

SUN PROTECTION SPF 50- avobenzene 2.9%, homosalate 4.0%, octisalate 4.9%, octocrylene 9.5% spray
Skinresource.md, L.L.C.

Drug Facts

Active Ingredient

Avobenzene – 2.9% - Homosalate – 4.0% - Octisalate – 4.9% - Octocrylene – 9.5%

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Seed Oil*, Polyester-8, Tocopherol.

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Keep out of reach of children

■ If product is swallowed, get medical help or contact a Poison Control Center right away

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Warnings

■ For external use only ■ Flammable Do not spray near heat, sparks, sources of ignition, or flames.
Protect this product from excessive heat and direct sun

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 skin irritation or redness occurs for more than 72 hours.

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AVOBENZONE - 2.9% - HOMOSALATE - 4.0% - OCTISALATE - 4.9% - OCTOCRYLENE - 9.0%

Purpose Sunscreen

Directions

- Apply liberally 15 minutes before sun exposure.

Reapply:

- After 80 minutes of swimming or sweating.
- Immediately after towel drying.
- At least every 2 hours.
- Children under 6 months:
Ask a doctor.



Directions • Apply liberally 15 minutes before sun exposure. Reapply: Every 2 hours during the day or after swimming or sweating. • Children under 6 months: ask a doctor.

Sun Protection Measure: 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 value of 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-sleeved shirts, pants, hats, and sun glasses.

Uses • Helps prevent sunburn • If used as directed with other sun protection measures, decreases the risk of skin cancer and early skin aging caused by the sun.

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Sun Protection Spray SPF 50

Prescribe it for Yourself.®



6 Fl. Oz. (177 mL)

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Questions? call (619) 776-3464 Option 3
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SUN PROTECTION SPF 50			
avobenzone 2.9%, homosalate 4.0%, octisalate 4.9%, octocrylene 9.5% spray			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83694-011
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7.08 g in 177 g
	OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	16.815 g in 177 g
	OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	8.673 g in 177 g
	AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	5.133 g in 177 g
Inactive Ingredients			
	Ingredient Name		Strength
	ETHYL FERULATE (UNII: 5B8915UELW)		

ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)	
HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL (UNII: 3W1JG795YI)	
DEHYDRATED ALCOHOL (UNII: 3K9958V90M)	
POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENOYL CAPPED) (UNII: T9296U138P)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ACRYLATES/OCTYLACRYLAMIDE COPOLYMER (75000 MW) (UNII: JU3XHR8VWK)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
CARTHAMUS TINCTORIUS (SAFFLOWER) SEED OIL (UNII: 65UEH262IS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83694-011-01	177 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	09/01/2024	

Labeler - Skinresource.md, L.L.C. (022050636)

Revised: 6/2025

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