ULTRA VIOLETTE SEQUIN SUPREME SPF 50- octocrylene, homosalate, octisalate, avobenzone lotion
Grace and Fire USA Inc.

ACTIVE INGREDIENTS

AVOBENZONE 3.0%

HOMOSALATE 7.0%

OCTISALATE 5.0%

OCTOCRYLENE 8.0%

Purpose

SUNSCREEN

USE

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 SECTION

For External Use Only

Do not use

on damaged or broken skin

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 swallowed, get medical help or contact a Poison Control Center right away

Directions

Directions for Sunscreen Use

- apply liberally 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, especally from 10 a.m. - 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months of age, Ask a doctor

Inactive Ingredients

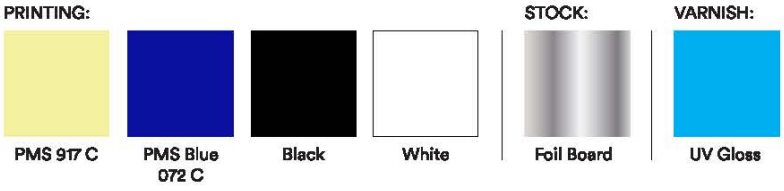
Water, Alcohol Denat., Butylene Glycol, Copernicia Cerifera (Carnauba) Wax, Glycerin, Diisopropyl Adipate, Mica, Cetearyl Alcohol, Caprylyl Methicone, Isododecane, Dimethicone, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Ceteareth-20, Nylon-6/12, Oryza Sativa (Rice) Bran Wax, Polysorbate 60, Saccharide Isomerate, Squalane, Carnosine, Tocopheryl Acetate, Terminalia Ferdinandiana (Kakadu Plum) Fruit Extract, Silica, PEG-40 Stearate, Citric Acid, Glyceryl Caprylate, Fragrance, Disodium EDTA, Pentylene Glycol, Caprylhydroxamic Acid, Triethanolamine, Lactic Acid, Sorbitan Isostearate, Sodium Citrate, Titanium Dioxide (CI 77891), Iron Oxides (CI 77492), Iron Oxides (CI 77491), Iron Oxides (CI 77499).

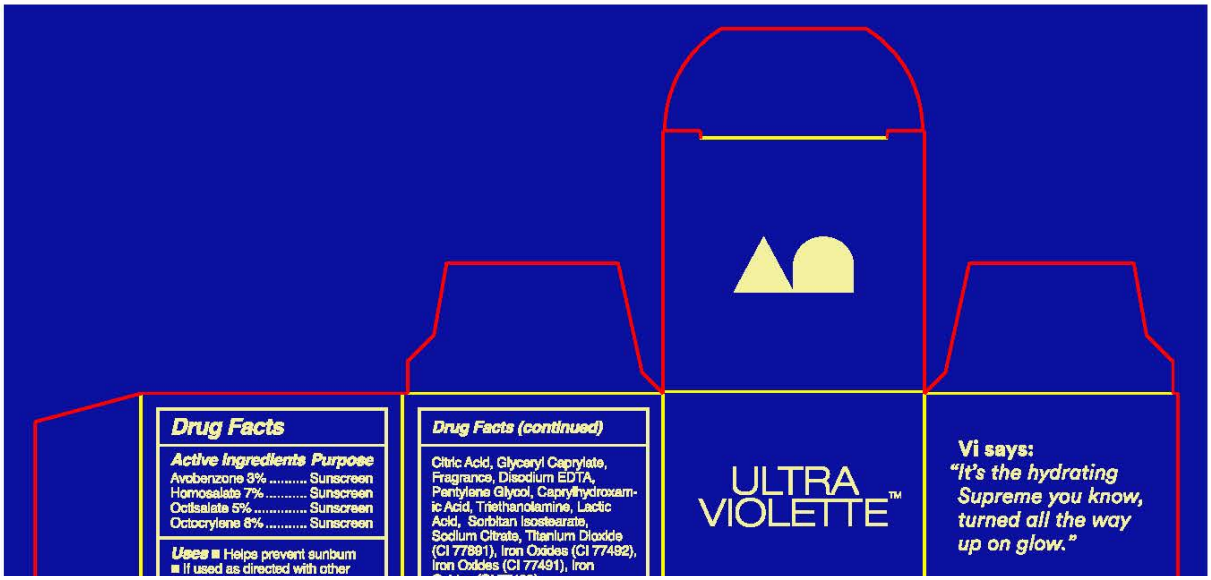
Other Information

Other information protect the product in this container from excessive heat and direct sun

PRINCIPLE DISPLAY PANEL.

