

**KIEHLS SINCE 1851 BROAD SPECTRUM SPF 30 SUNSCREEN POWERFUL
WRINKLE REDUCING- avobenzone, homosalate, octisalate and
octocrylene cream
L'Oreal USA Products Inc**

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 2.7%

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, cetearyl alcohol, isohexadecane, glyceryl stearate, cyclohexasiloxane, isononyl isononanoate, butyrospermum parkii (shea) butter, PEG-100 stearate, cetareth-20, phenoxyethanol, butylene glycol, ammonium polyacryloyldimethyl taurate, caprylyl glycol, tocopherol, chlorphenesin, xanthan gum, disodium EDTA, p-anisic acid, calcium PCA, adenosine, copper PCA, alteromonas ferment extract, corallina officinalis extract, citric acid, sodium hydroxide

Questions or comments?

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

SINCE *Kiehl's* 1851

1.7 fl. oz.
- 50 ml

**POWERFUL WRINKLE
REDUCING CREAM**

SUNSCREEN
Broad Spectrum SPF 30
with Copper PCA and Calcium PCA
Visibly Repairs Wrinkles and Refines Texture

Our visibly reparative treatment, formulated with Copper PCA and Calcium PCA, helps strengthen skin's elasticity, an important factor to reduce the appearance of wrinkles and help resist new ones. The result is a smoother skin surface, refined texture and less visible pores. When used along with other sun protective measures (see *Directions*), this SPF 30 cream provides broad-spectrum UV protection to help decrease the risk of early skin aging caused by the sun.

KIEHL'S SINCE 1851 LLC, NEW YORK, NY 10014
MADE IN U.S.A.

Dist., Kiehl's Canada, Montreal B4T 1K5
105 rue Duquesne 5491 Laval (Old Port) Quebec
TSA 10007 E 92667 ASNIERES CROIX
www.kiehls.com
S08601XX

Active Ingredients	Purpose
Azobenzene 3%	Sunscreen
Homosalate 2.7%	Sunscreen
Octisalate 5%	Sunscreen
Octocrylene 7%	Sunscreen

Drug Facts continued on bottom

3.487

6.974

Die # 17066_CIRCLE-15 PANELS

Size 1 100"

SIZE 1.100
Opening Side: Right

Position #'s 1 or 2.

1
2
rels

Closed view



DO NOT PRINT;
 INDICATES
 DAYLOTCODE
 0.5"x 0.4375"

**DRUG FACTS BOX For
 SPF ≥15 Broad Spectrum
 Non Water Resistant**

Title: 9pt
 Continued Title: 8pt
 Headings: 8pt
 Subheadings (bold): 6pt

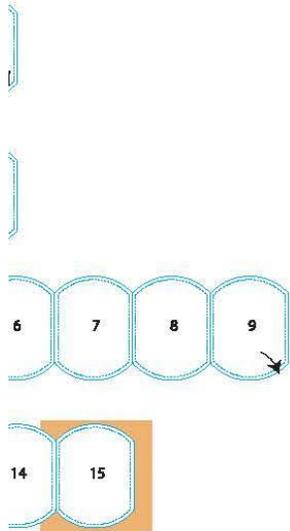


PLATE 2 - Back View

2 **Drug Facts (continued)**
Uses
 • helps prevent sunburn
 • if used as directed with other sun protection measures (see **Directions**), decreases the risk of

3 **Drug Facts (continued)**
 skin cancer and early skin aging caused by the sun
Warnings
 For external use only
 Do not use on damaged or

4 **Drug Facts (continued)**
 broken skin
 When using this product keep out of eyes. Rinse with water to remove.
Stop use and ask a doctor if rash occurs

5 **Drug Facts (continued)**
 Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
Directions

6 **Drug Facts (continued)**
 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

7 **Drug Facts (continued)**
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

8 **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.

