

REDNESS SOLUTIONS MAKEUP BROAD SPECTRUM SPF 15 WITH PROBIOTIC TECHNOLOGY- octinoxate, titanium dioxide, and zinc oxide lotion
CLINIQUE LABORATORIES LLC

REDNESS SOLUTIONS MAKEUP BROAD SPECTRUM SPF 15 WITH PROBIOTIC TECHNOLOGY

Drug Facts

Active ingredients

Octinoxate 5.9%
Titanium Dioxide 3.7%
Zinc Oxide 2.9%

Purpose

Sunscreen

Use

helps prevent sunburn

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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

For sunscreen use:

- apply liberally 15 minutes before sun exposure
- reapply at least every two hours
- use a water resistant sunscreen if swimming or sweating
- **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

Inactive ingredients

water/aqua/eau • methyl trimethicone • phenyl trimethicone • dimethicone • triethylhexanoin • butylene glycol • trimethylsiloxysilicate • peg-10 dimethicone • lauryl peg-9 polydimethylsiloxyethyl dimethicone • aluminum hydroxide • c12-15 alkyl benzoate • lactobacillus ferment • citrus grandis (grapefruit) peel extract • magnolia grandiflora bark extract • poria cocos sclerotium extract • astrocaryum murumuru seed butter • glycerin • caffeine • sodium myristoyl sarcosinate • caprylyl methicone • methicone • polyglyceryl-6 polyricinoleate • disteardimonium hectorite • acetyl glucosamine • isopropyl titanium triisostearate • dimethicone crosspolymer-3 • lecithin • tocopheryl acetate • laureth-7 • dimethicone/peg-10/15 crosspolymer • sodium chloride • dipropylene glycol • disodium edta • polyaminopropyl biguanide • phenoxyethanol • [+/- mica • titanium dioxide (ci 77891) • iron oxides (ci 77491, ci 77492, ci 77499)] [iln37592]

Other information

protect the product in this container from excessive heat and direct sun

PRINCIPAL DISPLAY PANEL - 30 ml Bottle Carton

CLINIQUE

redness
solutions

makeup
broad spectrum
SPF 15
with probiotic technology

1 FL.OZ.LIQ./30 ml e

FPO/UPC/COO

70JW-01-1114

Drug Facts (continued)

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Drug Facts (continued)

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Drug Facts (continued)

polyricinoleate • disteardimonium hectorite • acetyl glucosamine • isopropyl titanium trisostearate • dimethicone crosspolymer-3 • lecithin • tocopheryl acetate • laureth-7 • dimethicone/peg-10/15 crosspolymer • sodium citrate • dipropylene glycol • disodium edta • polyaminopropyl biguanide • phenoxyethanol • [+/- mica • titanium dioxide (ci 77891) • iron oxides (ci 77491, ci 77492, ci 77499)] [In:37592]

Drug Facts (continued)

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NEW YORK, N.Y. 10022
N.Y. • LONDON W1K 3BQ • PARIS
70JW



CLINIQUE

redness solutions

makeup broad spectrum SPF 15

with probiotic technology

1 FL.OZ.LIQ./30 ml e

Lab-certified to neutralize redness on contact. Allergy Tested, 100% Fragrance Free. Moderate coverage. Natural finish.

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PROBIOTIC TECHNOLOGY

octinoxate, titanium dioxide, and zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49527-070
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	59 mg in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	37 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	29 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
METHYL TRIMETHICONE (UNII: S73ZQI0GXM)	
WATER (UNII: 059QF0KO0R)	
PHENYL TRIMETHICONE (UNII: DROK5NOJ4R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TRIETHYLHEXANOIN (UNII: 7K3W1BIU6K)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
LAURYL PEG-9 POLYDIMETHYLSILOXYETHYL DIMETHICONE (UNII: 25G622K2RA)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
LIMOSILACTOBACILLUS REUTERI (UNII: 9913I24QEE)	
GRAPEFRUIT (UNII: O82C39RR8C)	
ASTROCARYUM MURUMURU SEED BUTTER (UNII: 12V64UPU6R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAFFEINE (UNII: 3G6A5W338E)	
SODIUM MYRISTOYL SARCOSINATE (UNII: J07237209D)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
METHICONE (20 CST) (UNII: 6777U11MKT)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
N-ACETYLGLUCOSAMINE (UNII: V956696549)	
ISOPROPYL TITANIUM TRIISOSTEARATE (UNII: 949E3KBJ1I)	
LAURETH-7 (UNII: Z95S6G8201)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MICA (UNII: V8A1AW0880)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49527-070-01	1 in 1 CARTON	12/01/2011	
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2011	

Labeler - CLINIQUE LABORATORIES LLC (044475127)

Registrant - Estee Lauder Companies Inc. (790802086)

Establishment

Name	Address	ID/FEI	Business Operations
Estee Lauder N.V.		370151326	manufacture(49527-070) , pack(49527-070) , label(49527-070)

Revised: 10/2023

CLINIQUE LABORATORIES LLC