## UNSEEN SUNSCREEN BODY SPF 40- avobenzone, homosalate, octisalate, octocrylene 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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## **Unseen Sunscreen Body SPF 40**

Avobenzone 3%

Homosalate 9%

Octisalate 5%

Octocrylene 10%

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.

- 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, Caprylyl Methicone, Dimethicone/Bis-Isobutyl PPG-20 Crosspolymer, Caprylic/CapricTriglyceride, Phenyl Trimethicone, Olea Europaea Olive Oil, Olea Europaea (Olive) Fruit Extract, Olea Europaea (Olive) Leaf Extract, Silica Dimethyl Silyate

## 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 is rash occurs

Unseen Sunscreen Body

SPF 40 PA +++

**Broad Spectrum Sunscreen** 

3.4 fl. oz. / 100 mL



avobenzone, homosalate, octisalate, octocrylene lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75936-605
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL	
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 g in 100 mL	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	9 g in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
DIMETHICONE/BIS-ISOBUTYL PPG-20 CROSSPOLYMER (UNII: O4I3UFO6ZF)		
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)		
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)		
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)		
OLEA EUROPAEA FRUIT VOLATILE OIL (UNII: 8E7358CX1J)		
OLEA EUROPAEA (OLIVE) OIL UNSAPONIFIABLES (UNII: XO45V955LT)		
ISODODECANE (UNII: A8289P68Y2)		
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)		
OLEA EUROPAEA LEAF (UNII: MJ95C3OH47)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:75936-605- 01	100 mL in 1 TUBE; Type 0: Not a Combination Product	10/28/2022		
2	NDC:75936-605- 02	30 mL in 1 TUBE; Type 0: Not a Combination Product	10/28/2022		
3	NDC:75936-605- 03	10 mL in 1 PACKET; Type 0: Not a Combination Product	10/28/2022		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	M020	10/28/2022		
OTC monograph final	M020	10/28/2022		

Revised: 10/2022 Supergoop, LLC