LANCOME PARIS ABSOLUE MAKEUP BROAD SPECTRUM SPF 20 SUNSCREEN ABSOLUTE REPLENISHING MAKEUP FOR DRY AND VERY DRY SKIN-octinoxate cream L'Oreal USA Products Inc

Drug Facts

Active ingredient

Octinoxate 7.5%

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 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, cyclohexasiloxane, hydrogenated polyisobutene, glycerin, cyclopentasiloxane, polyglyceryl-4 isostearate, petrolatum, nylon-12, cetyl PEG/PPG-10/1 dimethicone, hexyl laurate, disteardimonium hectorite, pentylene glycol, C18-36 acid glycol ester, linalool, paraffin, glycyrrhiza glabra (licorice) extract, diphenyl dimethicone, padina pavonica extract, cellulose gum, aluminum hydroxide, magnesium sulfate, limonene, disodium stearoyl glutamate, tristearin, dioscorea villosa (wild yam) root extract, acrylates copolymer, hexyldecanol, fragrance, methylparaben, phenoxyethanol, butylparaben, diazolidinyl urea, tocopherol; may contain: mica, titanium dioxide, iron oxides

Questions or comments?

1-800-LANCOME (1800-526-2663) Monday - Friday (9 a.m. - 5 p.m. EST)





LANCOME PARIS ABSOLUE MAKEUP BROAD SPECTRUM SPF 20 SUNSCREEN ABSOLUTE REPLENISHING MAKEUP FOR DRY AND VERY DRY SKIN

octinoxate cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-596	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
HYDROGENATED POLYISOBUTENE (450 MW) (UNII: 7YR4ZFS62E)	
GLYCERIN (UNII: PDC6A3C0OX)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
polyglyceryl-4 isostearate (UNII: 820DPX33S7)	
petrolatum (UNII: 4T6H12BN9U)	
nylon-12 (UNII: 446U8J075B)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 4) (UNII: 8INO2K35FA)	
hexyl laurate (UNII: 4CG9F9W01Q)	
disteardimonium hectorite (UNII: X687XDK09L)	
pentylene glycol (UNII: 50C1307PZG)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
Paraffin (UNII: I9O0E3H2ZE)	
DIPHENYL DIMETHICONE (300 CST) (UNII: ZF5SBB12ZZ)	
PADINA PAVONICA (UNII: 177U06NHZI)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)	
aluminum hydroxide (UNII: 5QB0T2IUN0)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	

LIMONENE, (+)- (UNII: GFD7C86Q1W)	
disodium stearoyl glutamate (UNII: 45ASM2L11M)	
tristearin (UNII: P6OCJ2551R)	
DIOSCOREA VILLOSA TUBER (UNII: IWY3IWX2G8)	
LUVISET 360 (UNII: 05BG6GY6YK)	
hexyldecanol (UNII: 151Z7P1317)	
methylparaben (UNII: A2I8C7HI9T)	
phenoxyethanol (UNII: HIE492ZZ3T)	
butylparaben (UNII: 3QPI1U3FV8)	
diazolidinyl urea (UNII: H5RIZ3MPW4)	
tocopherol (UNII: R0ZB2556P8)	
mica (UNII: V8A1AW0880)	
titanium dioxide (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:49967-596- 01	1 in 1 CARTON	11/01/2007		
1		35 mL in 1 JAR; Type 0: Not a Combination Product			
2	NDC:49967-596- 02	10 mL in 1 JAR; Type 0: Not a Combination Product	11/01/2007		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/01/2007	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment				
Na me	Address	ID/FEI	Business Operations	
L'Oreal USA, Inc.		185931458	manufacture(49967-596)	

Revised: 6/2024 L'Oreal USA Products Inc