REDNESS SOLUTIONS DAILY PROTECTIVE BASE BROAD SPECTRUM SPF 15-
titanium dioxide and zinc oxide cream
CLINIQUE LABORATORIES LLC

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.

REDNESS SOLUTIONS DAILY PROTECTIVE BASE BROAD SPECTRUM SPF 15

Drug Facts

Active ingredients

Titanium dioxide 6.5%

Zinc oxide 2.5%

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 • octyldodecyl neopentanoate • c12-15 alkyl benzoate • butylene glycol • tricaprylyl citrate • steareth-2 trioctyldodecyl citrate • glyceryl stearate • peg-100 stearate • barium sulfate • silica • ascophyllum nodosum extract • asparagopsis armata extract • sea whip extract • tocopheryl acetate • sucrose • pantethine • caffeine • dimethicone • cetearyl alcohol • isostearic acid • polyhydroxystearic acid • bisabolol • magnesium ascorbyl phosphate • lactobacillus ferment • sodium hyaluronate • sorbitol • phytosphingosine • caprylyl glycol • aluminum hydroxide • stearic acid • hexylene glycol • xanthan gum • disodium edta • phenoxyethanol • iron oxides (ci 77491, ci 77492, ci 77499) • chromium hydroxide green (ci 77289) [iln39531]

Other information

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

PRINCIPAL DISPLAY PANEL - 40 ml Bottle Carton

redness

CLINIQUE

redness
solutions

daily
protective base
broad spectrum
SPF 15

ALL SKIN TYPES

1.35 FL.OZ.LIQ./40 ml e

1,2,3,4

ALL SKIN TYPES

CULTECH INC

004164

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Drug Facts (continued)

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clinique.com

redness

CLINIQUE

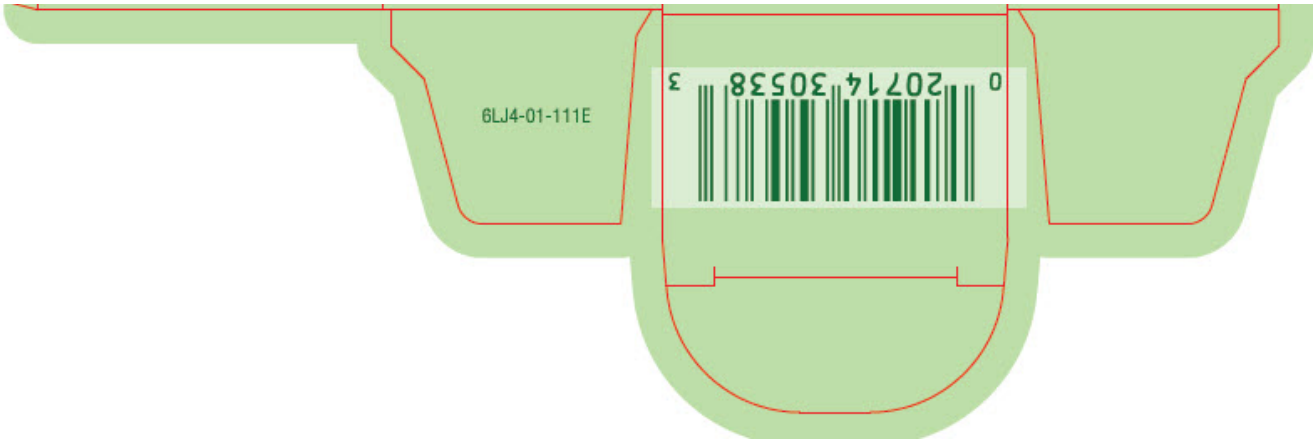
redness solutions

daily protective base
broad spectrum
SPF 15

ALL SKIN TYPES

1.35 FL.OZ.LIQ./40 ml e

Allergy Tested.
100% Fragrance Free.
Ophthalmologist Tested.
No chemical sunscreens.
Extra-gentle, physical, broad-spectrum daily protection from UVA/UVB exposure that can aggravate skins with Rosacea or reactive redness. Comforting makeup primer immediately corrects redness and evens skin tone with a sheer green tint. Oil-free. See enclosure.



REDNESS SOLUTIONS DAILY PROTECTIVE BASE BROAD SPECTRUM SPF 15

titanium dioxide and zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49527-019
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	65 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
METHYL TRIMETHICONE (UNII: S73ZQI0GXM)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
TRICAPRYLYL CITRATE (UNII: BXW1GAI4TA)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
BARIUM SULFATE (UNII: 25BB7EKE2E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
PSEUDOPTEROGORGIA ELISABETHAE (UNII: UDY3H1OUX5)	
SUCROSE (UNII: C151H8M554)	
PANTETHINE (UNII: 7K81IL792L)	
CAFFEINE (UNII: 3G6A5W338E)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
LEVOMENOL (UNII: 24WE03BX2T)	

MAGNESIUM ASCORBYL PHOSPHATE (UNII: 0R822556M5)	
LIMOSILACTOBACILLUS REUTERI (UNII: 9913I24QEE)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SORBITOL (UNII: 506T60A25R)	
PHYTOSPHINGOSINE (UNII: GIN46U9Q2Q)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
CHROMIUM HYDROXIDE GREEN (UNII: RV8FT8XF5R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49527-019-01	1 in 1 CARTON	12/01/2011	
1		40 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49527-019-02	15 mL in 1 TUBE; Type 0: Not a Combination Product	12/01/2022	

Marketing Information

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

Labeler - CLINIQUE LABORATORIES LLC (044475127)

Registrant - Estee Lauder Companies Inc. (790802086)

Establishment

Name	Address	ID/FEI	Business Operations
The Estee Lauder Inc		802599436	pack(49527-019) , label(49527-019) , manufacture(49527-019)

Establishment

Name	Address	ID/FEI	Business Operations
NORTHTEC LLC		943871157	pack(49527-019) , label(49527-019)