KIEHLS SINCE 1851 ULTRA FACIAL BROAD SPECTRUM SPF 30 SUNSCREEN-
avobenzone, homosalate, octisalate and octocrylene cream
L'Oreal USA Products Inc

Drug Facts

Active ingredient

Avobenzone 3%

Homosalate 5%

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

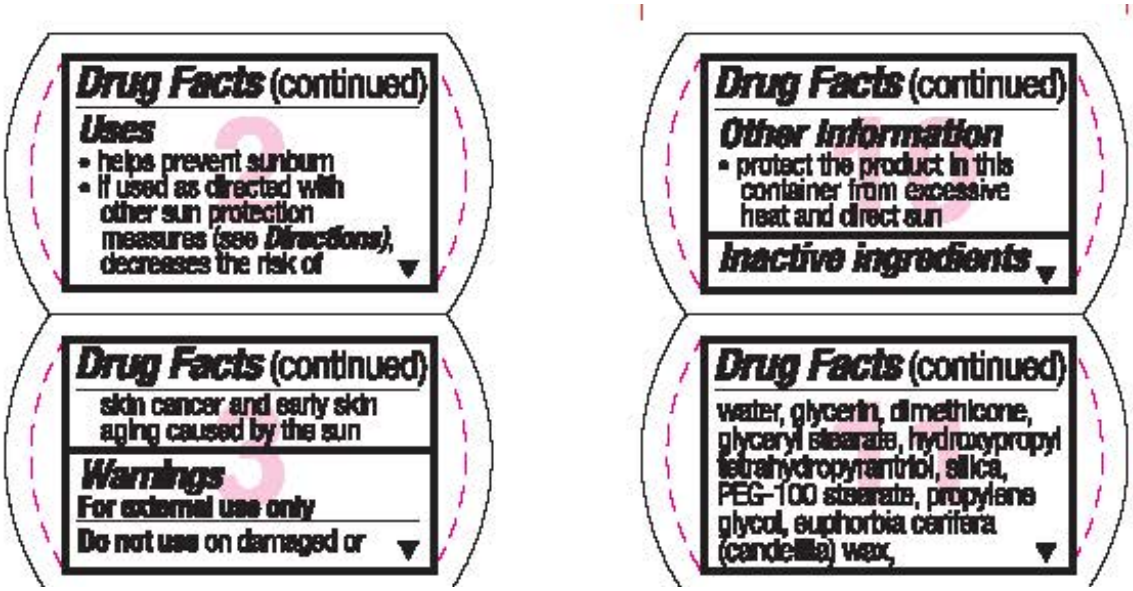
Inactive ingredients

water, glycerin, dimethicone, glyceryl stearate, hydroxypropyl tetrahydropyrantriol, silica, PEG-100 stearate, propylene glycol, euphorbia cerifera (candelilla) wax, phenoxyethanol, ammonium polyacryloyldimethyl taurate, stearic acid, dicaprylyl carbonate, cetyl alcohol, palmitic acid, capryloyl salicylic acid, caprylyl glycol, xanthan gum, dimethicone/vinyl dimethicone crosspolymer, fragrance, disodium EDTA, tocopherol, sodium hyaluronate, adenosine, linalool, sodium hydroxide, jasminum officinale (jasmine) flower extract, citronellol, geraniol, citral, benzyl alcohol, citric acid

Questions or comments?

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

Monday - Friday (9 a.m. to 5 p.m. EST)



Drug Facts (continued)

broken skin

When using this product keep out of eyes. Rinse with water to remove.

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Drug Facts (continued)

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Drug Facts (continued)

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- apply liberally 15 minutes before sun exposure
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Drug Facts (continued)

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Drug Facts (continued)

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.

Drug Facts (continued)

- wear long-sleeved shirts, pants, hats and sunglasses
- children under 6 months of age: Ask a doctor

Continued on back

Drug Facts (continued)

phenoxyethanol, ammonium polyacryloyldimethyl taurate, stearic acid, dicaprylyl carbonate, cetyl alcohol, palmitic acid, caprylcoyl salicylic acid, caprylyl glycol, xanthan

Drug Facts (continued)

gum, dimethicone/vinyl dimethicone crosspolymer, fragrance, disodium EDTA, tocopherol, sodium hyaluronate, adenosine,

Drug Facts (continued)

linalool, sodium hydroxide, jasminum officinale (jasmine) flower extract, citronellol, geraniol, citral, benzyl alcohol, citric acid

Drug Facts (continued)

Questions or comments?
Call toll free 1-800-946-4453

Fmla 685681 40 F.L. Code D174030/1

GLUE PANEL

Drug Facts Continued on reverse

Lift

DAY LOT
0.5"x0.875"

Drug Facts (continued)

Uses

- helps prevent sunburn
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Drug Facts (continued)

skin cancer and early skin aging caused by the sun

Warnings

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Do not use on damaged or

Drug Facts (continued)

broken skin

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Drug Facts (continued)

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Directions

.4"

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Drug Facts (continued)

Other Information

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Inactive ingredients

Drug Facts (continued)

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Drug Facts (continued)

Questions or comments?

Call toll free 1-800-946-4483

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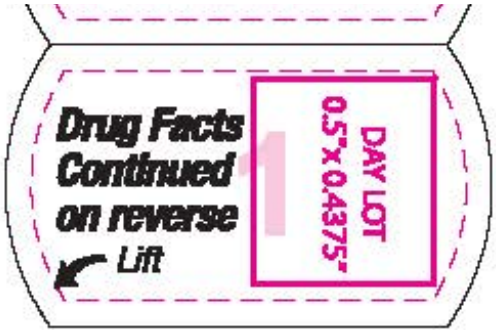
GLUE PANEL



KIEHLS SINCE 1851 ULTRA FACIAL BROAD SPECTRUM SPF 30 SUNSCREEN

avobenzone, homosalate, octisalate and octocrylene cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-941
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	50 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)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
HYDROXYPROPYL TETRAHYDROPYRANTRIOL (UNII: 4U3GMG1OT1)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CANDELILLA WAX (UNII: WL0328HX19)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (55000 MPA.S) (UNII: F01RIY4371)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
PALMITIC ACID (UNII: 2V16EO95H1)	
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (HARD PARTICLE) (UNII: H895X08VNQ)	

EDETATE DISODIUM (UNII: 7FLD91C86K)	
TOCOPHEROL (UNII: R0ZB2556P8)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
ADENOSINE (UNII: K72T3FS567)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
JASMINUM OFFICINALE FLOWER (UNII: 0Q8K841432)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
GERANIOL (UNII: L837108USY)	
CITRAL (UNII: T7EU0O9VPP)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-941-01	50 mL in 1 JAR; Type 0: Not a Combination Product	07/01/2015	
2	NDC:49967-941-02	125 mL in 1 JAR; Type 0: Not a Combination Product	07/01/2015	
3	NDC:49967-941-03	1 in 1 CARTON	07/01/2015	
3		75 mL in 1 JAR; Type 0: Not a Combination Product		
4	NDC:49967-941-04	3 mL in 1 PACKET; Type 0: Not a Combination Product	07/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	07/01/2015	

Labeler - L’Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L’Oreal USA, Inc.		185931458	manufacture(49967-941)

Revised: 12/2023 L’Oreal USA Products Inc