## SPF 50 SUNSCREEN- avobenzone, homosalate, octisalate, octocrylene lotion Consumer Product Partners, LLC

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Wild Tropics 059.000/059AA SPF 50 Sunscreen Lotion

### **Active ingredients**

Avobenzone 3%

Homosalate 10%

Octisalate 4.5%

Octocrylene 8%

### 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 out of eyes. Rinse with water to remove.

### Stop use and ask a doctor if

rash occurs

## Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

### Directions

apply liberally 15 minutes before sun exposure

- apply to all skin exposed to the sun
- 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 Broad Spectrum SPF value 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-sleeved shirts, pants, hats and sunglasses
- children under 6 months of age: Ask a doctor

# Other information

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

## Inactive ingredients

water, glycerin, aluminum starch octenylsuccinate, styrene/acrylates copolymer, polyester-7, silica, chlorphenesin, arachidyl alcohol, beeswax, neopentyl glycol diheptanoate, acrylates/C10-30 alkyl acrylate crosspolymer, behenyl alcohol, tocopherol, arachidyl glucoside, glyceryl stearate, PEG-100 stearate, potassium hydroxide, benzyl alcohol, disodium EDTA, fragrance

## Disclaimer

May stain or damage some fabrics or surfaces

## Adverse reaction

Manufactured by Vi-Jon, LLC One Swan Drive

Smyrna, TN 37167

Pat. Pend.

# Principal display panel

Wild Tropics™ SUNSCREEN LOTION BROAD SPECTRUM SPF 50 Octinoxate & Oxybenzone Free Hypoallergenic Dermatologist tested

## Water resistant (80 minutes)

## Fresh Banana Scent

**SPF 50** 

## 6 FL OZ (177 mL)



SPF 50 SUNSCREEN avobenzone, homosalate, octisalate, octocrylene lotion						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC	2:11344-059	
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ingred	lient Name		<b>Basis of Stren</b>	gth	Strength	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)			AVOBENZ ONE		30 mg in 1 mL	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)		SV4M95S)	HOMOSALATE		100 mg in 1 mL	

OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	45 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	80 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)	
STYRENE/ACRYLAMIDE COPOLYMER (500000 MW) (UNII: 5Z4DPO246A)	
POLYESTER-7 (UNII: 0841698D2F)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
DOCOSANOL (UNII: 9G10E216XY)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
PEG-100 STEARATE (UNII: YD01N1999R)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

# Packaging

#	Item Code Package Description		Marketing Start Date	Marketing End Date	
1	NDC:11344-059- 30	177 mL in 1 TUBE; Type 0: Not a Combination Product	03/01/2024		
Marketing Information					
	Marketing	Application Number or Monograph	Marketing Start	Marketing End	

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/01/2024	

Labeler - Consumer Product Partners, LLC (119091520)

**Registrant -** Consumer Product Partners, LLC (119091520)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
Consumer Product Partners, LLC		119091514	manufacture(11344-059)		