

**LA ROCHE POSAY NEW 50 BODY AND FACE SPF 50 MINERAL ANTHELIOS
GENTLE 100 PERCENT MINERAL UV FILTERS SOFT FINISH WITH CELL OX
SHIELD- titanium dioxide and zinc oxide lotion
L'Oreal USA Products Inc**

Drug Facts

Drug Facts

Titanium dioxide 5%

Zinc Oxide 15%

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:
 - after 80 minutes of swimming or sweating
 - immediately after towel drying
 - at least every 2 hours
- 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, C12-15 alkyl benzoate, isohexadecane, isononyl isononanoate, dicaprylyl ether, PEG-30 dipolyhydroxystearate, polyglyceryl-4 isostearate, dicaprylyl carbonate, ethylene/acrylic acid copolymer, triethylhexanoin, silica, poly C10-30 alkyl acrylate, citric acid, phenoxyethanol, sodium chloride, caprylyl glycol, diethylhexyl syringylidenemalonate, triethoxycaprylylsilane, aluminum hydroxide, stearic acid, aluminum stearate, chlorphenesin, disteardimonium hectorite, alumina, polyhydroxystearic acid, tocopherol, p-ansic acid, capryloyl salicylic acid, xanthan gum, disodium EDTA, propylene carbonate, caprylic/capric triglyceride, cassia alata leaf extract, maltodextrin

Questions or comments?

1-888-LRP-LABO 1-888-577-5226

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

text MUST NOT run over dashed line

LA ROCHE-POSAY
LABORATOIRE DERMATOLOGIQUE

NEW

50

Body & Face

BROAD SPECTRUM
SPF 50

105 mm

Mineral

ANTHELIOS SUNSCREEN - GENTLE LOTION

100% mineral UV filters
Soft finish lotion

with CELL-OX SHIELD®
UVA/UVB Protection + Antioxidants

Tested on Sensitive Skin. Fragrance-free
WATER RESISTANT (80 MINUTES)

4.0 FL.OZ. - 120 ml

INNOVATION

ANTHELIOS 50 Sunscreen - Gentle Lotion is ideal for sensitive skin and for those who want a sunscreen without chemical UV filters.

Properties:

1. Lightweight, soothing lotion

This gentle, oil-free sunscreen has a smooth application and blends easily into skin. Imparts a soft, powdery finish.

2. CELL-OX SHIELD® technology

100% Mineral UV Filters
UVA/UVB protection + antioxidants
Zinc Oxide and Titanium Dioxide deliver broad spectrum protection with antioxidants, Senna Alata and Vitamin E, to help neutralize free radicals.

ALLERGY TESTED. TESTED ON SENSITIVE SKIN
DERMATOLOGIST TESTED FOR SAFETY.
OIL-FREE. PARABEN-FREE. FRAGRANCE-FREE.
NON-COMEDOGENIC. WATER RESISTANT (80 MINUTES)

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1 upper layers of skin.

693261 24
Code F.I.L.: D223841/1



C/H/PE

0931618

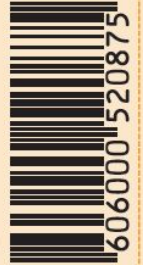
Made in USA with
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La Roche-Posay LLC,
10 Hudson Yards, New York,
NY 10001

La Roche-Posay Laboratoire
Dermatologique
86270 La Roche-Posay, France

TSA 75000 9384 ST OLEN CEDEX FR
www.laroche-posay.us



3 606000 520875

DECORATIVE AREA

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LABORATOIRE DERMATOLOGIQUE

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100% mineral UV filters
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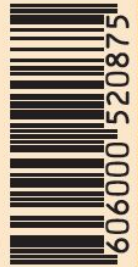
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LA ROCHE POSAY NEW 50 BODY AND FACE SPF 50 MINERAL ANTHELIOS GENTLE 100 PERCENT MINERAL UV FILTERS SOFT FINISH WITH CELL OX SHIELD

titanium dioxide and zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-091
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	50 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	150 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ISOHEXADECANE (UNII: 918X1OUF1E)	

ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)
DICAPRYLYL ETHER (UNII: 77JZM5516Z)
PEG-30 DIPOLYHYDROXYSTEARATE (UNII: 9713Q0S7FO)
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)
TRIETHYLHEXANOIN (UNII: 7K3W1BIU6K)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
CAPRYLYL GLYCOL (UNII: 00YIU5438U)
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
STEARIC ACID (UNII: 4ELV7Z65AP)
ALUMINUM STEARATE (UNII: U6XF9NP8HM)
CHLORPHENESIN (UNII: I670DAL4SZ)
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)
ALUMINUM OXIDE (UNII: LMI26O6933)
TOCOPHEROL (UNII: R0ZB2556P8)
P-ANISIC ACID (UNII: 4SB6Y7DMM3)
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)
XANTHAN GUM (UNII: TTV12P4NEE)
EDETATE DISODIUM (UNII: 7FLD91C86K)
PROPYLENE CARBONATE (UNII: 8D08K3S51E)
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)
SENNALATA LEAF (UNII: 4BXR6YZN92)
MALTODEXTRIN (UNII: 7CVR7L4A2D)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-091-01	120 mL in 1 TUBE; Type 0: Not a Combination Product	12/15/2018	
2	NDC:49967-091-02	90 mL in 1 TUBE; Type 0: Not a Combination Product	12/15/2018	
3	NDC:49967-091-03	5 mL in 1 PACKET; Type 0: Not a Combination Product	12/15/2018	
4	NDC:49967-091-04	5 mL in 1 TUBE; Type 0: Not a Combination Product	12/15/2018	
5	NDC:49967-091-05	150 mL in 1 TUBE; Type 0: Not a Combination Product	12/15/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/15/2018	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'OREAL USA, INC.		185931458	manufacture(49967-091)

Revised: 1/2024

L'Oreal USA Products Inc