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, squalene, dimethicone, PEG-100 stearate, glyceryl stearate, silica, octyldodecanol, stearic acid, phenoxyethanol, palmitic acid, tocopherol, dicaprylyl carbonate, steareth-100, acrylates/c10-30 alkyl acrylate crosspolymer, ophiopogon japonicus root extract, carbomer, chlorphenesin, capryloyl salicylic acid, caprylyl glycol, xanthan gum, dimethicone/vinyl dimethicone crosspolymer, disodium EDTA, sodium hydroxide, citrus aurantium dulcis (orange) peel oil, limonene, ectoin, hydrolyzed hyaluronic acid, myristic acid, mentha piperita (peppermint) oil, pseudoalteromonas ferment extract, ethylhexylglycerin, linalool, salicylic acid

Questions or comments?

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

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

4.2 fl. oz.
- 125 ml

SINCE KIEHL'S 1851

ULTRA FACIAL CREAM

SUNSCREEN

Broad Spectrum SPF 30

24-Hour, Everyday Hydrating Formula with Extracts of Glacial Glycoprotein and Fountain Plant

Inspired by our beloved "Greenland First Ascent" Ultra Facial Cream, this daily, light-textured, 24-hour moisturizer leaves skin feeling balanced—comfortable and hydrated—and helps protect against skin-damaging UVA and UVB rays with broad spectrum SPF 30. Formulated with Antarcticine™, a glycoprotein known to withstand extreme cold, and "Fountain Plant," a hydrating botanical that thrives in dry conditions, this superb moisturizer leaves skin soft, smooth, and healthy-looking.

Drug Facts

Active ingredients	Purpose
Avobenzone 3%	Sunscreen
Homosalate 5%	Sunscreen
Ocifsalate 5%	Sunscreen
Octocrylene 7%	Sunscreen

Drug Facts (continued)

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

DAY LOT CODE DO NOT PRINT

KIEHL'S SINCE 1851 LLC, NEW YORK, NY 10014
MADE IN U.S.A. • Dist. Kiehl's Canada, Montreal H4T 1K5
105 rue Danton 92691 Levallois Perret Cedex
TSA 75000 92584 ST OUEN CEDEX FR • www.kiehls.com

2086121

UPC

Lift

Itop

Drug Facts (continued)

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.

Drug Facts (continued)

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:

0.2500"
6.3500 mm

Base

4.1250"
104.7761 mm

Drug Facts (continued)

- 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, squalane, dimethicone, PEG-100 stearate, glyceryl stearate, silica, octyldodecanol, stearic acid, phenoxyethanol, palmitic acid, tocopherol, dicaprylyl carbonate, steareth-100,

Drug Facts (continued)

