

**CC SCREEN 100% MINERAL CC CREAM BROAD SPECTRUM SPF 50 426W-titanium dioxide, zinc oxide cream
Supergoop, 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.

CC Screen 100% Mineral CC Cream Broad Spectrum SPF 50 426W

Active Ingredients Purpose

Titanium Dioxide 4% Sunscreen

Zinc Oxide 20% Sunscreen

Uses

- Helps Prevent Sunburn
- If used as directed with other sun protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Keep out of reach of children. If product is swallowed, get medical help or contact a poison Control Center right away.

Stop use and ask a doctor if rash occurs or rash occurs

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.

Directions

- apply generously and evenly 15 minutes before sun exposure
- reapply at least every 2 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 your 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, C12-15 Alkyl Benzoate, Iron Oxides, Butyloctyl Salicylate, Isododecane, Propanediol, Glycerin, Cetyl Diglyceryl Tris(Trimethylsiloxy)silylethyl Dimethicone, Dimethicone, 1,2-Hexanediol, Pyrus Malus (Apple) Fruit Extract, Sodium Chloride, Yellow 5 Lakes, Silica, Triethoxycaprylylsilane, Dimethicone/Vinyl Dimethicone Crosspolymer, Mica, Hydroxyacetophenone, Diethylhexyl Syringlidenemalonate, Titanium Dioxide, Trisodium Ethylenediamine Disuccinate, Chlorphenesin, Xanthan Gum, Chondrus

Crispus Extract, Caprylic/Capric Triglyceride, Tin Oxide, Sodium Hyaluronate, Tocopherol

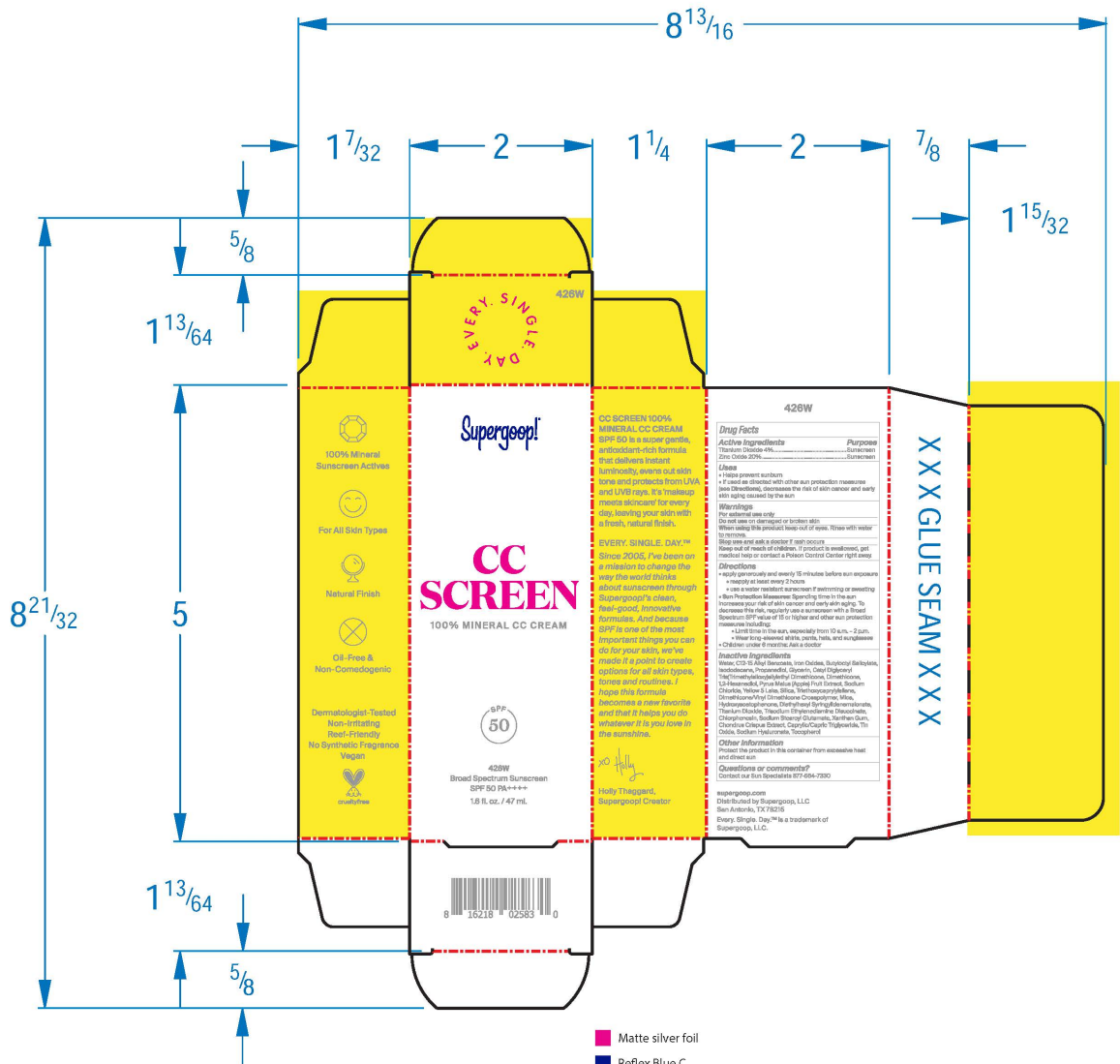
CC Screen

100% Mineral CC Cream 426W

SPF 50

Broad Spectrum SPF 50 PA+++

1.6 fl. oz./ 47 ml.



CC SCREEN 100% MINERAL CC CREAM BROAD SPECTRUM SPF 50 426W

titanium dioxide, zinc oxide cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:75936-239

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|----------------|
| ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) | ZINC OXIDE | 20 g in 100 mL |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP) | TITANIUM DIOXIDE | 4 g in 100 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| ISODODECANE (UNII: A8289P68Y2) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| .BETA.-TOCOPHEROL (UNII: 9U6A490501) | |
| CHLORPHENESIN (UNII: I670DAL4SZ) | |
| XANTHAN GUM (UNII: TTV12P4NEE) | |
| PROPANEDIOL (UNII: 5965N8W85T) | |
| ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ) | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| MICA (UNII: V8A1AW0880) | |
| FERRIC OXIDE YELLOW (UNII: EX438O2MRT) | |
| WATER (UNII: 059QF0K00R) | |
| BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3) | |
| 1,2-HEXANEDIOL (UNII: TR046Y3K1G) | |
| DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5) | |
| APPLE (UNII: B423VGH5S9) | |
| SODIUM CHLORIDE (UNII: 451W47IQ8X) | |
| MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U) | |
| STANNIC OXIDE (UNII: KM7N50LOS6) | |
| HYALURONATE SODIUM (UNII: YSE9PPT4TH) | |
| DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248) | |
| HYDROXYACETOPHENONE (UNII: G1L3HT4CMH) | |
| TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E) | |
| FERRIC OXIDE RED (UNII: 1K09F3G675) | |
| TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q) | |
| CHONDRUS CRISPUS CARRAGEENAN (UNII: UE856F2T78) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:75936-239-01 | 47 mL in 1 CARTON; Type 0: Not a Combination Product | 06/01/2020 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------|--|----------------------|--------------------|
| OTC monograph final | part352 | 05/21/2020 | |

Labeler - Supergoop, LLC (117061743)

Revised: 7/2023

Supergoop, LLC