LA ROCHE POSAY LABORATOIRE DERMATOLOGIQUE ANTHELIOS HA MINERAL BROAD SPECTRUM SPF 30 DAILY MOISTURIZING SUNSCREEN- titanium dioxide and zinc oxide cream
L'Oreal USA Products Inc

Drug Facts

Active ingredients

Titanium dioxide 5.5%

Zinc oxide 10%

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 after 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, glycerin, isohexadecane, isononyl isononanoate, dicaprylyl ether, PEG-30 dipolyhydroxystearate, c12-15 alkyl benzoate, polyglyceryl-4 isostearate, dimethicone, caprylyl methicone, ethylene/acrylic acid copolymer, caprylic/capric triglyceride, triethylhexanoin, silica, poly c10-30 alkyl acrylate, citric acid, phenoxyethanol, sodium chloride, caprylyl glycol, panthenol, diethylhexyl syringylidenemalonate, aluminum hydroxide, stearic acid, aluminum stearate, sodium hyaluronate, triethoxycaprylylsilane, chlorphenesin, disteardimonium hectorite, alumina, polyhydroxystearic acid, trisodium ethylenediamine disuccinate, tocopherol, p-anisic acid, capryloyl salicylic acid, xanthan gum, propylene carbonate, cassia alata leaf extract, maltodextrin

Questions or comments?

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

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

ANTHELIOS HA MINERAL

DAILY 30

SUNSCREEN

LA ROCHE-POSAY
LABORATOIRE DERMATOLOGIQUE

BROAD SPECTRUM SPF30

ANTHELIOS HA MINERAL

DAILY MOISTURIZING CREAM MINERAL SUNSCREEN
+ HYALURONIC ACID

Provides comfort and hydration
Helps protect against early skin aging caused by the sun*

Advanced Technology
with CELL-OX SHIELD®
UVA/UVB + ANTIOXIDANTS

Non-comedogenic
Fragrance-free. Paraben-free
Tested on sensitive skin

*when used as directed with other sun protection measures

1.7FL. OZ - 50 ml

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.

12M PAP MIX Packaging FSC FSC® B004460

Made in USA of US and Imported Ingredients.
1752816.
Distribution reserved to La Roche-Posay agreed distributors.
La Roche-Posay LLC, 10 Hudson Yards, New York, NY 10001.
La Roche-Posay Laboratoire Dermatologique
CAI 86270 La Roche-Posay, France
TSA 75000 93584 ST OUEN CEDEX FR
www.laroche-posay.us

3 606000 546028

ANTHELIOS HA MINERAL 30

Package:

Skin Type: All, especially dry and sensitive

Texture: Cream

Application: Face

PROPERTIES.

1. For all skin types, especially dry and sensitive.
Ideal for dry, sensitive skin and those who are concerned with premature skin aging and want a daily moisturizing sunscreen with 100% mineral UV filters.

2. Hydrates and helps reduce signs of premature skin aging caused by the sun*
A gentle, daily moisturizer with mineral UV filters and Hyaluronic Acid to help protect and hydrate skin. This non-greasy cream provides a comfortable feel and hydration, and helps protect against early skin aging caused by the sun*. Skin is left feeling soft, soothed, and hydrated.

3. CELL-OX SHIELD TECHNOLOGY 100% Mineral UV Filters.
Zinc Oxide and Titanium Dioxide deliver broad-spectrum protection.

Powerful Antioxidant Protection.
With Senna Alata, a tropical leaf extract, helps fight skin damaging free radicals caused by the sun that can accelerate skin aging.

RESULTS. Decreases the risk of skin cancer and early skin aging caused by the sun if used as directed with other sun protection measures (see **Directions**).

** When used as directed with other sun protection measures.*

OIL-FREE
FRAGRANCE-FREE
PARABEN-FREE
ALLERGY TESTED
NON-COMEDOGENIC
DERMATOLOGIST TESTED
FOR SAFETY

Drug Facts

Active ingredients	Purpose
Titanium dioxide 5.5%	Sunscreen
Zinc oxide 10%	Sunscreen

Uses
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Directions
For sunscreen use:
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- 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.
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693273 22 Code F.I.L.: D256875/1

