

**DAY L- avobenzene octisalate octinoxate lotion**  
**Kamins Dermatologics Inc.**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**CONSUMER LABELING**

Active ingredients:

AVOBENZONE 2.5 %

OCTINOXATE 7.5 %

OCTISALATE 4.0 %

Purpose: Sunscreen

Keep out of the reach of children.

A LIGHT, SILKY LOTION THAT DIFFUSES A BLEND OF MOISTURIZERS, ANTI-OXYDANT VITAMINS INTO THE SKIN TO ENHANCE DAY LONG COMFORT AND HYDRATION. THE SKIN'S OUTER LAYERS ARE COVERED WITH A PROTECTIVE SHIELD, PRODUCING A HEALTHY, LUMINESCENT GLOW.

Uses

- Helps prevent sunburn.
- Higher SPF gives more sunburn protection.
- Provides moderate protection against sunburn.

Warnings

For external use only.

When using this product, keep out of eyes. If contact occurs, rinse abundantly with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts.

If swallowed, seek medical assistance or contact a Poison Control Center immediately.

Directions:

Apply liberally and evenly each morning to cleansed face and neck, before sun exposure and as needed.

Children under 6 months of age: ask a doctor.

Inactive ingredients:

aqua/water/eau, acer saccharum (maple isolate), butylene glycol, hydrolyzed collagen, hydrolyzed elastin, octocrylene, cetyl alcohol, glyceryl stearate SE, cetearth-12, cyclotetrasiloxane, cyclopentasiloxane, PEG/PPG-18/18 dimethicone, cococaprylate/caprate, polysorbate-80, carbomer, tocopheryl acetate, retinyl palmitate, sodium hydroxide, phenoxyethanol, caprylyl glycol, sorbic acid, parfum/fragrance

B. KAMINS

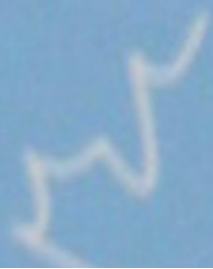
DAY LOTION SPF15

50mL / 1.7 fl.oz

DIN 02248016



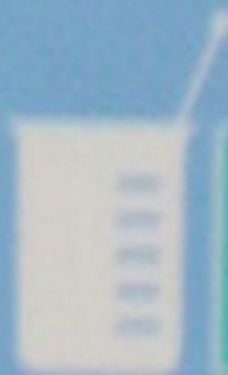
B. KAMINS  
laboratories



Day Lotion  
SPF 15  
Lotion

Crème  
de jour  
FPS 15

moisturize  
hydratant



50 ml / 1.7 fl. oz  
DIN 02248016

# DAY L

avobenzone octisalate octinoxate lotion

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63550-122
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	25 mg in 1 mL
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	40 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>SORBIC ACID</b> (UNII: X045WJ989B)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ACER SACCHARUM SAP</b> (UNII: 75UOH57984)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>HYDROLYZED ELASTIN, BOVINE, ALKALINE (1000 MW)</b> (UNII: ZR28QKN0WT)	
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6VM)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>CETEARETH-12</b> (UNII: 7V4MR24V5P)	
<b>CYCLOMETHICONE 4</b> (UNII: CZ227117JE)	
<b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)	
<b>POLYETHYLENE GLYCOL 1000</b> (UNII: U076Q6Q621)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>COCO-CAPRYLATE</b> (UNII: 4828G836N6)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>CARBOMER 934</b> (UNII: Z135WT9208)	
<b>ALPHA-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>VITAMIN A PALMITATE</b> (UNII: 1D1K0N0VVC)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	

## Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63550-122-03	50 mL in 1 BOX; Type 0: Not a Combination Product	03/01/2004	

B. KAMINS  
laboratories

Bio-Maple™

 Day Lotion

An ultra-lightweight oil-free  
lotion with vitamins A and E,  
that leaves the skin  
hydrated and soft looking  
all day long

B. KAMINS  
laboratories

Day Lotion

ESSENTIALS • ESSENTIELS

50 ml / 1.7 fl. oz



## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/01/2004	

**Labeler** - Kamins Dermatologics Inc. (254050784)

**Registrant** - Kamins Dermatologics Inc. (254050784)

## Establishment

Name	Address	ID/FEI	Business Operations
Odan Laboratories Ltd		208585604	manufacture(63550-122)

Revised: 12/2022

Kamins Dermatologics Inc.