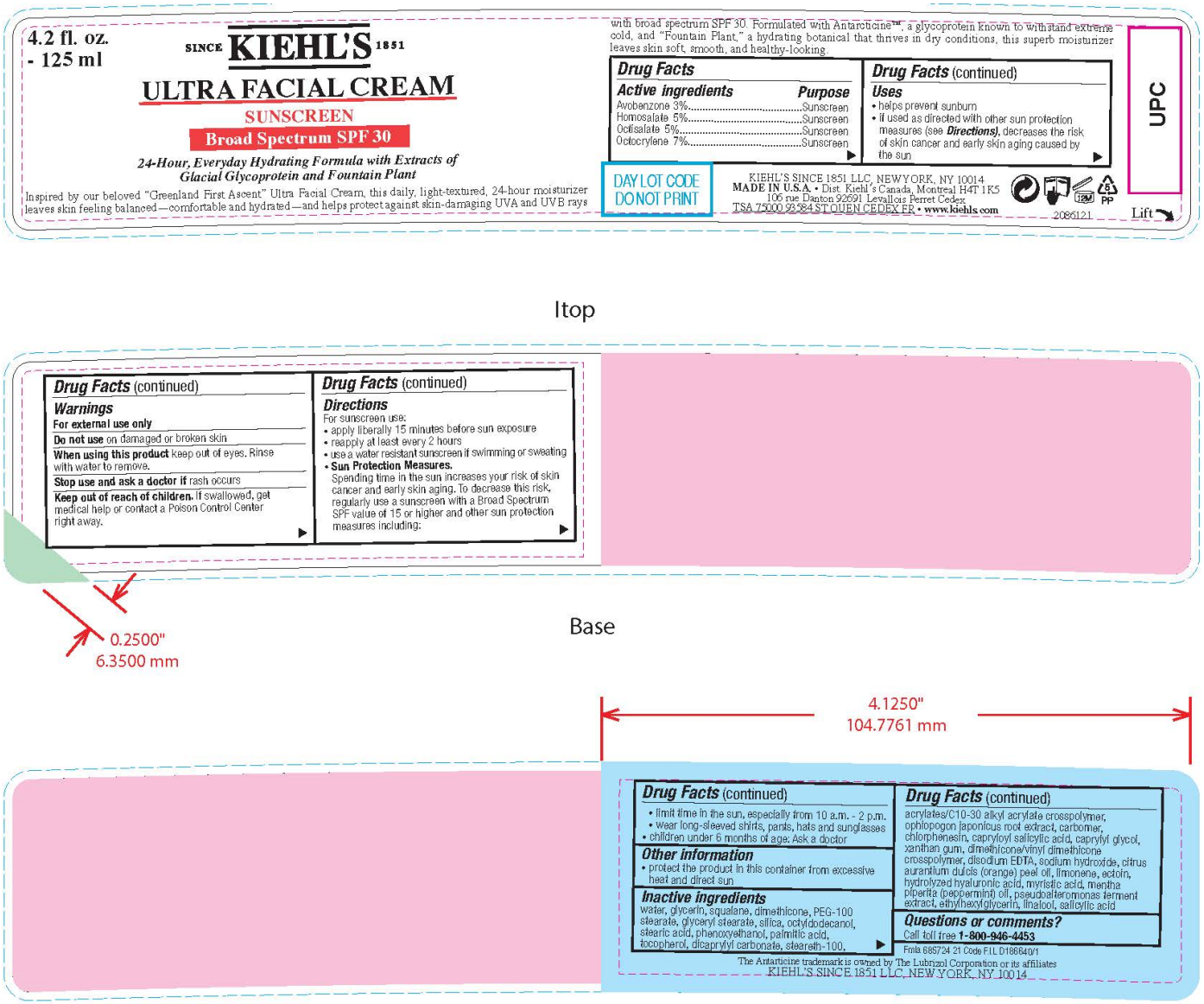
acrylates/C10-30 alkyl acrylate crosspolymer, ophiopogon japonicus root extract, carbomer, chlorphenesin, caprylyoyl salicylic acid, caprylyl glycol, xanthan gum, dimethicone/vinyl dimethicone crosspolymer, disodium EDTA, sodium hydroxide, citrus aurantium dulcis (orange) peel oil, limonene, ectoin, hydrolyzed hyaluronic acid, myristic acid, mentha piperita (peppermint) oil, pseudoalteromonas ferment extract, ethylhexylglycerin, linalool, salicylic acid

Questions or comments?

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

Fmla 685724 21 Code FIL D186640/1

KIEHL'S SINCE 1851 LLC, NEW YORK, NY 10014



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:49967-099
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)	
SQUALANE (UNII: GW89575KF9)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PEG-100 STEARATE (UNII: YD01N1999R)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PALMITIC ACID (UNII: 2V16EO95H1)	
TOCOPHEROL (UNII: R0ZB2556P8)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
STEARETH-100 (UNII: 4OH5W9UM87)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-099-01	50 mL in 1 JAR; Type 0: Not a Combination Product	11/01/2015	
2	NDC:49967-099-02	125 mL in 1 JAR; Type 0: Not a Combination Product	11/01/2015	
3	NDC:49967-099-03	3 mL in 1 PACKET; Type 0: Not a Combination Product	11/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/01/2015	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-099)

Revised: 12/2023 L'Oreal USA Products Inc