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DAILY 30

SUNSCREEN

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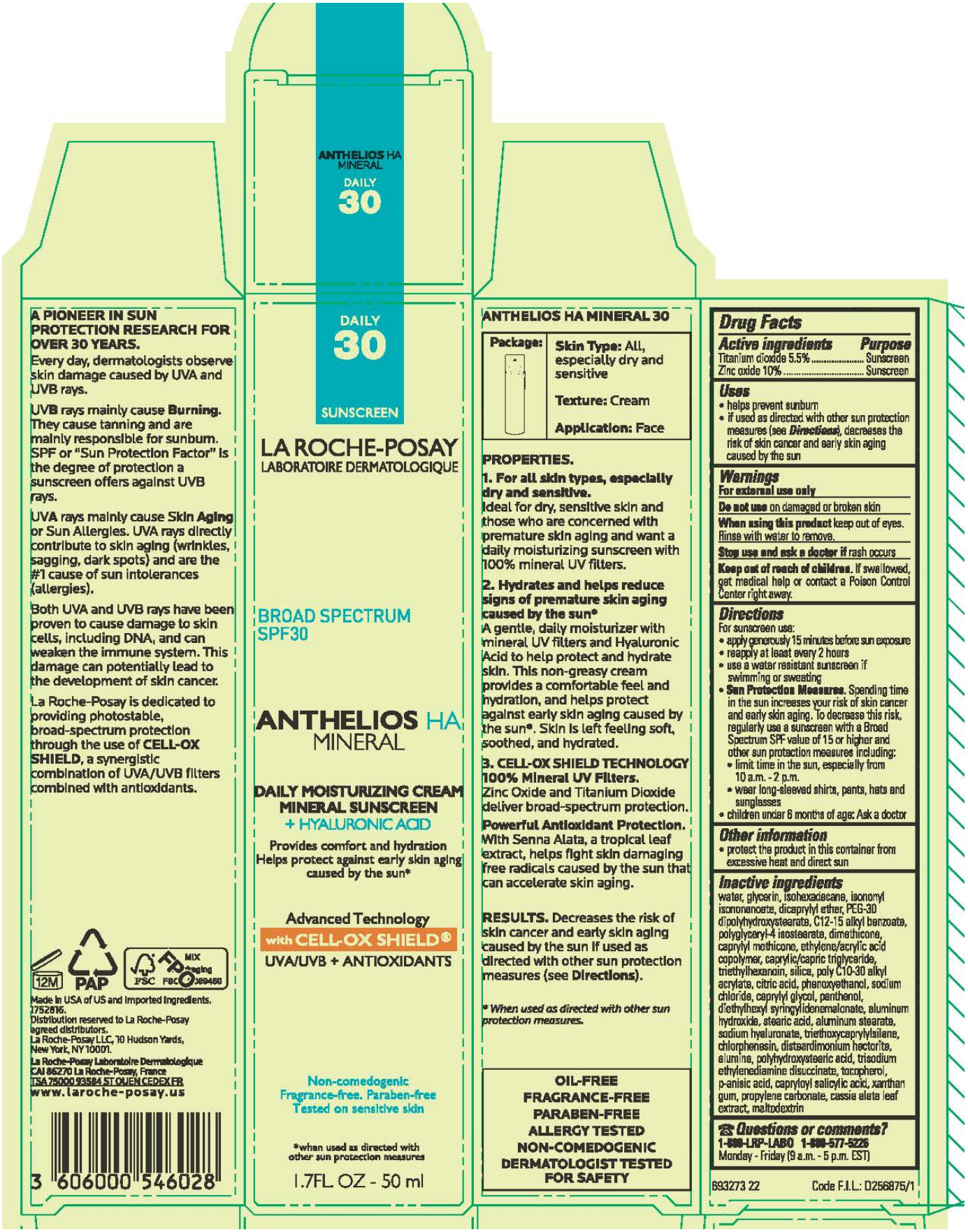

LA ROCHE POSAY LABORATOIRE DERMATOLOGIQUE ANTHELIOS HA MINERAL BROAD SPECTRUM SPF 30 DAILY MOISTURIZING

SUNSCREEN

titanium dioxide and zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-460
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	55 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	100 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
DICAPRYLYL ETHER (UNII: 77JZM5516Z)	
PEG-30 DIPOLYHYDROXYSTEARATE (UNII: 9713Q0S7FO)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
ACRYLIC ACID/ETHYLENE COPOLYMER (600 MPA.S) (UNII: 1PEZ3NLY6I)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
TRIETHYLHEXANOIN (UNII: 7K3W1BIU6K)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PANTHENOL (UNII: WV9CM0O67Z)	
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
ALUMINUM STEARATE (UNII: U6XF9NP8HM)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
TOCOPHEROL (UNII: R0ZB2556P8)	
P-ANISIC ACID (UNII: 4SB6Y7DMM3)	

CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
SENNA ALATA LEAF (UNII: 4BXR6YZN92)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-460-01	1 in 1 CARTON	12/01/2020	
1		50 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49967-460-02	1 in 1 CARTON	12/01/2020	
2		5 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:49967-460-03	2 mL in 1 PACKET; Type 0: Not a Combination Product	12/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2020	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, INC.		185931458	manufacture(49967-460)

Revised: 1/2024 L'Oreal USA Products Inc