

KIEHL'S ACTIVATED SUN PROTECTOR ULTRA LIGHT SUNSCREEN SPF 50 FOR BODY SWEAT AND WATER RESISTANT- octocrylene, octisalate, avobenzone, ecamsule and titanium dioxide lotion
L'Oreal USA Products Inc

Drug Facts

SINCE KIEHL'S 1851

ACTIVATED SUN PROTECTOR™
ULTRA LIGHT SUNSCREEN

SPF 50+
Broad Spectrum Very High Protection

UVA

Lotion For Body

Dermatologist Tested UVA/UVB Protection

Sweat & Water Resistant

5.0 fl. oz. - 150 ml

INGREDIENTS: Aqua/Water, C12-15 Alkyl Benzoate, Octocrylene, Glycerin, Propylene Glycol, Ethylhexyl Salicylate, Isohexadecane, Butyl Methoxydibenzoylmethane, Titanium Dioxide, Nylon-12, Zea Mays Starch/Coen Starch, Alcohol Denat., Bis-Ethylhexyloxyphenol Methoxyphenyl Triazine, PEG-100 Stearate, Potassium Cetyl Phosphate, Glyceryl Stearate, Synthetic Wax, Stearic Acid, Triethanolamine, Phenoxyethanol, Dimethicone, Lycium Barbarum Fruit Extract, Caprylyl Glycol, Terephthalylidene Dicamphor Sulfonic Acid, Palmitic Acid, Aluminum Hydroxide, Xanthan Gum, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Disodium EDTA, Tocopherol, Hydrolyzed Triticum Monococcum Seed Extract. Fmla 858190 3 Code F.I.L. D4E533/1

DIRECTIONS: Over-exposure to the sun is dangerous. Keep babies and young children out of direct sunlight. Do not stay too long in the sun, even while using a sunscreen product because it does not provide you 100% protection. Apply the sunscreen product just before exposure. Re-apply frequently and generously to maintain protection, especially after swimming, perspiring or travelling. Avoid the eye area. In case of contact with eyes, rinse them immediately and thoroughly.

MODO DE EMPLEO: La sobreexposición al sol es peligrosa. Proteger a los bebés y a los niños pequeños de la luz solar directa. No permanecer bajo el sol durante periodos prolongados, incluso si se utiliza un producto

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UNCOATED AREA

KIEHL'S ACTIVATED SUN PROTECTOR ULTRA LIGHT SUNSCREEN SPF 50 FOR BODY SWEAT AND WATER RESISTANT

octocrylene, octisalate, avobenzone, ecamsule and titanium dioxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-986
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	70 mg in 1 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	35 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL

TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	40 mg in 1 mL
ECAMSULE (UNII: M94R1PM439) (ECAMSULE - UNII:M94R1PM439)	ECAMSULE	15 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
BEMOTRIZINOL (UNII: PWZ1720CBH)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TROLAMINE (UNII: 9O3K93S3TK)	
STARCH, CORN (UNII: O8232NY3SJ)	
WATER (UNII: 059QF0K00R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
XANTHAN GUM (UNII: TTV12P4NEE)	
LYCIUM BARBARUM FRUIT (UNII: 930626MWDL)	
NYLON-12 (UNII: 446U8J075B)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-986-01	150 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
export only		03/01/2013	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'OREAL USA, INC.		185931458	manufacture(49967-986)