# MINERAL SHEER SCREEN BROAD SPECTRUM SUNSCREEN SPF 30- zinc oxide lotion 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.

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### Mineral Sheer Screen Broad Spectrum Sunscreen SPF 30

Active Ingredient Purpose

Zinc Oxide 17.5% Sunscreen

Sunscreen

Helps prevent sunburn.

If used as directed with other sun protection measures (see Directions), decreases the risk of skin cancer and early skin againg 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.

For external use only

Do not use on damaged or broken skin

When using this product kee out of eyes. Rinse with water to remove.

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

Water, Isododecane, Caprylic/Capric Triglyceride, Butyloctyl Salicylate, Propanedlol, Glycerin, Squalane, Trioctyldodecyl Citrate, Polyglycerln-3, Polyglyceryl-3 Lactate/Laurate, Polysilicone-11, Cetearyl Alcohol, Glyceryl Stearate, Sodium Stearoyl Glutamate, Butylene Glycol, Xanthan Gum, Polyacrylate Crosspolymer-6, Isostearic Acid, Lecithin, Polyglyceryl-3 Polyricinoleate, Polyhydroxystearic Acid, Caprylhydroxamic Acid, Citric Acid, Aloe Barbadensis Leaf Juice, Lespedeza Capitata Leaf/Stem Extract, Maltodextrin, Sodium Hyaluronate

Protect the product In this container from excessive heat and direct sun.

