LA ROCHE POSAY LABORATOIRE DERMATOLOGIQUE ANTHELIOS GLOW DAILY GLOW SUNSCREEN BRONZE CELL OX SHIELD BROAD SPECTRUM SPF 35- avobenzone, homosalate, octisalate and octocrylene liquid
L'Oreal USA Products Inc

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 13%

Octisalate 5%

Octocrylene 6%

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 generously 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
- store at 20ºC - 25ºC (68ºF - 77ºF)

Inactive ingredients

water, glycerin, dicaprylyl carbonate, alcohol denat., C12-22 alkyl acrylate/ hydroxyethylacrylate copolymer, dimethicone, silica, synthetic fluorphlogopite, zea mays (corn) starch, mica, iron oxides, phenoxyethanol, titanium dioxide, dimethicone/PEG-10/15 crosspolymer, hydroxyacetophenone, pentaerythrityl tetra-di-t-butyl hydroxyhydrocinnamate, acrylamide/sodium acryloyldimethyltaurate copolymer, xanthan gum, arachidyl alcohol, chlorphenesin, caprylyl glycol, isohexadecane, behenyl alcohol, tocopherol, pentylene glycol, tetrasodium glutamate diacetate, diethylhexyl syringylidenemalonate, trisodium ethylenediamine disuccinate, arachidyl glucoside, citric acid, polysorbate 80, ammonium polyacryloyldimethyl taurate, cassia alata leaf extract, maltodextrin, sorbitan oleate, dipropylene glycol, glycine soja (soybean) oil, caprylic/capric triglyceride, tin oxide, disodium stearoyl glutamate, sodium citrate, aluminum hydroxide

Questions or comments?

1-888-LRP-LAB0 1-888-577-5226

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

ANTHELIOS GLOW BRONZE

LA ROCHE-POSAY
LABORATOIRE DERMATOLOGIQUE

NEW

BROAD SPECTRUM SPF 35

35

ANTHELIOS GLOW
DAILY GLOW SUNSCREEN

INSTANT GLOWING, RADIANT SKIN
Helps protect against sun damage*
Primes skin for all day brighter look

BRONZE

CELL-OX SHIELD®
UVA/UVB Protection + Antioxidants

Fragrance-free
Suitable for sensitive skin

*when used as directed with other sun protection measures

1.35 FL.OZ. - 40 ml

Drug Facts (continued)

ammonium polyacryloyldimethyl taurate, cassia alata leaf extract, maltodextrin, sorbitan oleate, dipropylene glycol, glycine soja (soybean) oil, caprylic/capric triglyceride, tin oxide, disodium stearoyl glutamate, sodium citrate, aluminum hydroxide

Questions or comments?

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2067530 14 Code F.LL. V70065118/1

A PIONEER IN SUN PROTECTION RESEARCH FOR OVER 30 YEARS.

Every day, dermatologists observe skin damage caused by UVA and UVB rays.

UVB rays mainly cause **Burning.** They cause tanning and are mainly responsible for sunburn. SPF or "Sun Protection Factor" is the degree of protection a sunscreen offers against UVB rays.

UVA rays mainly cause Skin **Aging** or Sun Allergies. UVA rays directly contribute to skin aging (wrinkles, sagging, dark spots) and are the #1 cause of sun intolerances (allergies).

Both UVA and UVB rays have been proven to cause damage to skin cells, including DNA, and can weaken the immune system. This damage can potentially lead to the development of skin cancer.

La Roche-Posay is dedicated to providing photostable, broad-spectrum protection through the use of **CELL-OX SHIELD**, a synergistic combination of UVA/UVB filters combined with antioxidants.

21
PAP
12M

FSC Mix Packaging FSC™ C095159

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ANTHELIOS GLOW 35 DAILY GLOW SUNSCREEN

Package:	Skin Type: All Texture: Light Lotion Application: Face

PROPERTIES.

1. Designed to Instantly Illuminate the Skin and Create a Sun-Kissed Glow. This lightweight lotion leaves a radiant finish and has a subtle, flexible tint ideal for daily use. It helps protect against the effects of daily sun exposure. Blends seamlessly into the skin and provides hydration.

2. Formulated for Adaptable and Multi-Purpose Use. Designed for use with and without makeup for "glow from within" skin. Use alone to instantly illuminate skin, or use as a primer to enhance makeup finish. With consistent daily usage, improves the skin's natural radiance.

3. CELL-OX SHIELD TECHNOLOGY UVA/UVB Protection + Antioxidants

Broad Spectrum UV Filters Combines photostable UVA/UVB filters chosen for their synergistic effect to deliver broad-spectrum protection.

Powerful Antioxidant Protection With Senna Alata, a tropical leaf extract known to defend skin cells[1] against damaging free radicals caused by the sun that can accelerate skin aging.

[1] upper layers of skin

FRAGRANCE-FREE
NON-COMEDOGENIC
ALLERGY TESTED
SUITABLE FOR SENSITIVE SKIN
DERMATOLOGIST TESTED FOR SAFETY

Made in USA of US and/or Imported Ingredients
Distribution reserved to La Roche-Posay agreed distributors
L'Oréal USA S/D, Inc., Dist.,
10 Hudson Yards,
New York, NY 10001.
3612624669005

La Roche-Posay
Laboratoire Dermatologique
CAI 86270 La Roche-Posay, France
TSA 75000 93584 ST OUEN CEDEX FR
www.laroche-posay.us

Drug Facts

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Octisalate 5%	Sunscreen
Octocrylene 6%	Sunscreen

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LA ROCHE POSAY LABORATOIRE DERMATOLOGIQUE ANTHELIOS GLOW DAILY GLOW SUNSCREEN BRONZE CELL OX SHIELD BROAD SPECTRUM SPF 35

avobenzone, homosalate, octisalate and octocrylene liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82046-768
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	130 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	60 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
ALCOHOL (UNII: 3K9958V90M)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SILICA (UNII: ETJ7Z6XBU4)	
ZEA MAYS (CORN) STARCH (UNII: O8232NY3SJ)	
MICA (UNII: V8A1AW0880)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
DIMETHICONE/PEG-10/15 CROSSPOLYMER (UNII: 21AS8B1BSS)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
PENTAERYTHRITYL TETRA-DI-T-BUTYL HYDROXYHYDROCINNAMATE (UNII: 255PIF62MS)	
ACRYLAMIDE/SODIUM ACRYLOYLDIMETHYLTAURATE COPOLYMER (120000 MPA.S AT 1%) (UNII: 5F4963KLHS)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ARACHIDYL ALCOHOL (UNII: 1QR1QRA9BU)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
BEHENYL ALCOHOL (UNII: 9G1OE216XY)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
ARACHIDYL GLUCOSIDE (UNII: 6JVW35JOOJ)	
CITRIC ACID (UNII: 2968PHW8QP)	

POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (UNII: F01RIY4371)	
SENNA ALATA LEAF (UNII: 4BXR6YZN92)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SORBITAN OLEATE (UNII: 06XEA2VD56)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
SOYBEAN OIL (UNII: 241ATL177A)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
TIN OXIDE (UNII: KM7N50LOS6)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82046-768-01	1 in 1 CARTON	04/28/2026	
1		40 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/28/2026	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'OREAL USA PRODUCTS, INC.		624244349	manufacture(82046-768)

Establishment

Name	Address	ID/FEI	Business Operations
Unette Corporation		011401882	pack(82046-768)

Revised: 4/2026 L'Oreal USA Products Inc