KIEHLS SINCE 1851 DERMATOLOGIST SOLUTIONS SUPER FLUID DAILY UV DEFENSE BROAD SPECTRUM SPF 50 PLUS SUNSCREEN ANTIPOLLUTION LIGHTWEIGHT FORMULA- avobenzone, homosalate, octisalate, and octocrylene lotion
L'Oreal USA Products Inc.

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 15%

Octisalate 5%

Octocrylene 10%

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

Flammble until dry. Do not use near fire, flame or heat.

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

- shake well before 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. - 2p.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, dimethicone, alcohol denat., silica, styrene/acrylates copolymer, butyloctyl salicylate, PEG-30 dipolyhydroxystearate, polymethylsilsesquioxane, tocopherol, phenoxyethanol, PEG-8 laurate, lauryl PEG/PPG-18/18 methicone, sodium chloride, caprylyl glycol, isostearyl alcohol, poly c10-30 alkyl acrylate, p-anisic acid, pentaerythrityl tetra-di-t-butyl hydroxyhydrocinnamate, trisodium ethylenediamine disuccinate, sodium dodecylbenzenesulfonate

Our advanced sunscreen, formulated with effective UVA/UVB filters and pollution-deflecting technology, provides protection against environmental aggressors. When used as directed with other sun protection measures (see **Directions**), this lightweight fluid helps prevent early skin aging caused by the sun* and visibly improves rough skin. Our formula feels gentle on skin, leaving a soft, matte finish and no residue.
Dermatologist-tested for safety and suitable for sensitive skin.

Kiehl's Dermatologist Solutions are highly advanced targeted treatments developed by Kiehl's Since 1851. Relying on our extensive skincare expertise and botanical knowledge, Kiehl's chemists partner with an international team of leading dermatologists to deliver powerful, yet safe skincare solutions.

SINCE 1851

KIEHL'S

DERMATOLOGIST SOLUTIONS™

Super Fluid Daily UV Defense

SUNSCREEN

Broad Spectrum SPF 50+

ANTI-POLLUTION

Advanced UVA/UVB Technology
Helps Prevent Early Skin Aging
Caused by the Sun*

Lightweight Formula for All Skin Types

SHAKE WELL

Fragrance-Free
Non-Comedogenic
Paraben-Free

1.7 fl. oz. - 50 ml

*When used with sun protection measures (see **Directions**)

FSC LOGO
FPO

PAP

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KIEHL'S SINCE 1851 LLC
NEW YORK, NY 10014
Ingredients sourced worldwide, made in U.S.A.
Dist. Kiehl's Canada, Montreal H4T 1K5
14 rue Royale 75008 Paris · LONDON W6 8AZ
TSA 75000 93584 ST OUEN CEDEX FR
www.kiehls.com
2093628

3 605971 618123

Drug Facts

Active ingredients	Purpose
Avobenzone 3%	Sunscreen
Homosalate 15%	Sunscreen
Octisalate 5%	Sunscreen
Octocrylene 10%	Sunscreen

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Warnings
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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:
- shake well before 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
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Inactive ingredients
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Questions or comments?
Call toll free **1-800-946-4453**

Fmla 685777 15 F.LL. Code D276520/1 PAT: patents.kiehls.com

KIEHLS SINCE 1851 DERMATOLOGIST SOLUTIONS SUPER FLUID

DAILY UV DEFENSE BROAD SPECTRUM SPF 50 PLUS SUNSCREEN ANTIPOLLUTION LIGHTWEIGHT FORMULA

avobenzone, homosalate, octisalate, and octocrylene lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-812
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	150 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALCOHOL (UNII: 3K9958V90M)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STYRENE/ACRYLAMIDE COPOLYMER (500000 MW) (UNII: 5Z4DPO246A)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
PEG-30 DIPOLYHYDROXYSTEARATE (UNII: 9713Q0S7FO)	
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PEG-8 LAURATE (UNII: 762O8IWA10)	
LAURYL PEG/PPG-18/18 METHICONE (UNII: ZJ5S27D9NX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ISOSTEARYL ALCOHOL (UNII: Q613OCQ44Y)	
P-ANISIC ACID (UNII: 4SB6Y7DMM3)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-812-01	1 in 1 CARTON	10/01/2022	
1		125 mL in 1 TUBE; Type 0: Not a Combination Product		
	NDC:49967-812			

2	NDC:49967-812-02	1 in 1 CARTON	10/01/2022	
2		50 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:49967-812-03	125 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/01/2022	

Labeler - L'Oreal USA Products Inc. (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
Dimensional Merchandising Inc.		076693183	manufacture(49967-812) , pack(49967-812)

Revised: 12/2023 L'Oreal USA Products Inc.