

**MOISTURIZER MAPLE DAY SPF15- moisturizer maple day spf15 cream
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.

Moisturizer Maple day Cream SPF 15

Purpose section: Sunscreen

A super emollient hydrating day cream with sunscreen to nourish deeply dry and dehydrated skin and supplement the skin's diminished natural oils.

Uses:

Helps prevent sunburn

Warnings

Skin Cancer/Skin Aging Alert

Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to help sunburn, not skin cancer or early skin aging.

For external use only.

Do not use on damaged or broken skin.

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

Stop use and ask a doctor if rash occurs.

Keep out of the reach and sight of children.

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

Directions

Apply liberally and evenly, each morning to cleansed face and neck, 15 minutes before sun exposure and as needed

Reapply at least every 2 hours

Use a water-resistant sunscreen if swimming or sweating

Children under 6 months of ages: ask a doctor.

Storage:

Protect the product in the container from excessive heat and direct sun.

Store between 15-30 °C

Principal display Panel:

B. KAMINS

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laboratories

Bio-Maple™

 **Moisturizer
Maple Day
Cream SPF 15**

A super emollient hydrating day
cream with sunscreen to nourish
deeply dry and dehydrated skin
and supplement the skin's
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Moisturizer Maple Day Cream SPF 15

HYDRATION ● HYDRATATION

DIN 02244222 | 50 ml / 1.7 fl.oz



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50mL / 1.7 fl.oz

MOISTURIZER MAPLE DAY SPF15

moisturizer maple day spf15 cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63550-108
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZ ONE - UNII:95OOS7VE0Y)	OXYBENZONE	45 mg in 1 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM LACTATE (UNII: TU7HW0W0QT)	50 mg in 1 g
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	1.7 mg in 1 g
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	4 mg in 1 g
SQUALANE (UNII: GW89575KF9)	35 mg in 1 g
SORBIC ACID (UNII: X045WJ989B)	4 mg in 1 g
1,1-DIPHENYLUREA (UNII: H33D27I551)	50 mg in 1 g
WATER (UNII: 059QF0KO0R)	35.8 mg in 1 g
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	80 mg in 1 g

GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	12.5 mg in 1 g
PEG-75 STEARATE (UNII: OT38R0N74H)	12.5 mg in 1 g
CETETH-20 (UNII: I835H2IHHX)	12.5 mg in 1 g
STEARETH-20 (UNII: L0Q8IK9E08)	12.5 mg in 1 g
RICINUS COMMUNIS SEED (UNII: 7EK4SFN1TX)	20 mg in 1 g
ACER SACCHARUM LEAF (UNII: VP8WP1ASR7)	50 mg in 1 g
TOCOPHEROL (UNII: R0ZB2556P8)	10 mg in 1 g
CHOLESTEROL (UNII: 97C5T2UQ7J)	10 mg in 1 g
MONOBASIC POTASSIUM PHOSPHATE (UNII: 4J9FJ0HL51)	4.3 mg in 1 g
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	40 mg in 1 g
FRAGRANCE GREEN APPLE ORC2001072 (UNII: U9GH30P956)	2.1 mg in 1 g
BOVINE TYPE II COLLAGEN (TRACHEAL CARTILAGE) (UNII: 76TK29UQEZ)	40 mg in 1 g
CETYL ALCOHOL (UNII: 936JST6JCN)	5 mg in 1 g
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	15 mg in 1 g
SODIUM CHLORIDE DIHYDRATE (UNII: 70521YQ49G)	10 mg in 1 g
RETINOL (UNII: G2SH0XKK91)	0.9 mg in 1 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)	4 mg in 1 g

Product Characteristics

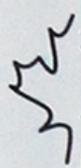
Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63550-108-01	50 g in 1 BOX; Type 0: Not a Combination Product	11/01/2001	

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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	11/01/2001	

Labeler - Kamins Dermatologics Inc. (254050784)

Registrant - Odan Laboratories Ltd (208585604)

Establishment

Name	Address	ID/FEI	Business Operations
Odan Laboratories Ltd		208585604	manufacture(63550-108) , analysis(63550-108)

Revised: 12/2022

Kamins Dermatologics Inc.