KIEHLS SINCE 1851 ACTIVATED SUN PROTECTOR BROAD SPECTRUM SPF 50 SUNSCREEN WATER RESISTANT 80 MINUTES- avobenzone, homosalate, octisalate, octocrylene and oxybenzone lotion
L'Oreal USA Products Inc

Drug Facts

Active ingredient

Avobenzone 3%

Homosalate 10.72%

Octisalate 3.21%

Octocrylene 6%

Oxybenzone 3.86%

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:
  - after 80 minutes of 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, dimethicone, alcohol denat., styrene/acrylates copolymer, acrylates/dimethicone copolymer, phenoxyethanol, caprylyl glycol, sodium polyacrylate, silica, PEG-8 laurate, acrylates/C10-30 alkyl acrylate crosspolymer, tocopherol, menthyl lactate, disodium EDTA

Questions or comments?

Call toll free **1-800-946-4453**

SINCE KIEHL'S 1851

ACTIVATED SUN PROTECTOR™

SUNSCREEN

Broad Spectrum SPF 50

Water-Light Lotion for Face and Body

Cooling and Refreshing Weightless Formula

UVA/UVB Protection

Water Resistant (80 minutes)

Our refreshing, virtually weightless sunscreen lotion is formulated with an advanced sun-filter system that provides broad spectrum protection, and transforms to a unique, water-like texture on skin. With antioxidant Vitamin E, our formula helps to reinforce the skin's natural defense and neutralize skin-damaging free radicals. Fragrance-Free. Paraben-Free. Non-Comedogenic. Safe for Sensitive Skin.

5.0 fl. oz. – 150 ml

Drug Facts

Active Ingredients	Purpose
Avobenzone 3%	Sunscreen
Homosalate 10.72%	Sunscreen
Octisalate 3.21%	Sunscreen
Octocrylene 6%	Sunscreen
Oxybenzone 3.66%	Sunscreen

Uses

- helps prevent sunburn
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Questions or comments?

Call toll free **1-800-848-4453**

Fmla 858430 F.L.L. Code D175968/1

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2020309

BARCODE FPO

KIEHLS SINCE 1851 ACTIVATED SUN PROTECTOR BROAD SPECTRUM SPF 50 SUNSCREEN WATER RESISTANT 80 MINUTES

avobenzone, homosalate, octisalate, octocrylene and oxybenzone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-047
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	107.2 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	32.1 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	60 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	38.6 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALCOHOL (UNII: 3K9958V90M)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
PEG-8 LAURATE (UNII: 762O8IWA10)	
TOCOPHEROL (UNII: R0ZB2556P8)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-047-01	150 mL in 1 TUBE; Type 0: Not a Combination Product	02/15/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/15/2015	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-047)

Revised: 12/2023 L'Oreal USA Products Inc