GLOW SCREEN BODY BROAD SPECTRUM SUNSCREEN SPF 40- avobenzone, homosalate, octisalate, octocrylene cream
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.

Glow Screen Body Broad Spectrum Sunscreen SPF 40

Avobenzone 3%.....Sunscreen

Homosalate 7%.....Sunscreen

Octisalate 5%.....Sunscreen

Octocrylene 10%.....Sunscreen

- 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.

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.

- 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 decreases this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection meausres 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.

Aletris Farinosa Root Extract, Bisabolol, Butyloctyl Salicylate,Calcium Sodium Borosilicate, Caprylhydroxamic Acid, Caprylic/Capric Triglyceride, Capryly Glycol, Cetearyl Alcohol, Coco-Caprylate/Caprate, Coco-Glucoside, Coconut Alkanes, Erythritol, Ethyl Ferulate, Ethylhexyl Methoxycrylene, Glycerin, Glyceryl Stearate, Hydrogenated Polycyclopentadiene, Methyl Dlhydroabietate, Mica, Polyester-8, Propanedlol, Sodium Gluconate, Sodium Polyacrylate Starch, Sodium Stearoyl Glutamate, Titanium Dioxide, Tocopherol, Water, Xanthan Gum

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

Supergoop!

Glow Screen Body

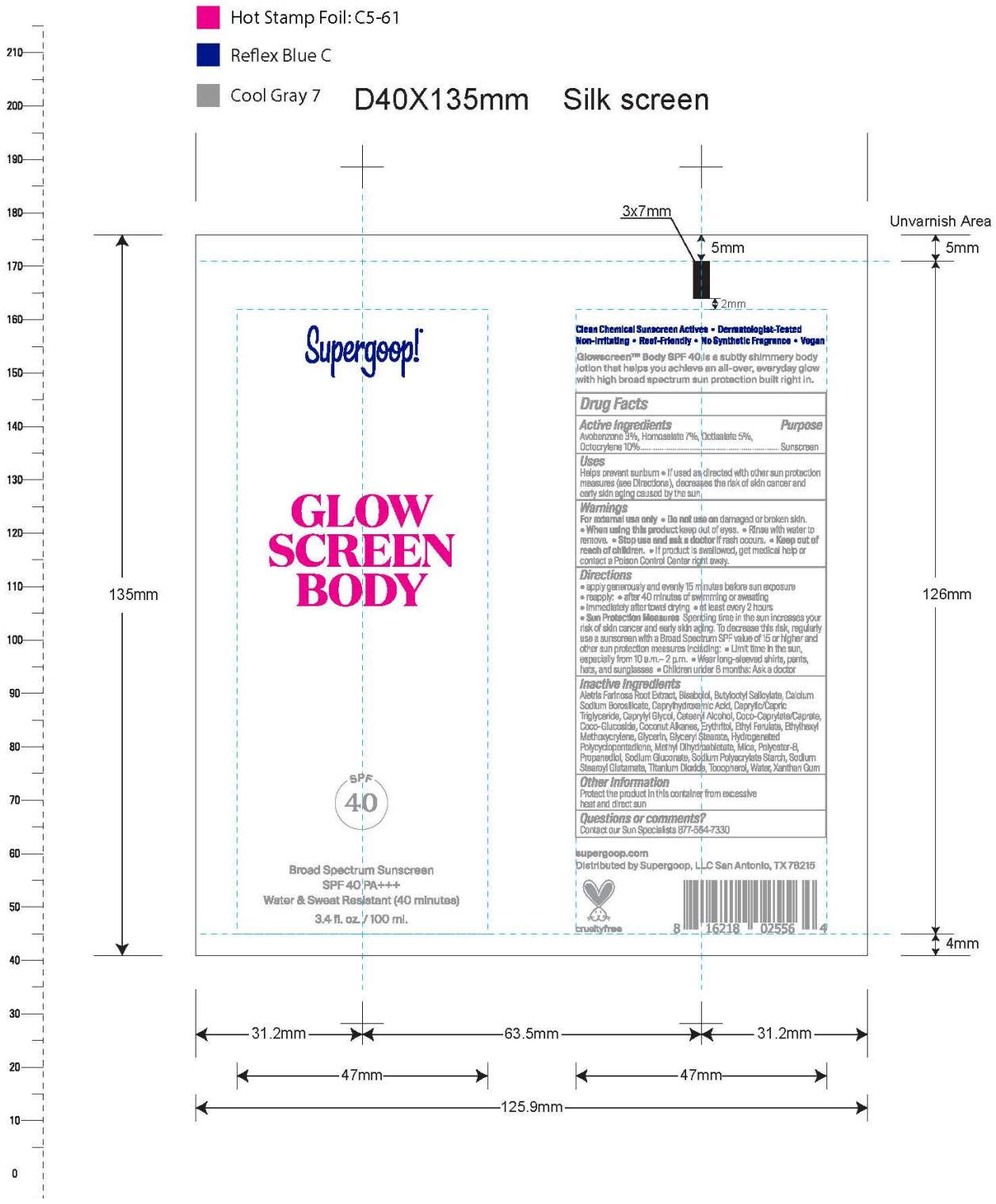
SPF 40

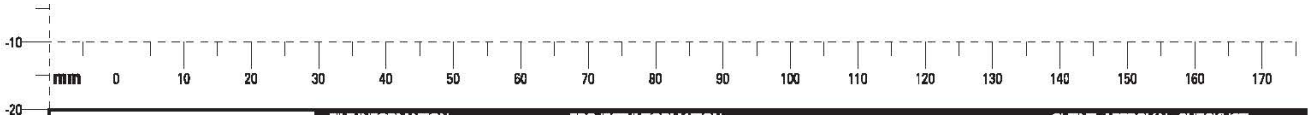
Broad Spectrum Sunscreen

SPF 40 PA+++

Water & Sweat Resistant (40 minutes)

3.4 fl. oz. / 100 ml.





