SKINCEUTICALS LIGHT MOISTURE UV DEFENSE BROAD SPECTRUM SPF 50 SUNSCREEN- avobenzone, homosalate, octisalate and octocrylene lotion L'Oreal USA Products Inc

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 10%

Octisalate 5%

Octocrylene 7%

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

For sunscreen use:

- apply liberally 15 minutes before sun exposure
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- 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 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

dwater, styrene/acrylates copolymer, dimethicone, polymethylsilsesquioxane, butyloctyl salicylate,

glycerin, alcohol denat., poly c10-30 alkyl acrylate, caprylyl methicone, trisiloxane, acrylates/c10-30 alkyl acrylate crosspolymer, acrylates/dimethicone copolymer, caprylic/capric triglyceride, caprylyl glycol, diethylhexyl, syringylidenemalonate, disodium EDTA, glyceryl stearate, inulin lauryl carbamate, p-anisic acid, PEG-100 stearate, PEG-8 laurate, phenoxyethanol, potassium cetyl phosphate, tocopherol, triethanolamine, xanthan gum

Questions or comments?

Call **1-800-811-1660**

Monday - Friday (9 a.m. - 5 p.m. CST)





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avobenzone, homosalate, octisalate and octocrylene lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:49967-888

| Active Ingredient/Active Moiety | | | | |
|--|--------------------------|----------------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX) | AVOBENZ ONE | 30 mg in 1 mL | | |
| HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII: V06SV4M95S) | HOMOSALATE | 100 mg in 1 mL | | |
| OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W) | OCTISALATE | 50 mg in 1 mL | | |
| OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM) | OCTOCRYLENE | 70 mg in 1 mL | | |

| Inactive Ingredients | |
|---|----------|
| Ingredient Name | Strength |
| WATER (UNII: 059QF0KO0R) | |
| STYRENE/ACRYLAMIDE COPOLYMER (500000 MW) (UNII: 5Z4DPO246A) | |
| DIMETHICONE (UNII: 92RU3N3Y1O) | |
| POLYMETHYLSILSESQUIOXANE (11 MICRONS) (UNII: Z570VEV8XK) | |
| BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| ALCOHOL (UNII: 3K9958V90M) | |
| CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL) | |
| TRISILOXANE (UNII: 9G1ZW13R0G) | |
| CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO) | |
| MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) | |
| CAPRYLYL GLYCOL (UNII: 00YIU5438U) | |
| DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248) | |
| EDETATE DISODIUM (UNII: 7FLD91C86K) | |
| GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4) | |
| P-ANISIC ACID (UNII: 4SB6Y7DMM3) | |
| PEG-100 STEARATE (UNII: YD01N1999R) | |
| PEG-8 LAURATE (UNII: 76208IWA10) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |
| POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT) | |
| TOCOPHEROL (UNII: R0ZB2556P8) | |
| TROLAMINE (UNII: 903K93S3TK) | |
| XANTHAN GUM (UNII: TTV12P4NEE) | |

| Packaging | | | | |
|-----------|----------------------|---|-------------------------|-----------------------|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:49967-888- 01 | 1 in 1 CARTON | 03/01/2018 | 09/22/2025 |
| 1 | | 200 mL in 1 TUBE; Type 0: Not a Combination Product | | |
| 2 | NDC:49967-888- 02 | 1 in 1 CARTON | 03/01/2018 | 09/22/2025 |
| 2 | | 4 mL in 1 TUBE; Type 0: Not a Combination Product | | |
| | | | | |

| Marketing Information | | | | |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | M020 | 03/01/2018 | 09/22/2025 | |
| | | | | |

Labeler - L'Oreal USA Products Inc (002136794)

| Establishment | | | | | |
|-----------------------|---------|-----------|----------------------------|--|--|
| Name | Address | ID/FEI | Business Operations | | |
| Cosmetic Essence, LLC | | 032565959 | manufacture(49967-888) | | |

Revised: 12/2023 L'Oreal USA Products Inc