AUSTRALIAN GOLD BOTANICAL SUNSCREEN BROAD SPECTRUM SPF 30 NATURAL- avobenzone, homosalate, octisalate, octocrylene spray
Prime Packaging Inc.

Botanical Sunscreen Broad Spectrum SPF 30 Natural

Active Ingredients

Avobenzone 3%

Homosalate 10%

Octisalate 5%

Octocrylene 2.75%

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 aging caused by the sun.

Warnings

For external use only

Do not use on damaged or broken skin

When using this product keep away from face to avoid breathing it

- keep out of eyes. Rinse with water to remove

Stop use and ask a doctor if rash occurs

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

Flammable: Avoid fire, flame, heat and smoking. **Contents under presssure.** Do not puncture or incinerate. Store at temperatures below 120ºC (50ºC).

Directions

- shake well before use
- Apply liberally 15 minutes before sun exposure and rub into skin
- hold container 4 to 6 inches from the skin to apply
- do not spray directly into face. Spray on hands then apply to face
- do not apply in windy conditions
- use in well-ventilated area

- 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 a broad spectrum SPF 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

SD Alcohol 40-B, Glycerin, VA/Butyl Maleate/Isobornyl Acrylate Copolymer, Fragrance (Parfum), Water/Aqua/Eau, Polyester-8, Tocopheryl Acetate, Eucalyptus Globulus Leaf Extract, Poryphyra Umbilicalis Extract, Terminalia Ferdinandiana (Kakadu Plum) Fruit Extract

Other Information

- protect this product from excessive heat and direct sun
- May stain some fabrics or surfaces

Question or comments?

Call toll free 1-855-LIV-GOLD (548-4653)

Botanical Sunscreen Broad Spectrum SPF 30 Natural

Australian Gold®

Drug Facts

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*Organic
**Vitamin E

crueltyfree andvegan

No Added Parabens, Dye Free, Oil Free, Sulfate Free, Petrolatum Free & No CFCs

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www.AustralianGold.com

A70800
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0 54402 51050 6

DOT 2P M5655

ORGANIC ALCOHOL

HYPO-ALLERGENIC

OXYBENZONE FREE

PABA FREE

PHTHALATE FREE

GLUTEN FREE

DERMATOLOGIST TESTED

PEDIATRICIAN TESTED

NO ANIMAL TESTING

Australian Gold®

BOTANICAL 30

NATURAL SPRAY SUNSCREEN
POWDER DRY, BREATHABLE FEEL

WATER RESISTANT (80 MINUTES)
BROAD SPECTRUM SPF 30

6 FL OZ (177 mL)
Net Wt. 5.3 OZ (152 g)

AUSTRALIAN GOLD BOTANICAL SUNSCREEN BROAD SPECTRUM SPF 30 NATURAL

avobenzone, homosalate, octisalate, octocrylene spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13630-0110
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	26.04 mg in 1 mL

HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	86.8 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	43.4 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	23.87 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYL ACRYLATE/C16-C20 ALKYL METHACRYLATE/METHACRYLIC ACID/METHYL METHACRYLATE COPOLYMER (UNII: 7K68DGG29P)	
ALCOHOL (UNII: 3K9958V90M)	
POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENOYL CAPPED) (UNII: T9296U138P)	
PORPHYRA UMBILICALIS (UNII: 14AN0J70WO)	
KAKADU PLUM (UNII: 0ZQ1D2FDLI)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
GLYCERIN (UNII: PDC6A3C0OX)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13630-0110-4	177 mL in 1 CAN; Type 0: Not a Combination Product	07/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	07/15/2016	

Labeler - Prime Packaging Inc. (805987059)

Registrant - Prime Packaging Inc. (805987059)

Establishment

Name	Address	ID/FEI	Business Operations
Prime Enterprises Inc		101946028	manufacture(13630-0110) , analysis(13630-0110)

Establishment

Name	Address	ID/FEI	Business Operations
Prime Packaging Inc.		805987059	pack(13630-0110) , label(13630-0110)

Revised: 5/2024

Prime Packaging Inc.