Mineral Sheer Screen

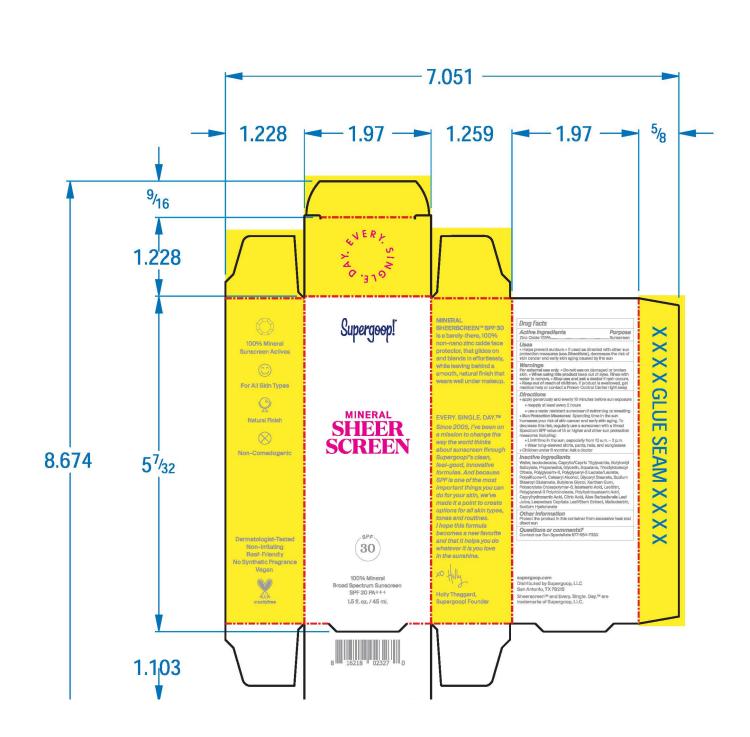
**SPF 30** 

100% Mineral

**Broad Spectrum Sunscreen** 

SPF 30 PA+++

1.5 fl. oz./45 ml

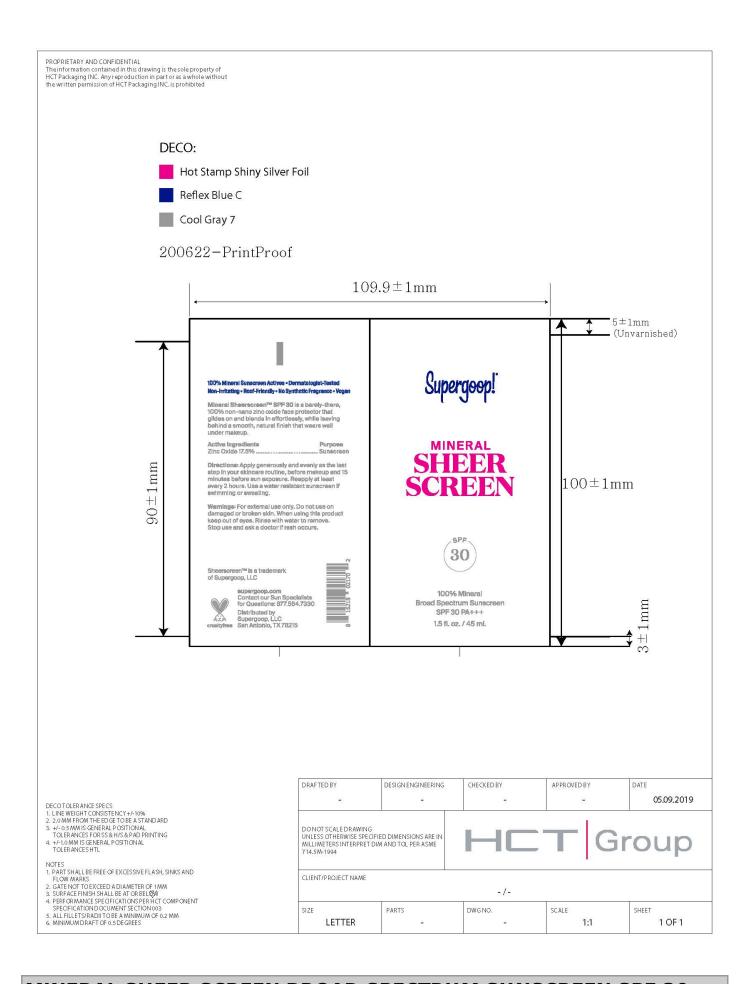




Hot Stamp Silver Foil
Reflex Blue C

Cool Gray 7

PMS 102 U



#### **Product Information**

HUMAN OTC DRUG NDC:75936-305 **Product Type** Item Code (Source)

**Route of Administration TOPICAL** 

#### **Active Ingredient/Active Moiety**

**Ingredient Name Basis of Strength** Strength

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) ZINC OXIDE 17.5 g in 100 mL

#### **Inactive Ingredients**

Ingredient Name	Strength
ISODODECANE (UNII: A8289P68Y2)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

**GLYCERIN** (UNII: PDC6A3C0OX)

TRIOCTYLDODECYL CITRATE (UNII: 35X8CT063R) **GLYCERYL MONOSTEARATE** (UNII: 2300U9XXE4)

**BUTYLENE GLYCOL** (UNII: 3XUS85K0RA)

POLYGLYCERYL-3 PENTARICINOLEATE (UNII: 7Q00K5D0T4)

POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)

CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)

AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)

**SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)** 

POLYGLYCERIN-3 (UNII: 4A0NCJ6RD6)

DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

**HYALURONATE SODIUM** (UNII: YSE9PPT4TH) **BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)** 

PROPANEDIOL (UNII: 5965N8W85T)

**SQUALANE** (UNII: GW89575KF9) ISOSTEARIC ACID (UNII: X33R8U0062)

LECITHIN, SOYBEAN (UNII: 1DI56QDM62)

CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)

XANTHAN GUM (UNII: TTV12P4NEE)

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

MALTODEXTRIN (UNII: 7CVR7L4A2D)

WATER (UNII: 059QF0KO0R)

#### **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75936-305- 01	45 mL in 1 TUBE; Type 0: Not a Combination Product	01/12/2021	
٦	NDC:75936-305-	15 mL in 1 TUBE; Type 0: Not a Combination	01/12/2021	

I	4	02	Product	U1/12/2U21	
	3	NDC:75936-305- 03	10 mL in 1 TUBE; Type 0: Not a Combination Product	01/12/2021	
	4	NDC:75936-305- 04	5 mL in 1 TUBE; Type 0: Not a Combination Product	01/12/2021	
	5	NDC:75936-305- 05	20 mL in 1 TUBE; Type 0: Not a Combination Product	08/26/2021	

Marketing In			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/12/2021	

## Labeler - Supergoop LLC (117061743)

Revised: 10/2022 Supergoop LLC