

**KIEHL'S SINCE 1851 ULTRA FACIAL MOISTURIZER BROAD SPECTRUM SPF 30
SUNSCREEN FOR ALL SKIN TYPES- avobenzone, homosalate, octisalate and
octocrylene lotion
L'Oreal USA Products Inc**

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 7%

Octisalate 5%

Octocrylene 5%

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, propylene glycol, dicaprylyl ether, glycerin, PEG-100 stearate, glyceryl stearate, squalane, hydrogenated polyisobutene, phenoxyethanol, sorbitan oleate, caprylyl glycol, acrylates/c10-30 alkyl acrylate crosspolymer, triethanolamine, beeswax, tocopherol, disodium EDTA, stearic acid, myristyl alcohol, prunus amygdalus dulcis (sweet almond) oil, prunus armeniaca (apricot) kernel oil, persea gratissima (avocado) oil

Questions or comments?

Call toll free **1-800-946-4453**

SINCE **KIEHL'S** 1851

**ULTRA FACIAL
MOISTURIZER**

SUNSCREEN
Broad Spectrum
SPF 30

for all skin types

Our uniquely lightweight formulation is readily absorbed into skin, leaving it soft, smooth and nurtured. This superb formula also helps protect against skin-damaging UVA and UVB rays with broad spectrum SPF 30.

2.5 fl. oz. - 75 ml

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Homosalate 7%	Sunscreen
Octisalate 5%	Sunscreen
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Fmle 685001/16 Code FLL D54062/2
KIEHL'S SINCE 1851 LLC
NEW YORK, NY 10014
MADE IN U.S.A.
Dist. Kiehl's Canada
Montreal H4T 1K51
106 rue Danton 92691
Levallois Perret Cedex
TSA 10007
E 92667 ASSNIERES CROIX
www.kiehls.com
509002XX

3605970387976

Lift

**Daylot Box
Does Not Print**

Space
0-3125

Drug Facts (continued)

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Drug Facts (continued)

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avobenzone, homosalate, octisalate and octocrylene lotion				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-797	
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	70 mg in 1 mL	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)		OCTISALATE	50 mg in 1 mL	
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)		OCTOCRYLENE	50 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
DICAPRYLYL ETHER (UNII: 77JZM5516Z)				
GLYCERIN (UNII: PDC6A3C0OX)				
PEG-100 STEARATE (UNII: YD01N1999R)				
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)				
SQUALANE (UNII: GW89575KF9)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
CAPRYLYL GLYCOL (UNII: 00YIU5438U)				
TOCOPHEROL (UNII: R0ZB2556P8)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
MYRISTYL ALCOHOL (UNII: V42034O9PU)				
ALMOND OIL (UNII: 66YXD4DKO9)				
APRICOT KERNEL OIL (UNII: 54JB35T06A)				
AVOCADO OIL (UNII: 6VNO72PFC1)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-797-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	

2	NDC:49967-797-02	125 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	
3	NDC:49967-797-03	75 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	
4	NDC:49967-797-04	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	
5	NDC:49967-797-05	3 mL in 1 PACKET; Type 0: Not a Combination Product	03/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/01/2013	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-797)

Revised: 12/2023

L'Oreal USA Products Inc