

**PLAY MINERAL BROAD SPECTRUM SPF 50- titanium dioxide, zinc oxide lotion  
Supergoop, LLC**

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**Play 100% Mineral Lotion Broad Spectrum Sunscreen SPF 50**

Active Ingredients Purpose

Titanium Dioxide 2% Sunscreen

Zinc Oxide 24% 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.

***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 liberally 15 minutes before sun exposure

Reapply:

- after 80 minutes of swimming or sweating
- Immediately after towel drying
- At least every 2 hours

**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 15 or higher and other sun protection measures including:

- Limit time in the sun, especially from 10a.m. – 2p.m.
- Wear long-sleeve shirts, pants, hats, and sunglasses
- Children under 6 months: Ask a doctor

***Inactive Ingredients***

C9-12 Alkane, Caprylic/Capric Triglyceride, Cetyl Esters, Coco-Caprylate/Caprato, Haematococcus Pluvalis Extract, Helianthus Annuus (Sunflower) Seed Wax, Hydrogenated Rapeseed Oil, Hydrogenated Rapeseed Oil, Hydrolyzed Jojoba Esters, Isopropyl Palmitate, Jojoba Esters, Perilla Ocymoides (Perilla) Seed Oil, Polyhydroxystearic Acid, Punica Granatum (Pomegranate) Seed Oil, Sea Water, Silica, Stearic Acid, Water

Play 100% Mineral Lotion

with green Algae  
 Broad Spectrum Sunscreen SPF 50  
 Water Resistant (80 Minutes)



**PLAY MINERAL BROAD SPECTRUM SPF 50**  
 titanium dioxide, zinc oxide lotion

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**Product Information**

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:75936-175 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength | Strength       |
|---|-------------------|----------------|
| <b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP) | TITANIUM DIOXIDE  | 2 g in 100 mL  |
| <b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)             | ZINC OXIDE        | 24 g in 100 mL |

### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>CETYL ESTERS WAX</b> (UNII: D072FFP9GU)                     |          |
| <b>COCO-CAPRYLATE/CAPRATE</b> (UNII: 8D9H4QU99H)               |          |
| <b>PERILLA FRUTESCENS SEED OIL</b> (UNII: 322MS57V7Z)          |          |
| <b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)                      |          |
| <b>HYDROLYZED JOJOBA ESTERS (ACID FORM)</b> (UNII: UDR641JW8W) |          |
| <b>HAEMATOCOCCUS PLUVIALIS</b> (UNII: 31TOFF0472)              |          |
| <b>HELIANTHUS ANNUUS SEED WAX</b> (UNII: 42DG15CHXV)           |          |
| <b>HYDROGENATED RAPESEED OIL</b> (UNII: K168T6Y0YU)            |          |
| <b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)                  |          |
| <b>POLYHYDROXYSTEARIC ACID (2300 MW)</b> (UNII: YXH47AOU0F)    |          |
| <b>POMEGRANATE SEED OIL</b> (UNII: OUI45XV0T6)                 |          |
| <b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)                         |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                                |          |
| <b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)                      |          |
| <b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)           |          |

### Packaging

| # | Item Code        | Package Description                                   | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:75936-175-01 | 100 mL in 1 BOTTLE; Type 0: Not a Combination Product | 05/16/2019           |                    |
| 2 | NDC:75936-175-02 | 30 mL in 1 BOTTLE; Type 0: Not a Combination Product  | 05/16/2019           |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M020                                     | 05/16/2019           |                    |

**Labeler** - Supergoop, LLC (117061743)

### Establishment

| Name | Address | ID/FEI | Business Operations |
|------|---------|--------|---------------------|
|      |         |        |                     |

Revised: 1/2026

Supergoop, LLC