9 **Drug Facts (continued)**
 • wear long-sleeved shirts, pants, hats and sunglasses
 • children under 6 months of age: Ask a doctor
 ← Continued on back

PLATE 1 - Front View

10 **Drug Facts (continued)**
Other Information
 • protect the product in this container from excessive heat and direct sun
Inactive ingredients

11 **Drug Facts (continued)**
 water, glycerin, cetearyl alcohol, isohexadecane, glyceryl stearate, cyclohexazone, isononyl isononanoate, butyrospermium parkii (shea) butter,

12 **Drug Facts (continued)**
 PEG-100 stearate, ceteareth-20, phenoxethanol, butylene glycol, ammonium polyacryloyldimethyl taurate, caprylyl glycol, tocopherol, chlorphenesin, xanthan

13 **Drug Facts (continued)**
 gum, disodium EDTA, p-anisic acid, calcium PCA, adenosine, copper PCA, altomonas ferment extract, coralina officinale extract, citric acid,

14 **Drug Facts (continued)**
 sodium hydroxide
Questions or comments?
 Call toll free 1-800-946-4453
 Fmta 86562 31 F.J.L. Code D53887/3

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1 **GLUE PANEL**
 tinted plate

1 **Drug Facts Continued on reverse**
 ← Lift

DO NOT PRINT;
 INDICATES
 DAYLOTCODE
 0.5"x 0.4375"



Die # 17066_CIRCLE-15 PANELS
Size 1.188"
Opening Side: Right

Position #'s 1 or 2.

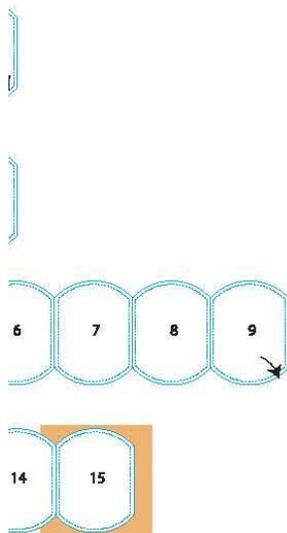
1
2
rels

Closed view



**DRUG FACTS BOX For
 SPF ≥15 Broad Spectrum
 Non Water Resistant**

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Directions

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 • use if water resistant sunscreen if swimming or sweating

7 **Drug Facts (continued)**
 • 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

PLATE 2 Back View

10 **Drug Facts (continued)**
Other information
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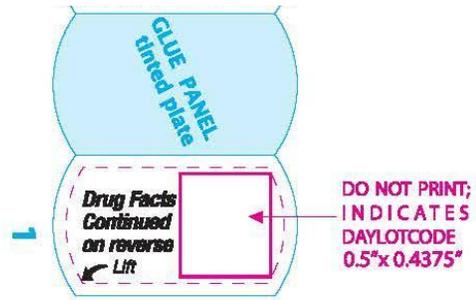
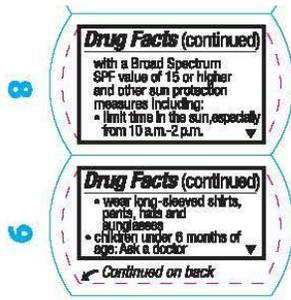
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 PEG-100 stearate, ceteareth-20, phenoxethanol, butylene glycol, ammonium polyacryloyldimethyl taurate, caprylyl glycol, tocopherol, chlorphenesin, xanthan

13 **Drug Facts (continued)**
 gum, disodium EDTA, p-aminic acid, calcium PCA, ascorbyl tetraisostearate, copper PCA, allantoin, ferulic acid, corallina officinalis extract, citric acid,

14 **Drug Facts (continued)**
 sodium hydroxide
Questions or comments?
 Call toll free 1-800-996-4453
 Fm1a 88562 31 F.I.L. Code D53887/3

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PLATE 1 Front View



KIEHLS SINCE 1851 BROAD SPECTRUM SPF 30 SUNSCREEN POWERFUL WRINKLE REDUCING

avobenzone, homosalate, octisalate and octocrylene cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-501
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	27 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)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
XANTHAN GUM (UNII: TTV12P4NEE)	
p-ANISIC ACID (UNII: 4SB6Y7DMM3)	
ADENOSINE (UNII: K72T3FS567)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
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#	Item Code	Package Description	Date	Date
1	NDC:49967-501-01	50 mL in 1 JAR; Type 0: Not a Combination Product	12/01/2012	
2	NDC:49967-501-02	75 mL in 1 JAR; Type 0: Not a Combination Product	12/01/2012	
3	NDC:49967-501-03	3 mL in 1 PACKET; Type 0: Not a Combination Product	12/01/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2012	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-501) , pack(49967-501)

Establishment

Name	Address	ID/FEI	Business Operations
Universal Packaging Systems, Inc.		177711082	pack(49967-501)

Revised: 12/2023

L'Oreal USA Products Inc