

**CERAVE DEVELOPED WITH DERMATOLOGISTS HYDRATING SUNSCREEN SPF 50  
FACE LIGHTWEIGHT, NON-GREASY FEEL 3 ESSENTIAL CERAMIDES  
NIACINAMIDE MINERAL- titanium dioxide and zinc oxide lotion  
L'Oreal USA Products Inc**

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**Drug Facts**

**Active ingredients**

Titanium dioxide 9%

Zinc oxide 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

**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.-2p.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, C12-15 alkyl benzoate, dimethicone, isododecane, styrene/acrylates copolymer, glyceryl stearate, butyloctyl salicylate, dicaprylyl carbonate, propanediol, stearic acid, aluminum hydroxide, PEG-100 stearate, sorbitan stearate, niacinamide, PEG-8 laurate, ceramide NP, ceramide AP, ceramide EOP, sobitan isostearate, carbomer, cetearyl alcohol, cetareth-20, triethoxycaprylylsilane, dimethiconol, sodium citrate, sodium lauroyl lactylate, sodium dodecylbenzenesulfonate, myristic acid, sodium hyaluronate, cholesterol, palmitic acid, phenoxyethanol, chlorphenesin, tocopherol, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, caprylyl glycol, citric acid, panthenol, xanthan gum, phytosphingosine, polyhydroxystearic acid, polysorbate 60, ethylhexylglycerin

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### Questions or comments?

Toll-free number 1-888-768-2915

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CeraVe<sup>®</sup>

DEVELOPED WITH DERMATOLOGISTS

MINERAL SUNSCREEN

BROAD SPECTRUM SPF 50  
Hydrating Sunscreen

50  
FACE



Cerave<sup>®</sup> LLC,  
New York, NY 10001  
Made in USA of US and/or  
Imported Ingredients  
www.cerave.com

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NEW

CeraVe<sup>®</sup>

DEVELOPED WITH DERMATOLOGISTS

Hydrating  
Sunscreen

BROAD SPECTRUM SPF 50

50  
FACE

Lightweight, non-greasy feel

3 essential ceramides & niacinamide

MINERAL SUNSCREEN

2.5 FL OZ (75 ml)



Developed with dermatologists, its formula – with 3 essential ceramides – helps restore and maintain the protective skin barrier.



Broad Spectrum Protection

Helps protect your skin against harmful UVA and UVB rays



100% Mineral Sun Filters

Reflects and blocks the sun's damaging rays



Lightweight & Oil Free

Won't clog pores



Niacinamide

Helps calm skin

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BROAD SPECTRUM SPF 50  
Hydrating Sunscreen  
50 FACE



CeraVe® LLC.  
New York, NY 10001  
Made in USA of US and/or  
Imported Ingredients  
www.cerave.com

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NEW

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titanium dioxide and zinc oxide lotion

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-959
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	90 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	70 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ALKYL (C12-15) BENZOATE</b> (UNII: A9EJ3J61HQ)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>ISODODECANE</b> (UNII: A8289P68Y2)	
<b>STYRENE/ACRYLAMIDE COPOLYMER (500000 MW)</b> (UNII: 5Z4DPO246A)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>BUTYLOCTYL SALICYLATE</b> (UNII: 2EH13UN8D3)	
<b>DICAPRYLYL CARBONATE</b> (UNII: 609A3V1SUA)	
<b>PROPANEDIOL</b> (UNII: 5965N8W85T)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>ALUMINUM HYDROXIDE</b> (UNII: 5QB0T2IUN0)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>SORBITAN MONOSTEARATE</b> (UNII: NVZ4I0H58X)	
<b>NIACINAMIDE</b> (UNII: 25X51I8RD4)	
<b>PEG-8 LAURATE</b> (UNII: 762O8IWA10)	
<b>CERAMIDE NP</b> (UNII: 4370DF050B)	
<b>CERAMIDE AP</b> (UNII: F1X8L2B00J)	
<b>CERAMIDE 1</b> (UNII: 5THT33P7X7)	
<b>SORBITAN ISOSTEARATE</b> (UNII: 01S2G2C1E4)	
<b>CARBOXYPOLYMETHYLENE</b> (UNII: 0A5MM307FC)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>POLYOXYL 20 CETOSTEARYL ETHER</b> (UNII: YRC528SWUY)	
<b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SODIUM LAUROYL LACTYLATE</b> (UNII: 7243K85WFO)	
<b>SODIUM DODECYLBENZENESULFONATE</b> (UNII: 554127163Y)	
<b>MYRISTIC ACID</b> (UNII: 0I3V7S25AW)	
<b>HYALURONATE SODIUM</b> (UNII: YSE9PPT4TH)	
<b>CHOLESTEROL</b> (UNII: 97C5T2UQ7J)	
<b>PALMITIC ACID</b> (UNII: 2V16EO95H1)	
<b>phenoxyethanol</b> (UNII: HIE492ZZ3T)	
<b>CHLORPHENESIN</b> (UNII: I670DAL4SZ)	
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)	
<b>HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%)</b> (UNII: 86FQE96TZ4)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>CITRIC ACID ACETATE</b> (UNII: DSO12WL7AU)	
<b>PANTHENOL</b> (UNII: WV9CM0O67Z)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>PHYTOSPHINGOSINE</b> (UNII: GIN46U9Q2Q)	
<b>POLYHYDROXYSTEARIC ACID STEARATE</b> (UNII: 8KQ7I65XZE)	
<b>POLYSORBATE 60</b> (UNII: CAL22UVI4M)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-959-01	1 in 1 CARTON	02/01/2019	
1		75 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:49967-959-02	5 mL in 1 TUBE; Type 0: Not a Combination Product	02/01/2019	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/01/2019	

**Labeler** - L'Oreal USA Products Inc (002136794)

## Establishment

Name	Address	ID/FEI	Business Operations
L'OREAL USA PRODUCTS, INC.		624244349	manufacture(49967-959)

Revised: 12/2023

L'Oreal USA Products Inc