berlin™ PACKAGING

PROOF NUMBER	PROOF SIZE
1	100%

FILE INFORMATION

FILE NAME SG_MineralSunscreen_2017

CLIENT

STARTED 12/06/18 CGP

MODIFIED 00/00/00 CGP

SOFTWARE ai by Illustrator CC

EXPROT PDF

PROJECT INFORMATION

TUBE DIAMETER ∅ 40 mm

TUBE LENGTH 135 mm

PRINT INFORMATION

OFFSET COLOR

SILK-SCREEN COLOR

☐ TUBE COLOR

Hot Stamping

☐ Silver ☐ Gold

☐ Other Metallic Color

5 mm unvarnished from bottom

INTERNAL APPROVAL

Design _____ Proof _____

UPC Test _____ Verify _____

RELEASE DATE

DIELINE DOES NOT PRINT

CLIENT APPROVAL CHECKLIST

☐ Dieline ☐ Copy ☐ Colors ☐ UPC

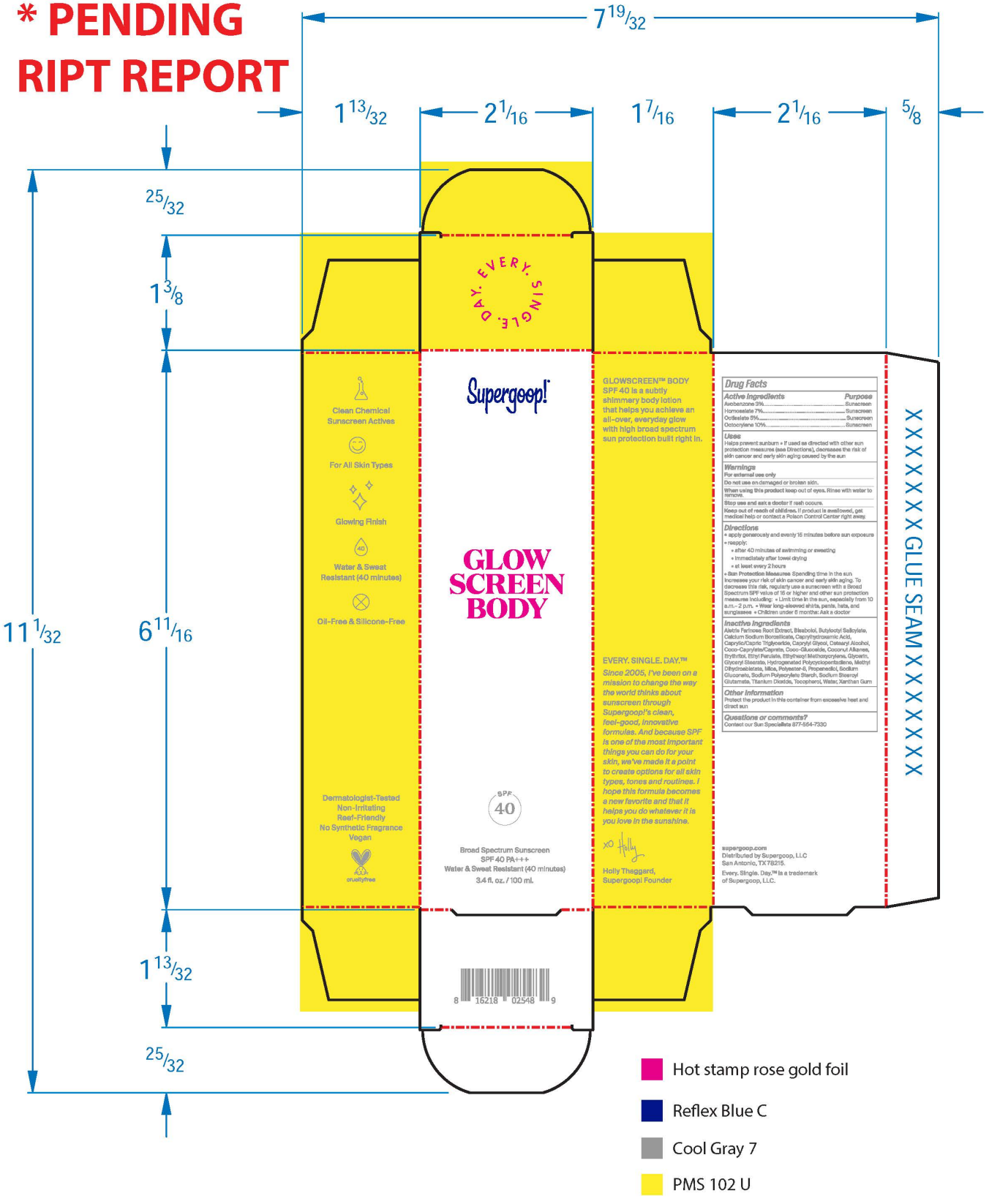
☐ Photography ☐ Illustration ☐ Eye Mark

client signature

date

IMPORTANT: This art has been checked and proofed for accuracy. It is the responsibility of the client to make all final approvals before the release of this art. PLEASE DOUBLE CHECK FOR ACCURACY.

*** PENDING RIPT REPORT**



GLOW SCREEN BODY BROAD SPECTRUM SUNSCREEN SPF 40

avobenzone, homosalate, octisalate, octocrylene cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75936-350
