

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

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**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)] [ln39833]

## 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  
C321922604

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

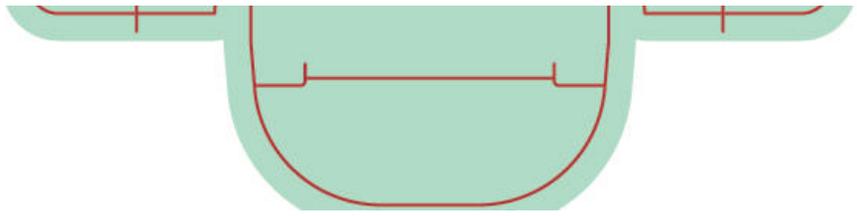
CC  
cream

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

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

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
<b>MICA</b> (UNII: V8A1AW0880)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERROSFERRIC OXIDE</b> (UNII: XM0M87F357)	
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)	
<b>POLYGLYCERYL-6 POLYRICINOLEATE</b> (UNII: YPM0ZOC2HR)	
<b>MAGNESIUM POTASSIUM ALUMINOSILICATE FLUORIDE</b> (UNII: YK3DC63Y5M)	
<b>THERMUS THERMOPHILUS LYSATE</b> (UNII: 775R692494)	
<b>POLYGLYCERYL-10 PENTASTEARATE</b> (UNII: PMX5872701)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SQUALANE</b> (UNII: GW89575KF9)	
<b>OCTYLDODECYL NEOPENTANOATE</b> (UNII: X8725R883T)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>PROPANEDIOL</b> (UNII: 5965N8W85T)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>DOCOSANOL</b> (UNII: 9G1OE216XY)	
<b>PEG-40 STEARATE</b> (UNII: ECU18C66Q7)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>TREHALOSE</b> (UNII: B8WCK70T7I)	
<b>CAFFEINE</b> (UNII: 3G6A5W338E)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>LINOLEIC ACID</b> (UNII: 9KJL21T0QJ)	

<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)
<b>SORBITOL</b> (UNII: 506T60A25R)
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)
<b>AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER</b> (UNII: W59H9296ZG)
<b>SODIUM STEAROYL LACTYLATE</b> (UNII: IN99IT31LN)
<b>HYALURONATE SODIUM</b> (UNII: YSE9PPT4TH)
<b>TETRAHEXYLDECYL ASCORBATE</b> (UNII: 9LBV3F07AZ)
<b>PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE)</b> (UNII: 255PIF62MS)
<b>ISOPROPYL TITANIUM TRIISOSTEARATE</b> (UNII: 949E3KBJ1I)
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)
<b>SODIUM DEHYDROACETATE</b> (UNII: 8W46YN971G)
<b>STANNIC OXIDE</b> (UNII: KM7N50LOS6)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)

<b>Packaging</b>				
#	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

<b>Marketing Information</b>			
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)

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

<b>Establishment</b>			
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)

Revised: 8/2025

CLINIQUE LABORATORIES LLC