SEQUIN SCREEN
USA 50ML, CARTON ART
SES50C1US – FA3





ULTRA

VIOLETTE

SEQUIN SUPREME SPF 50

GLOW -BOOSTING

PRIMING SUNSCREEN

UV

FOR A DEWY

HYDRATED FINISH

Broad Spectrum SPF 50

Sunscreen PA++++

50ml 1.7fl.oz

ULTRA VIOLETTE SEQUIN SUPREME SPF 50

octocrylene, homosalate, octisalate, avobenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84803-107
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	80 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
HYDRATED SILICA (UNII: Y6O7T4G8P9)	0.25 mg in 1 mL
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	0.021 mg in 1 mL
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	1.5 mg in 1 mL
T-BUTYL ALCOHOL (UNII: MD83SFE959)	0.005 mg in 1 mL
POLYSORBATE 60 (UNII: CAL22UVI4M)	0.07 mg in 1 mL
PENTYLENE GLYCOL (UNII: 50C1307PZG)	0.075 mg in 1 mL
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	0.05 mg in 1 mL
CETEARETH-20 (UNII: YRC528SWUY)	0.5 mg in 1 mL
COPERNICIA CERIFERA (CARNAUBA) WAX (UNII: R12CBM0EIZ)	1.9 mg in 1 mL
PEG-40 STEARATE (UNII: ECU18C66Q7)	0.25 mg in 1 mL
CI 77492 (UNII: EX438O2MRT)	0.03 mg in 1 mL
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	0.1 mg in 1 mL
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	2 mg in 1 mL
CARNOSINE (UNII: 8HO6PVN24W)	0.03 mg in 1 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	0.58 mg in 1 mL
ORYZA SATIVA (RICE) BRAN (UNII: R60QEP13IC)	0.1 mg in 1 mL
WATER (UNII: 059QF0KO0R)	19.25 mg in 1 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.77 mg in 1 mL
DIISOPROPYL ADIPATE (UNII: P7E6YFV72X)	1.5 mg in 1 mL
KAKADU PLUM (UNII: 0ZQ1D2FDLI)	0.02 mg in 1 mL
CITRIC ACID (UNII: 2968PHW8QP)	0.003 mg in 1 mL

HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	0.51 mg in 1 mL
ALCOHOL (UNII: 3K9958V90M)	2.49 mg in 1 mL
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	1 mg in 1 mL
ISODODECANE (UNII: A8289P68Y2)	1 mg in 1 mL
DIMETHICONE (UNII: 92RU3N3Y1O)	0.75 mg in 1 mL
NYLON 612 (18000 MW) (UNII: X288170XH1)	0.5 mg in 1 mL
LACTIC ACID (UNII: 33X04XA5AT)	0.025 mg in 1 mL
CI 77491 (UNII: 1K09F3G675)	0.008 mg in 1 mL
CI 77499 (UNII: XM0M87F357)	0.0035 mg in 1 mL
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	0.025 mg in 1 mL
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	1 mg in 1 mL
MICA (UNII: V8A1AW0880)	1.42 mg in 1 mL
GLYCERYL CAPRYLATE (UNII: TM2TZD4G4A)	0.375 mg in 1 mL
SQUALANE (UNII: GW89575KF9)	0.35 mg in 1 mL
SACCHARIDE ISOMERATE (UNII: W8K377W98I)	0.31 mg in 1 mL
FRAGRANCE 13576 (UNII: 5EM498GW35)	0.075 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84803-107-50	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2025	
2	NDC:84803-107-15	15 mL in 1 TUBE; Type 0: Not a Combination Product	11/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/01/2025	

Labeler - Grace and Fire USA Inc. (119357605)

Revised: 10/2025 Grace and Fire USA Inc.