KIEHLS SINCE 1851 DERMATOLOGIST SOLUTIONS AUTOTONE DISCOLORATION AND UV SOLUTION BROAD SPECTRUM SPF 30 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

water, C15-19 alkane, niacinamide, glycerin, butylene glycol, silica, titanium dioxide, hydroxyacetophenone, pentaerythrityl tetra-di-t-butyl hydroxyhydrocinnamate, potassium hydroxide, sodium thiosulfate, taraxacum officinale (dandelion) rhizome/root extract, trisodium ethylenediamine disuccinate, 2-mercaptonicotinoyl glycine, butyrospermum parkii (shea) butter, ammonium acryloyldimethyltaurate/VP copolymer, arachidyl alcohol, arachidyl glucoside, behenyl alcohol, caprylyl glycol, carbomer, citric acid, mica, pentylene glycol, poly c10-30 alkyl acrylate, polyglyceryl-4 caprate, sodium dilauramidoglutamide lysine, tin oxide, xanthan gum, red 4, phenoxyethanol

Questions or comments?

Call toll free 1-800-946-4453

Drug Facts (continued)

water to remove

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Directions

For sunscreen use:

- . apply generously and evenly 15 minutes before sun exposure
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Drug Facts (continued)

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Fmla 685904 3 F.I.L. Code V70039198 PAT: patent.kiehls.com







KIEHL'S SINCE 1851 LLC, NEW YORK, NY 10014 + 399659 Ingredients sourced worldwide, made in U.S.A.
DIST, KIEHL'S CANADA, MONTREAL HAT 1KS
14 PUE ROYALE 7500B PARIS + CONDON W12 7SA
ISA 75000 93584 ST OUEN CEDEX FR + www.kiehls.com







DERMATOLOGIST SOLUTIONS N

Auto-Tone Discoloration & UV Solution

SUNSCREEN Broad Spectrum SPF 30

with 0.5% Melasyl

Clinically Demonstrated* to Visibly Improve the Appearance of Discoloration for a Brighter, More Even Skin Tone

1.35 fl. oz. - 40 ml

*Based on expert assessment in a clinical study

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avobenzone, homosalate, octisalate and octocrylene lotion

Product Information

HUMAN OTC DRUG NDC:49967-069 **Product Type** Item Code (Source)

Route of Administration TOPICAL

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) | |
| C15-19 ALKANE (UNII: CI87N1IM01) | |
| NIACINAMIDE (UNII: 25X51I8RD4) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| BUTYLENE GLYCOL (UNII: 3XUS85K0RA) | |
| SILICON DIOXIDE (UNII: ETJ7Z 6XBU4) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| HYDROXYACETOPHENONE (UNII: G1L3HT4CMH) | |
| PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS) | |
| POTASSIUM HYDROXIDE (UNII: WZ H3C48M4T) | |
| SODIUM THIOSULFATE (UNII: HX1032V43M) | |
| TARAXACUM OFFICINALE ROOT (UNII: 9DE5YCO0RU) | |
| TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q) | |
| SHEA BUTTER (UNII: K49155WL9Y) | |
| AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG) | |
| ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU) | |
| ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ) | |
| DOCOSANOL (UNII: 9G10E216XY) | |
| CAPRYLYL GLYCOL (UNII: 00YIU5438U) | |
| CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) | |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) | |
| MICA (UNII: V8A1AW0880) | |
| PENTYLENE GLYCOL (UNII: 50C1307PZG) | |
| BEHENYL ACRYLATE POLYMER (UNII: D64PM5UT4U) | |
| POLYGLYCERYL-4 CAPRATE (UNII: 3N873UN885) | |
| SODIUM DILAURAMIDOGLUTAMIDE LYSINE (UNII: MNJ7VPT2R5) | |
| STANNIC OXIDE (UNII: KM7N50LOS6) | |
| XANTHAN GUM (UNII: TTV12P4NEE) | |
| FD&C RED NO. 4 (UNII: X3W0AM1JLX) | |
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |

| Packaging | | | | | |
|-----------|----------------------|---|-------------------------|-----------------------|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | |
| 1 | NDC:49967-069- 01 | 1 in 1 CARTON | 06/15/2024 | | |
| 1 | | 40 mL in 1 TUBE; Type 0: Not a Combination Product | | | |
| 2 | NDC:49967-069- 02 | 1 in 1 CARTON | 06/15/2024 | | |
| 2 | | 15 mL in 1 TUBE; Type 0: Not a Combination Product | | | |
| 3 | NDC:49967-069- 03 | 2 mL in 1 PACKET; Type 0: Not a Combination Product | 06/15/2024 | | |
| | | | | | |

| Marketing Information | | | |
|-----------------------|---|-------------------------|-----------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC Monograph Drug | M020 | 06/15/2024 | |
| | | | |

Labeler - L'Oreal USA Products Inc (002136794)

| Establishment | | | | |
|-------------------|---------|-----------|------------------------|--|
| Name | Address | ID/FEI | Business Operations | |
| L'Oreal USA, Inc. | | 185931458 | manufacture(49967-069) | |

Revised: 6/2024 L'Oreal USA Products Inc