681-1801 or usa@standardprocedure.co

Standard Procedure

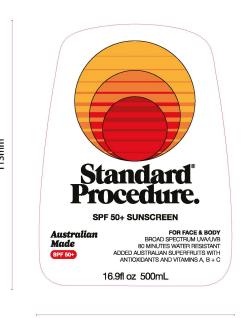
SPF 50+ sunscreen

For face and body

Broad Spectrum UVA/UVB

80 Minutes Water Resistant

Added Australian Superfruits with Antioxidants and Vitamins A, B + C





83mm

# STANDARD PROCEDURE SPF 50 SUNSCREEN FACE AND BODY

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70157-027
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 mL	
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	8 g in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
PAPAYA (UNII: KU94FIY6JB)	
BEESWAX (UNII: 2ZA36H0S2V)	
CETEARETH-20 (UNII: YRC528SWUY)	
TRIETHANOLAMINE (UNII: 903K93S3TK)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
CARBOMER (UNII: 0A5MM307FC)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CITRIC ACID (UNII: 2968PHW8QP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
MANGO (UNII: 162913NR86)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
PINEAPPLE (UNII: 2A88ZO081O)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70157- 027-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2025	
2	NDC:70157- 027-02	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2025	
3	NDC:70157- 027-03	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2025	
4	NDC:70157- 027-04	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/30/2025	

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

OTC Monograph Drug	M020	05/30/2025	

# **Labeler -** Baxter Laboratories (740537709)

Revised: 5/2025 Baxter Laboratories