

MOISTURE SURGE CC BROAD SPECTRUM SPF 30 HYDRATING COLOUR CORRECTOR- octinoxate, octisalate, titanium dioxide, and zinc oxide cream
CLINIQUE LABORATORIES LLC

MOISTURE SURGE CC

BROAD SPECTRUM SPF 30 HYDRATING COLOUR CORRECTOR

Drug Facts

Active ingredients

Octinoxate 7.5%
Octisalate 5.0%
Titanium dioxide 5.2%
Zinc oxide 3.2%

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 • squalane • octyldodecyl neopentanoate • butylene glycol • propanediol • glyceryl stearate • behenyl alcohol • peg-40 stearate • polyglyceryl-10 pentastearate • aloe barbadensis leaf water • thermus thermophilus ferment • trehalose • caffeine • glycerin • linoleic acid • lecithin • stearic acid • hydrogenated lecithin • sorbitol • polyglyceryl-6 polyricinoleate • tocopheryl acetate • ammonium acryloyldimethyltaurate/vp copolymer • sodium stearyl lactylate • sodium hyaluronate • tetrahexyldecyl ascorbate • pentaerythrityl tetra-di-t-butyl hydroxyhydrocinnamate • isopropyl titanium triisostearate • synthetic fluorophlogopite • xanthan gum • silica • alumina • sodium dehydroacetate • tin oxide • disodium edta • phenoxyethanol • [+/- mica • titanium dioxide (ci 77891) • iron oxides (ci 77492) • iron oxides (ci 77491) • iron oxides (ci 7799)] [iln39833]

Other information

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

CLINIQUE LABS., DIST.
NEW YORK, N.Y. 10022

PRINCIPAL DISPLAY PANEL - 40 ml Bottle Carton

C

CLINIQUE

moisture surge

CC

cream

hydrating

colour corrector

broad spectrum

SPF 30

ALL SKIN TYPES

1.4 FL.OZ.LIQ./40 ml e

1, 2, 3, 4
ALL SKIN TYPES

S.3

7WCP-01-1111



87683
C3219222604

87683



**Allergy Tested.
100% Fragrance Free.**

Next-generation perfecting formula delivers colour-correction, hydration and UVA/UVB protection for a more flawless look today and tomorrow—all with a refreshingly lightweight feel. Special optics help correct dullness, sallowness and other concerns to enhance skin's radiance. In shades that provide natural-looking coverage. Wears comfortably alone or under foundation. Oil-free. See enclosure.



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Drug Facts (cont'd)

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NEW YORK, N.Y. 10022
LONDON W1K 3BQ • PARIS
MADE IN BELGIUM
7WCP



clinique.com



CLINIQUE

moisture surge

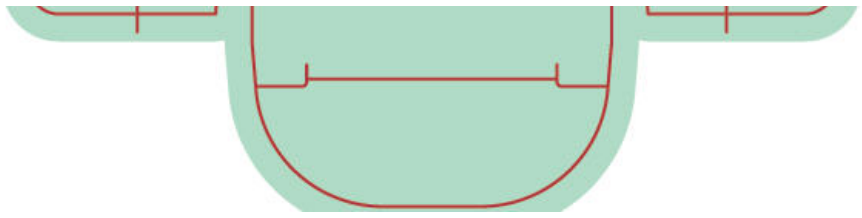
CC
cream

hydrating
colour corrector
broad spectrum
SPF 30

ALL SKIN TYPES

1.4 FL.OZ.LIQ./40 ml e

7WCP-01-1111



MOISTURE SURGE CC BROAD SPECTRUM SPF 30 HYDRATING COLOUR CORRECTOR

octinoxate, octisalate, titanium dioxide, and zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49527-029
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	52 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	32 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
THERMUS THERMOPHILUS LYSATE (UNII: 775R692494)	
POLYGLYCERYL-10 PENTASTEARATE (UNII: PMX5872701)	
WATER (UNII: 059QF0KO0R)	
SQUALANE (UNII: GW89575KF9)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PROPANEDIOL (UNII: 5965N8W85T)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
DOCOSANOL (UNII: 9G1OE216XY)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TREHALOSE (UNII: B8WCK70T7I)	
CAFFEINE (UNII: 3G6A5W338E)	
GLYCERIN (UNII: PDC6A3C0OX)	
LINOLEIC ACID (UNII: 9KJL21T0QJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SORBITOL (UNII: 506T60A25R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
SODIUM STEAROYL LACTYLATE (UNII: IN99IT31LN)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	

PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)
ISOPROPYL TITANIUM TRIISOSTEARATE (UNII: 949E3KBJ1I)
XANTHAN GUM (UNII: TTV12P4NEE)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
ALUMINUM OXIDE (UNII: LMI26O6933)
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)
STANNIC OXIDE (UNII: KM7N50LOS6)
EDETATE DISODIUM (UNII: 7FLD91C86K)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
MICA (UNII: V8A1AW0880)
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)
FERRIC OXIDE RED (UNII: 1K09F3G675)
FERROSFERRIC OXIDE (UNII: XM0M87F357)
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)
POLYGLYCERYL-6 POLYRICINOLEATE (UNII: YPM0ZOC2HR)
MAGNESIUM POTASSIUM ALUMINOSILICATE FLUORIDE (UNII: YK3DC63Y5M)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49527-029-01	1 in 1 CARTON	10/05/2022	
1		40 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49527-029-02	15 mL in 1 TUBE; Type 0: Not a Combination Product	10/05/2022	12/31/2023
3	NDC:49527-029-03	7 mL in 1 TUBE; Type 0: Not a Combination Product	10/05/2022	12/31/2023

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/05/2022	

Labeler - CLINIQUE LABORATORIES LLC (044475127)

Establishment			
Name	Address	ID/FEI	Business Operations
Estee Lauder Cosmetics Ltd		202952982	manufacture(49527-029)

Establishment			
Name	Address	ID/FEI	Business Operations
Estee Lauder Cosmetics Ltd.		204132062	pack(49527-029) , label(49527-029) , manufacture(49527-029)

Establishment

Name	Address	ID/FEI	Business Operations
The Estee Lauder Inc		802599436	manufacture(49527-029) , label(49527-029) , pack(49527-029)

Establishment

Name	Address	ID/FEI	Business Operations
PADC		949264774	label(49527-029) , pack(49527-029)

Revised: 1/2024

CLINIQUE LABORATORIES LLC