

**LANCOME PARIS TEINT IDOLE ULTRA WEAR BREATHABLE COVERAGE
FOUNDATION BROAD SPECTRUM SPF 25 SUNSCREEN- octinoxate and titanium
dioxide liquid
SICOS ET CIE**

Drug Facts

Active ingredients

Octinoxate 6.7%

Titanium dioxide 5.1%

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

Flammable until dry.

Do not use near fire, flame, or heat.

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:

- shake well
- 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, dimethicone, isododecane, alcohol denat., trimethylsiloxy silicate, butylene glycol, PEG-10 dimethicone, perlite, synthetic fluorophlogopite, moringa oleifera seed extract, polymnia sonchifolia root juice, calcium aluminum borosilicate, glycerin, dipentaerythrityl tetrahydroxystearate/tetraisostearate, alpha-glucan oligosaccharide, sodium hyaluronate, silica, silica silylate, HDI/trimethylol hexyllactone crosspolymer, cellulose, aluminum hydroxide, magnesium sulfate, nylon-12, disodium phosphate, disodium stearyl glutamate, isopropyl lauroyl sarcosinate, hydrogen dimethicone, citric acid, diisopropyl sebacate, bis-PEG/PPG-14/14 dimethicone, lactobacillus, maltodextrin, disteardimonium hectorite, BHT, tocopherol, phenoxyethanol; may contain: titanium dioxide, iron oxides

Questions or comments?

1-800-LANCOME

(1-800-526-2663)

Monday - Friday

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

Drug Facts (continued)

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6933800 V297134/4

LONG WEAR FOUNDATION
Breathable and buildable full coverage.
Natural matte finish.

Paraben-free
Non-comedogenic
Non-sensitizing
Tested under dermatological control for safety



RECYCLABLE GLASS BOTTLE MADE WITH 25% RECYCLED MATERIAL.

Anti Diversion



LANCÔME SUPPORTS FOREST MANAGEMENT THAT RESPECTS PEOPLE AND NATURE.

THIS PRODUCT CAN BE SOLD ONLY BY APPROVED DISTRIBUTORS
LANCÔME PARIS 14 RUE ROYALE 75008 PARIS LANCÔME
LUXURY PRODUCTS LLC 10 HUDSON YARDS, NEW YORK, NY
10001 - IMP. BY LANCÔME CANADA, MONTREAL, H4T 1K5 -
LONDON W6 0AZ - TS&T 600 9365451 QUEBEC Q1E 0E6

30 ml 1.0 FL. OZ.
MADE IN FRANCE
LANCÔME.COM

LANCÔME
PARIS

LANCÔME
PARIS



TEINT IDOLE
ULTRA WEAR

SUNSCREEN
BROAD SPECTRUM SPF 25
UP TO 24H WEAR FOUNDATION
BREATHABLE COVERAGE

COVERAGE
● ● ● ● ●
100W

TRANSFER-RESISTANT
30 ml - 1.0 FL. OZ.

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SV 3614273793025
GENCOD 80%

Drug Facts (continued)

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Anti Diversion



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LANCOME PARIS, 14 RUE ROYALE 75008 PARIS LANCOME LUXURY PRODUCTS LLC, 10 HUDSON YARDS, NEW YORK, NY 10001 - IMP. BY LANCOME CANADA, MONTREAL, H4T 1K5 - LONDON W6 8AZ - TSA760003565451 OULENCELEEE

30 ml 1.0 FL. OZ.

MADE IN FRANCE
LANCOME.COM

LANCOME PARIS

LANCÔME PARIS



TEINT IDOLE
ULTRA WEAR

SUNSCREEN
BROAD SPECTRUM SPF 25
UP TO 24H WEAR FOUNDATION
BREATHABLE COVERAGE

COVERAGE



100W

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 MIX Packaging PSC. Fsc® C200050

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30 ml 
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LANCOME.COM

LANCÔME

PARIS



**TEINT IDOLE
ULTRA WEAR**

**SUNSCREEN
BROAD SPECTRUM SPF 25**

UP TO 24H WEAR FOUNDATION
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SV 3614273793025
GENCOD 80%

C1 - Internal use

LANCOME PARIS TEINT IDOLE ULTRA WEAR BREATHABLE

COVERAGE FOUNDATION BROAD SPECTRUM SPF 25 SUNSCREEN

octinoxate and titanium dioxide liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51150-302
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	67 mg in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	51 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ISODODECANE (UNII: A8289P68Y2)	
ALCOHOL (UNII: 3K9958V90M)	
TRIMETHYLSILOXYSILICATE (M/Q 0.6-0.8) (UNII: 5041RX63GN)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
PERLITE (UNII: 0SG101ZGK9)	
MAGNESIUM POTASSIUM ALUMINOSILICATE FLUORIDE (UNII: YK3DC63Y5M)	
MORINGA OLEIFERA SEED (UNII: TIX5482832)	
SMALLANTHUS SONCHIFOLIUS ROOT JUICE (UNII: M9S7HX36CT)	
CALCIUM ALUMINUM BOROSILICATE (UNII: 3JRB8A35M0)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIPENTAERYTHRITYL TETRAHYDROXYSTEARATE/TETRAISOSTEARATE (UNII: 230K0823CE)	
.ALPHA.-GLUCAN OLIGOSACCHARIDE (UNII: S95658MI3W)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HEXAMETHYLENE DIISOCYANATE/TRIMETHYLOL HEXYLLACTONE CROSSPOLYMER (UNII: WB5K9Y35Y9)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
NYLON-12 (UNII: 446U8J075B)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
ISOPROPYL LAUROYL SARCOSINATE (UNII: LYR06W430J)	
HYDROGEN DIMETHICONE (13 CST) (UNII: 4QGR4P2YOI)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DIISOPROPYL SEBACATE (UNII: J8T3X564IH)	
BIS-PEG/PPG-14/14 DIMETHICONE (UNII: X2I70H0QJE)	
EMMENTAL LACTOBACILLUS (UNII: B253LSV04X)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	

BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51150-302-01	1 in 1 CARTON	12/12/2022	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:51150-302-02	100 mL in 1 TUBE; Type 0: Not a Combination Product	12/12/2022	
3	NDC:51150-302-03	1 mL in 1 PACKET; Type 0: Not a Combination Product	12/12/2022	01/22/2027
4	NDC:51150-302-04	0.4 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/12/2022	12/28/2026

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/12/2022	

Labeler - SICOS ET CIE (276993581)

Establishment			
Name	Address	ID/FEI	Business Operations
SICOS ET CIE		276993581	manufacture(51150-302) , pack(51150-302)

Establishment			
Name	Address	ID/FEI	Business Operations
BPS60		272259304	pack(51150-302)

Establishment			
Name	Address	ID/FEI	Business Operations
Socoplan		276221405	pack(51150-302)