

**COPPERTONE TANNING SUNSCREEN SPF 8- avobenzone 2%, homosalate 5%, octisalate 4.5%, octocrylene 3% lotion**  
**Beiersdorf Inc**

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**Coppertone Tanning Lotion SPF 8**

***Drug Facts***

***Active ingredients***

Avobenzone 2%, Homosalate 5%, Octisalate 4.5%, Octocrylene 3%

***Purpose***

Sunscreen

***Use***

■ helps prevent sunburn

***Warnings***

**Skin Cancer/Skin Aging Alert:** Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to help prevent sunburn, not skin cancer or early skin aging.

**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 product is swallowed, get medical help or contact a Poison Control Center right away.

***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

### ***Other information***

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

### ***Inactive ingredients***

water, aluminum starch octenylsuccinate, glycerin, silica, phenoxyethanol, isododecane, ethylhexylglycerin, polyester-27, styrene/acrylates copolymer, acrylates/C10-30 alkyl acrylate crosspolymer, arachidyl alcohol, neopentyl glycol diheptanoate, glyceryl stearate, PEG-100 stearate, tocopherol, beeswax, fragrance, behenyl alcohol, arachidyl glucoside, potassium hydroxide, disodium EDTA, sodium ascorbyl phosphate

### ***Questions***

1-866-288-3330

Coppertone<sup>®</sup> Sunscreen Lotion

Tanning

Lightweight & Moisturizing 8

No Dyes, PABA, Oxybenzone

Water Resistant (80 Minutes)

Broad Spectrum SPF 8

8 FL OZ (237 mL)



## COPPERTONE TANNING SUNSCREEN SPF 8

avobenzone 2%, homosalate 5%, octisalate 4.5%, octocrylene 3% lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66800-4086
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	3 g in 100 g
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2 g in 100 g
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.5 g in 100 g
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	5 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>FRAGRANCE FLORAL ORC0902236</b> (UNII: R66Z4YW3X0)	
<b>ISODODECANE</b> (UNII: A8289P68Y2)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>POTASSIUM HYDROXIDE</b> (UNII: WZ3H3C48M4T)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PHENOXYETHANOL</b> (UNII: H1E492ZZ3T)	

<b>POLYESTER-7</b> (UNII: 0841698D2F)
<b>DOCOSANOL</b> (UNII: 9G10E216XY)
<b>STYRENE/ACRYLAMIDE COPOLYMER (MW 500000)</b> (UNII: 5Z4DPO246A)
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)
<b>ACRYLATES CROSSPOLYMER-6</b> (UNII: 4GXD0Q3OS3)
<b>ARACHIDYL GLUCOSIDE</b> (UNII: 6JVW35JOOJ)
<b>DISODIUM EDTA-COPPER</b> (UNII: 6V475AX06U)
<b>SODIUM ASCORBYL PHOSPHATE</b> (UNII: 836SJG51DR)
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)
<b>ARACHIDYL ALCOHOL</b> (UNII: 1QR1QRA9BU)
<b>NEOPENTYL GLYCOL DIHEPTANOATE</b> (UNII: 5LKW3C543X)
<b>GLYCERYL STEARATE SE</b> (UNII: FCZ5MH785I)
<b>SYNTHETIC BEESWAX</b> (UNII: 08MNR5YE2R)
<b>ALUMINUM STARCH OCTENYLSUCCINATE</b> (UNII: I9PJ0O6294)

### Product Characteristics

<b>Color</b>	white (White to Off-White)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66800-4086-1	237 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/03/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/03/2020	

**Labeler** - Beiersdorf Inc (001177906)