

**UNSEEN SUNSCREEN BROAD SPECTRUM SPF 40- avobenzone, homosalate, octinoxate, octisalate cream
Supergoop, LLC**

Unseen Sunscreen Broad Spectrum SPF 40

Active Ingredients	Purpose
Avobenzone 3%	Sunscreen
Homosalate 8%	Sunscreen
Octocrylene 4%	Sunscreen
Octisalate 5%	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

Directions

- apply generously and evenly 15 minutes before sun exposure
- reapply:

after 40 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 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: ask a doctor

Inactive Ingredients

Isododecane, Dimethicone Crosspolymer, Dimethicone/Bis-Isobutyl PPG-20 Crosspolymer, Polymethylsilsesquioxane, Isohexadecane, Dicrapyl Carbonate, Meadowfoam Estolide, Caprylic/Capric Triglyceride, Polyester-7, Neopentyl Glycol Diheptanoate, Lithothamnion Calcareum Extract, Butyrospermum Parkii (Shea) Butter,

Jojoba Esters, Mannitol, Boswellia Serrata Resin Extract, Lecithin, Microcrystalline Cellulose, Diatomaceous Earth, Zinc Sulfate, Silica, Tocopherol

EVERY SINGLE DAY

SPF IS THE #1 THING you can do for your skin

Supergoop!

UNSEEN SUNSCREEN™
SPF 50 is our iconic 100% invisible SPF for face that's weightless, colorless and scentless.

Bestseller!

ZERO WHITE CAST!

A BIRTHDAY GIFT FROM ONE ICON TO ANOTHER

It's your birthday wish come true!

Drug Facts

Active Ingredients	Purpose
Avobenzone 3%, Homosalate 7%	Sunscreen
Octisalate 5%, Octocrylene 9%	

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.

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
• apply generously and evenly 15 minutes before sun exposure
• reapply
• after 40 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 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: Ask a doctor

Inactive Ingredients
Isododecane, Dimethicone/Bis-Isobutyl PPG-20 Crosspolymer, Dimethicone/Vinyl Dimethicone Crosspolymer, Dimethicone, Dimethicone Crosspolymer, Caprylic/Capric Triglyceride, Butyrospermum Parkii (Shea) Butter, Physalis Angulata Extract, Himerthalia Elongata Extract, Meadowfoam Estolide, Tocopherol, Jojoba Esters, Polymethylsilsesquioxane, Caprylyl Glycol, Diethylhexyl Syringlideneisobutylate, Silica, Bisabolol, Dipotassium Glycyrrhizate.

Other Information Protect the product in this container from excessive heat and direct sun

Questions or comments?
Contact our Sun Specialists 877-564-7330

supergoop.com
Distributed by Supergoop LLC
San Antonio, TX 78215

ARTWORK PURPOSE

UNSEEN SUNSCREEN BROAD SPECTRUM SPF 40

avobenzone, homosalate, octinoxate, octisalate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75936-164
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	8 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	4 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ISODODECANE (UNII: A8289P68Y2)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	
DIMETHICONE/BIS-ISOBUTYL PPG-20 CROSSPOLYMER (UNII: O4I3UFO6ZF)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
POLYESTER-7 (UNII: 0841698D2F)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
POLYMETHYLSILSESQUIOXANE (11 MICRONS) (UNII: Z570VEV8XK)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PHYMATOLITHON CALCAREUM (UNII: 6J1M3WA0ZK)	
SHEA BUTTER (UNII: K49155WL9Y)	
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)	
MANNITOL (UNII: 3OWL53L36A)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
LECITHIN, SUNFLOWER (UNII: 834K0WOS5G)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
DIATOMACEOUS EARTH (UNII: 2RF6EJ0M85)	
ZINC SULFATE (UNII: 89DS0H96TB)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TOCOPHEROL (UNII: ROZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75936-164-02	1 in 1 CARTON	10/12/2017	
1	NDC:75936-164-01	50 mL in 1 PACKAGE; Type 0: Not a Combination Product		
2	NDC:75936-164-06	15 mL in 1 TUBE; Type 0: Not a Combination Product	10/12/2017	
3	NDC:75936-164-03	10 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/12/2017	
4	NDC:75936-164-04	5 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/12/2017	
5	NDC:75936-164-05	1.5 mL in 1 PACKAGE; Type 0: Not a Combination Product	10/12/2017	
6	NDC:75936-164-07	30 mL in 1 TUBE; Type 0: Not a Combination Product	08/17/2020	
7	NDC:75936-164-08	20 mL in 1 CARTON; Type 0: Not a Combination Product	02/15/2021	

8	NDC:75936-164-09	73.9 mL in 1 TUBE; Type 0: Not a Combination Product	10/27/2017	
9	NDC:75936-164-11	1 in 1 CARTON	10/10/2017	
9	NDC:75936-164-10	50 mL in 1 TUBE; Type 0: Not a Combination Product		
10	NDC:75936-164-13	1 in 1 CARTON	08/07/2024	
10	NDC:75936-164-12	15 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/10/2017	

Labeler - Supergoop, LLC (117061743)

Revised: 8/2024

Supergoop, LLC