

**ULTRA VIOLETTE FUTURE SCREEN BROAD SPECTRUM SPF 50- zinc  
oxide lotion  
Grace and Fire USA Inc.**

-----

**ACTIVE INGREDIENTS**

ZINC OXIDE 20%

**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

**FOR EXTERNAL USE ONLY**

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

**Directions**

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

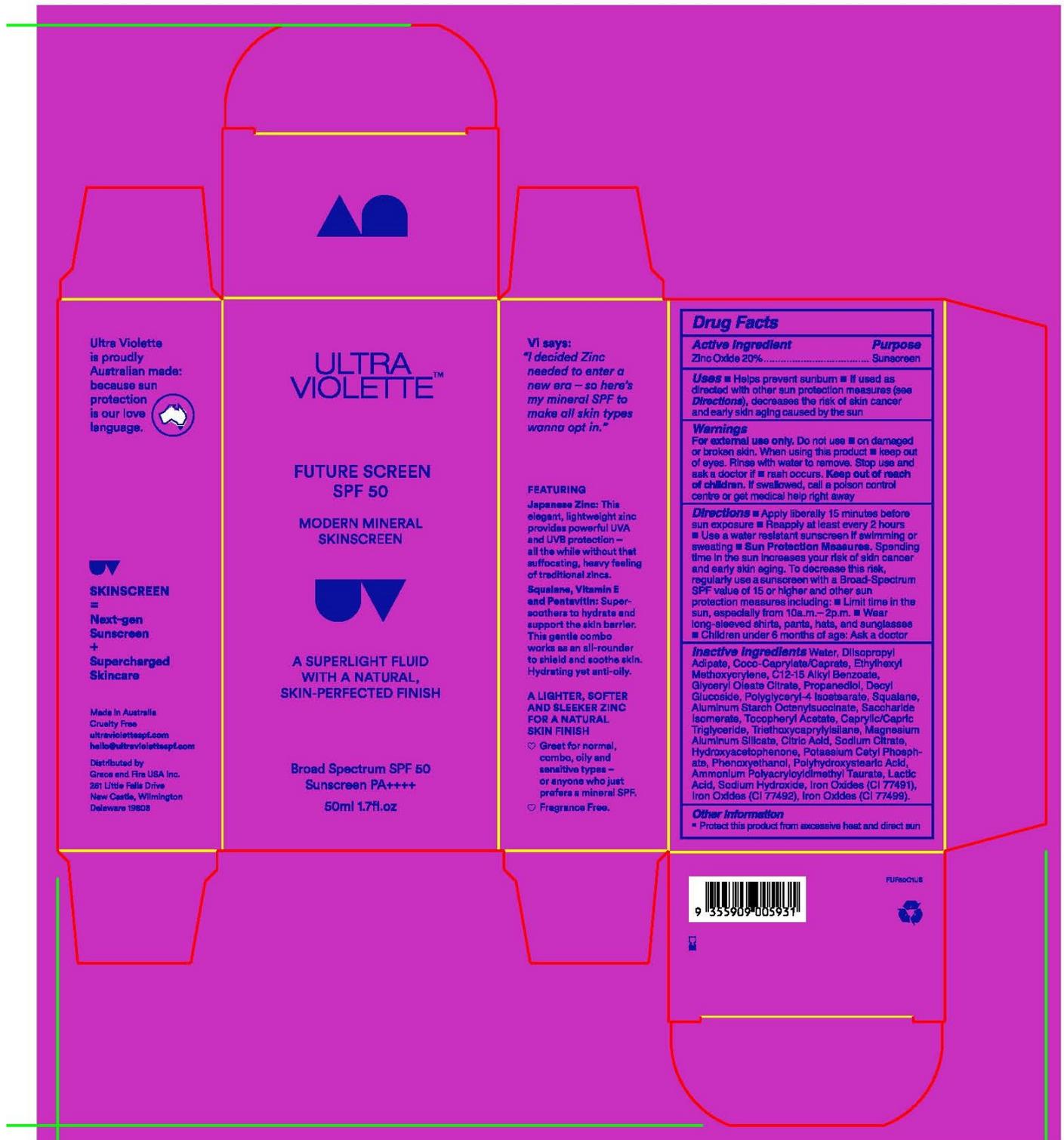
**Inactive Ingredients**

Water, Diisopropyl Adipate, Coco-Caprylate/Caprates, Ethylhexyl Methoxycrylene, C12-15 Alkyl Benzoate, Glyceryl Oleate Citrate, Propanediol, Decyl Glucoside, Polyglyceryl-4 Isostearate, Squalane, Aluminum Starch Octenylsuccinate, Saccharide Isomerate, Tocopheryl Acetate, Caprylic/Capric Triglyceride, Triethoxycaprylylsilane, Magnesium Aluminum Silicate, Citric Acid, Sodium Citrate, Hydroxyacetophenone, Potassium Cetyl Phosphate, Phenoxyethanol, Polyhydroxystearic Acid, Ammonium Polyacryloyldimethyl Taurate, Lactic Acid, Sodium Hydroxide, Iron Oxides (CI 77491), Iron Oxides (CI 77492), Iron Oxides (CI 77499)

### **Other Information**

Protect this product from excessive heat and direct sun

### **Principal Display Panel**



Ultra Violette  
 Future Screen Broad Spectrum SPF 50  
 1.7 FL. OZ. (50 mL)

<p><b>ULTRA VIOLETTE FUTURE SCREEN BROAD SPECTRUM SPF 50</b>          zinc oxide lotion</p>
<p><b>Product Information</b></p>

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84803-103
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	200 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
C12-15 ALKYL BENZOATE (UNII: A9EJ3J61HQ)	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
CI 77499 (UNII: XM0M87F357)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SQUALANE (UNII: GW89575KF9)	
CI 77492 (UNII: EX438O2MRT)	
LACTIC ACID (UNII: 33X04XA5AT)	
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
WATER (UNII: 059QF0KO0R)	
GLYCERYL OLEATE CITRATE (UNII: NLE5KIG74K)	
SACCHARIDE ISOMERATE (UNII: W8K377W98I)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
DIISOPROPYL ADIPATE (UNII: P7E6YFV72X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
PROPANEDIOL (UNII: 5965N8W85T)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
CITRIC ACID (UNII: 2968PHW8QP)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (UNII: F01RIY4371)	
CI 77491 (UNII: 1K09F3G675)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84803-103-50	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2024	
2	NDC:84803-103-15	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2024	
3	NDC:84803-103-21	20 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/19/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/01/2024	

**Labeler** - Grace and Fire USA Inc. (119357605)

Revised: 1/2026

Grace and Fire USA Inc.