

SUPER LYSINE PLUS STRAWBERRY LIP TREATMENT- octinoxate and oxybenzone lipstick
Quantum, Inc.

Super Lysine +[®] Strawberry Lip Treatment

Drug Facts

<i>Active Ingredients</i>	<i>Purpose</i>
Octinoxate 7.0%	Sunscreen
Oxybenzone (Benzophenone-3) 5.0%	Sunscreen

Uses

Helps protect against sunburn and chapped lips.

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 prevent sunburn, not skin cancer or early skin aging.
- ***For external use only.***
- ***Do not use*** on damaged or broken skin.
- ***Stop use and consult a doctor*** if rash occurs.
- ***Avoid contact with eyes.*** If in eyes, rinse with warm water to remove.
- ***Keep out of reach of children.*** If swallowed, get medical help or contact a Poison Control Center.

Directions

Apply liberally 15 minutes before sun exposure. Use a water resistant sunscreen if swimming or sweating. Reapply at least every 2 hours. Children under 6 months: ask a doctor.

Other Information

Protect this product from excessive heat and direct sun.

Inactive Ingredients

Beeswax, Calendula Officinalis Extract, Carthamus Tinctorius (Safflower) Seed Oil, Cholecalciferol (Vitamin D), Copernicia Cerifera (Carnauba) Wax, Echinacea Purpurea

Extract, Gum Benzoin Tincture, Hydrastis Canadensis (Goldenseal) Leaf, Lysine, Melaleuca Alternifolia (Tea Tree) Oil, Melaleuca Cajuputi (Cajeput) Oil, Natural Flavor, Propolis Extract, Purified Water, Retinyl Palmitate (Vitamin A), Ricinus Communis (Castor) Seed Oil, Simmondsia Chinensis (Jojoba) Seed Oil, Tocopherol (Vitamin E), Tocopheryl Acetate (Vitamin

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PRINCIPAL DISPLAY PANEL - 5 g Cylinder Blister Pack

Quantum[®]
HEALTH

PEEL LABEL
FOR DRUG FACTS

SuperLysine⁺[®]

Strawberry Flavor
LIP TREATMENT & PROTECTANT

NET WT 5g / .17 OZ

SPF 21
SUNSCREEN

Quantum[®]
HEALTH

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R0716

SUPER LYSINE PLUS STRAWBERRY LIP TREATMENT

octinoxate and oxybenzone lipstick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70084-240
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	70 mg in 1 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
ECHINACEA PURPUREA WHOLE (UNII: QI7G114Y98)	
BENZOIN RESIN (UNII: GK21SBA74R)	
GOLDENSEAL (UNII: ZW3Z11D0JV)	
LYSINE HYDROCHLORIDE (UNII: JNJ23Q2COM)	
TEA TREE OIL (UNII: VIF565UC2G)	
MELALEUCA CAJUPUTI LEAF OIL (UNII: 5O59RMR6DU)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
SAFFLOWER (UNII: 4VBL71TY4Y)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
WATER (UNII: 059QF0KO0R)	
CASTOR OIL (UNII: D5340Y2I9G)	
JOJOBA OIL (UNII: 724GKU717M)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Product Characteristics

Color	white (Brownish-White)	Score	
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Shape		Size	
Flavor	STRAWBERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70084-240-01	5 g in 1 CYLINDER; Type 0: Not a Combination Product	04/02/2001	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/02/2001	

Labeler - Quantum, Inc. (044806305)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Deep Inc.		189788201	manufacture(70084-240)

Revised: 10/2025

Quantum, Inc.