

KIEHLS SINCE 1851 FACIAL FUEL DAILY ENERGIZING MOISTURE TREATMENT FOR MEN BROAD SPECTRUM SPF 20 SUNSCREEN- avobenzone, homosalate, octisalate and octocrylene lotion
L'Oreal USA Products Inc

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 8.8%

Octisalate 4.9%

Octocrylene 5.9%

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, dimethicone, glycerin, alcohol denat., tocopherol, aloe barbadensis leaf juice, phenoxyethanol, sodium hydroxide, caprylyl glycol, sodium polyacrylate, fragrance, acrylates/c10-30 alkyl acrylate crosspolymer, ascorbyl glucoside, acrylates/dimethicone copolymer, caffeine, sodium hyaluronate, xanthan gum, menthyl lactate, disodium EDTA, limonene, ceramide NP, castanea sativa (chestnut) seed extract, pentylene glycol, hydrolyzed soy protein, linalool, citric acid, potassium sorbate, citral, sodium benzoate, ethylhexylglycerin

Questions or comments?

Call toll free 1-800-946-4453

SINCE **KIEHL'S** 1851

For a Refuel Land at Kiehl's



FACIAL FUEL

DAILY ENERGIZING MOISTURE
TREATMENT FOR MEN
SUNSCREEN

**BROAD SPECTRUM
SPF 20**

Our energizing, lightweight water gel moisturizer instantly refreshes and revitalizes the look of dull, fatigued skin while providing protection against the sun's harmful UVA/UVB rays. Delivering all day hydration, with an invigorating aroma, this moisturizer provides anti-pollution protection helping to defend against oxidative stress. Our formula with Vitamin C and E, Chestnut Extract and Caffeine leaves skin feeling refueled and re-energized. Appropriate for all skin types.

6.8 fl. oz. - 200 ml

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Fmla 685783 35 F.I.L. Code D224591/1

KIEHL'S SINCE 1851 LLC, NEW YORK, NY 10014
Ingredients sourced worldwide, made in U.S.A.
Dist., Kiehl's Canada, Montreal H4T 1K5
106 rue Danton 92691 Levallois Perret Cedex
TSA 75000 93584 ST. QUEN CEDEX FR
www.kiehls.com
1045871



UPC

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avobenzone, homosalate, octisalate and octocrylene lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-608
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	88 mg in 1 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	49 mg in 1 mL

Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	59 mg in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALCOHOL (UNII: 3K9958V90M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ASCORBYL GLUCOSIDE (UNII: 2V52RONHXW)	
CAFFEINE (UNII: 3G6A5W338E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
XANTHAN GUM (UNII: TTV12P4NEE)	
METHYL LACTATE, (-)- (UNII: 0379G9C44S)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
CERAMIDE NP (UNII: 4370DF050B)	
SPANISH CHESTNUT (UNII: 2MT5XMR2YW)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
HYDROLYZED SOY PROTEIN (ENZYMATIC; 2000 MW) (UNII: 1394NXB9L6)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
CITRAL (UNII: T7EU009VPP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-608-01	1 in 1 CARTON	03/29/2018	
1		125 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:49967-608-02	1 in 1 CARTON	03/29/2018	
2		15 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:49967-608-03	1 in 1 CARTON	12/10/2018	
3		75 mL in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:49967-608-04	1 in 1 CARTON	12/10/2018	

4		200 mL in 1 TUBE; Type 0: Not a Combination Product	
5	NDC:49967-608-05	3 mL in 1 PACKET; Type 0: Not a Combination Product	12/10/2018

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/29/2018	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA Products Inc		185931458	manufacture(49967-608) , pack(49967-608)

Revised: 12/2023

L'Oreal USA Products Inc