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PROPANEDIOL (UNII: 5965N8W85T)	
WATER (UNII: 059QF0KO0R)	
ERYTHRITOL (UNII: RA96B954X6)	
ETHYL FERULATE (UNII: 5B8915UELW)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ALETRIS FARINOSA ROOT (UNII: O021JGR97X)	
LEVOMENOL (UNII: 24WE03BX2T)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
COCO GLUCOSIDE (UNII: ICS790225B)	
COCONUT ALKANES (UNII: 1E5KJY107T)	
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
MICA (UNII: V8A1AW0880)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
TOCOPHEROL (UNII: R0ZB2556P8)	
POLYESTER-8 (1400 MW, CYANODIPHENYLPROPENOYL CAPPED) (UNII: T9296U138P)	
XANTHAN GUM (UNII: TTV12P4NEE)	
METHYL DIHYDROABIETATE (UNII: 7666FJ0J9F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75936-350-01	100 mL in 1 TUBE; Type 0: Not a Combination Product	01/20/2021	

2	NDC:75936-350-02	10 mL in 1 TUBE; Type 0: Not a Combination Product	01/20/2021	
3	NDC:75936-350-03	15 mL in 1 TUBE; Type 0: Not a Combination Product	01/20/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/20/2021	

Labeler - Supergoop, LLC (117061743)

Revised: 10/2022 Supergoop, LLC