

**TRIPLE PREP SPF 40- avobenzene, octisalate, octocrylene, homosalate lotion  
Supergoop, LLC**

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**Triple Prep Weightless multitasking moisturizer SPF 40**

**Active Ingredients:** Avobenzene 3%, Homosalate 7%, Octisalate 5%, Octocrylene 6%

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.

**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

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 sunglasses

Children under 6 months: Ask a doctor

**Inactive Ingredients:** Water, Isododecane, Butyloctyl Salicylate, Calcium Aluminum Borosilicate, Glycerin, Polymethylsilsesquioxane, Polyglycerin-3, Polyglyceryl-3 Lactate/Laurate, Ammonium Acryloyldimethyltaurate/VP Copolymer, Ectoin, Acrylates Crosspolymer, Gluconolactone, Arginine, Butylene Glycol, Sodium Benzoate, Polyacrylate Crosspolymer-6, Bisabolol, Tocopherol, Citric Acid, Niacinamide, Sodium Dilauramidoglutamide Lysine, Sodium Hyaluronate, Calcium Gluconate, Sodium Polyglutamate, Hydrolyzed Rhodophyceae Extract

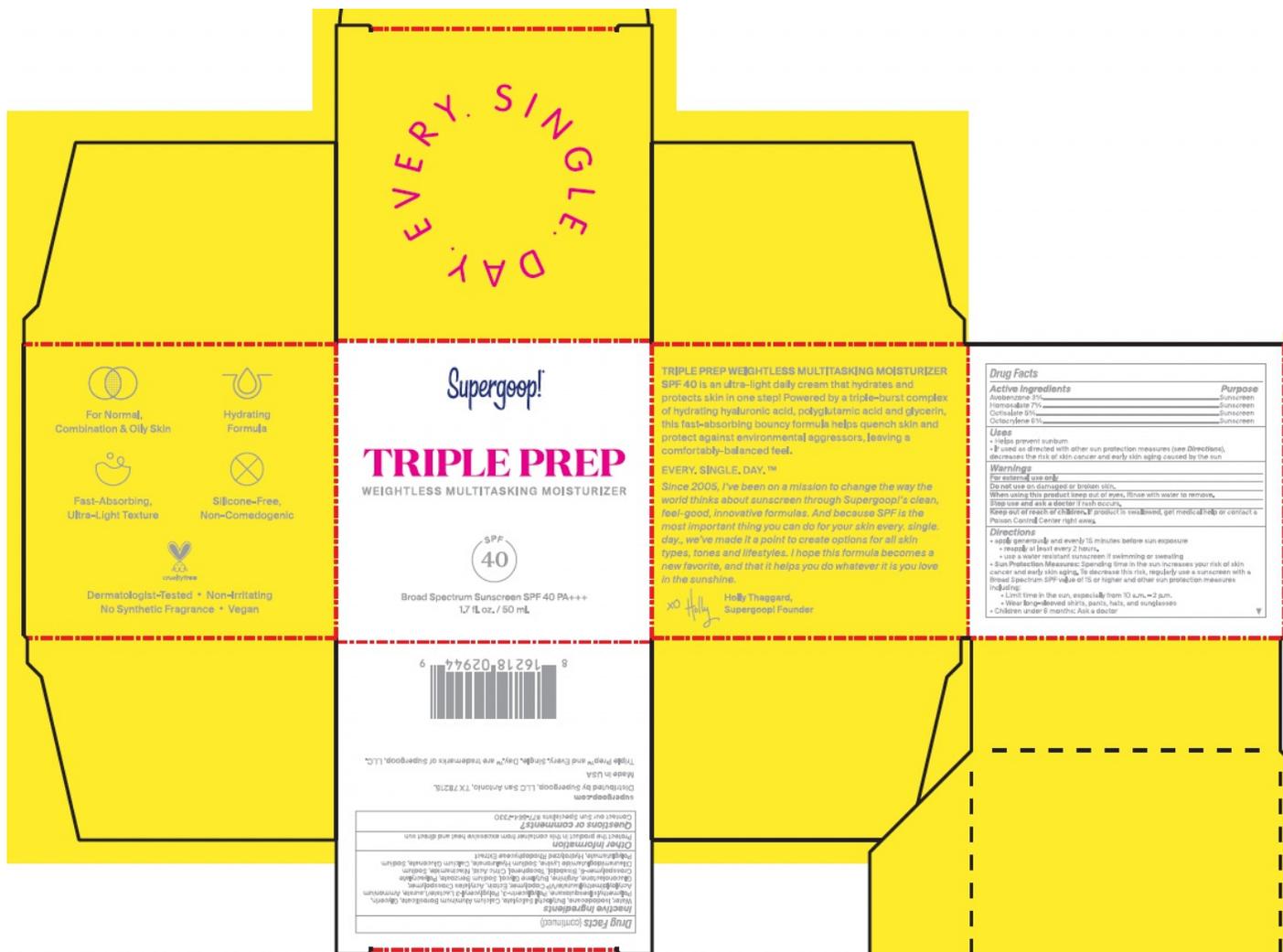
TRIPLE PREP

Weightless Multitasking Moisturizer

SPF 40

Broad Spectrum Sunscreen SPF 40 PA +++

1.7 fl. oz. / 50 mL



## TRIPLE PREP SPF 40

avobenzone, octisalate, octocrylene, homosalate lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:75936-621
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZ ONE	3 g in 100 mL
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	6 g in 100 mL
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	

<b>ARGININE</b> (UNII: 94ZLA3W45F)
<b>SODIUM DILAURAMIDOGlutAMIDE LYSINE</b> (UNII: MNJ7VPT2R5)
<b>POLYMETHYLSILSESQUIOXANE (11 MICRONS)</b> (UNII: Z570VEV8XK)
<b>AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER</b> (UNII: W59H9296ZG)
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)
<b>ECTOINE</b> (UNII: 7GXZ3858RY)
<b>LEVOMENOL</b> (UNII: 24WE03BX2T)
<b>CALCIUM SODIUM PHOSPHATE</b> (UNII: 7850XT8YDT)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>POLYGLYCERIN-3</b> (UNII: 4A0NCJ6RD6)
<b>POLYGLYCERYL-3 LAURATE</b> (UNII: Y9ZSR39D0E)
<b>HYALURONATE SODIUM</b> (UNII: YSE9PPT4TH)
<b>SODIUM LAUROYL GLUTAMATE</b> (UNII: NCX1UU2D33)
<b>ACRYLATES CROSSPOLYMER-4</b> (UNII: GEV2EL4D9G)
<b>GLUCONOLACTONE</b> (UNII: WQ29KQ9POT)
<b>NIACINAMIDE</b> (UNII: 25X5118RD4)
<b>AMMONIUM ACRYLOYLDIMETHYLTAURATE, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S)</b> (UNII: Q7UI015FF9)
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)
<b>CALCIUM GLUCONATE</b> (UNII: SQE6VB453K)
<b>WATER</b> (UNII: 059QF0K00R)
<b>ISODODECANE</b> (UNII: A8289P68Y2)
<b>BUTYLOCTYL SALICYLATE</b> (UNII: 2EH13UN8D3)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75936-621-02	1 in 1 BOX	07/12/2023	
1	NDC:75936-621-01	50 mL in 1 JAR; 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	07/12/2023	

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