

**SOLEIL SET SUNSCREEN SPF 50- avabenzone 3% homosalate 10% octisalate  
5% spray**

**The Soleil Group**

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**Soleil Toujours SPF 50 Spray**

**Active ingredients**

Avobenzone 3.00%  
Homosalate 10.00%  
Octisalate 5.00%.

**Purpose**

Sunscreen  
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Sunscreen

**Uses**

- helps prevent sunburn•

**Warnings**

**For external use only.**

**Flammable:**

Do not use near heat, flame or while smoking.

**Do not use**

on damaged or broken skin.

**When using this product**

- keep out of eyes. Rinse with water to remove. • Do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120°F.

**Stop use and ask doctor if**

rash occurs.

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center immediately

**Directions**

- spray liberally and spread evenly by hand 15 minutes before sun exposure• reapply: • after 80 minutes of swimming or sweating • immediately after towel drying • at least every 2 hours
- 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 a well-ventilated area
- 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 a long-sleeve shirt, pants, hats, and sunglasses.
- children under 6 months: Ask a doctor.

## Inactive Ingredieints

Alcohol Denat.  
 Butyloctyl Salicylate  
 Fragrance  
 Oryza Sativa (Rice) Bran Oil  
 Polyester-8  
 VA/Butyl Maleate/Isobornyl Acrylate Copolymer



<b>SOLEIL SET SUNSCREEN SPF 50</b>			
avabenzone 3% homosalate 10% octisalate 5% spray			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69630-006

Route of Administration TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 g
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (ETHYLHEXYL SALICYLATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>FRAGRANCE VERBENA JUNIPERBERRY ORC1403606</b> (UNII: A59PZZ5O26)	
<b>POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENOYL CAPPED)</b> (UNII: T9296U138P)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>ALOE</b> (UNII: V5VD430YW9)	
<b>ASCORBYL PALMITATE</b> (UNII: QN83US2B0N)	
<b>BISABOLOL</b> (UNII: 24WE03BX2T)	
<b>CAMELLIA OLEIFERA LEAF</b> (UNII: 5077EL0C60)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PORPHYRIDIUM PURPUREUM</b> (UNII: K2P8K2558N)	
<b>SODIUM HYALURONATE</b> (UNII: YSE9PPT4TH)	
<b>TETRAHEXYLDECYL ASCORBATE</b> (UNII: 9LBV3F07AZ)	
<b>.ALPHA.-TOCOPHEROL ACETATE, DL-</b> (UNII: WR1WPI7EW8)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69630-006-02	48 g in 1 BOTTLE; Type 0: Not a Combination Product	08/06/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/06/2025	

**Labeler** - The Soleil Group (079694651)

**Registrant** - Inspec Solutions (081030372)

### Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions		081030372	manufacture(69630-006)

