

**BIOSSANCE SQUALANE AND DAILY MINERAL SUNSCREEN BROAD SPECTRUM
SPF 50- titanium dioxide, zinc oxide cream
THG Beauty USA LLC**

Titanium Dioxide 5.14%

Zinc Oxide 5.74%

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 exposure to the sun.

For external use only.

Do not use on damaged or broken skin.

Stop use and ask a doctor if rash occurs.

When using this product keep out of eyes. Rinse with water to remove.

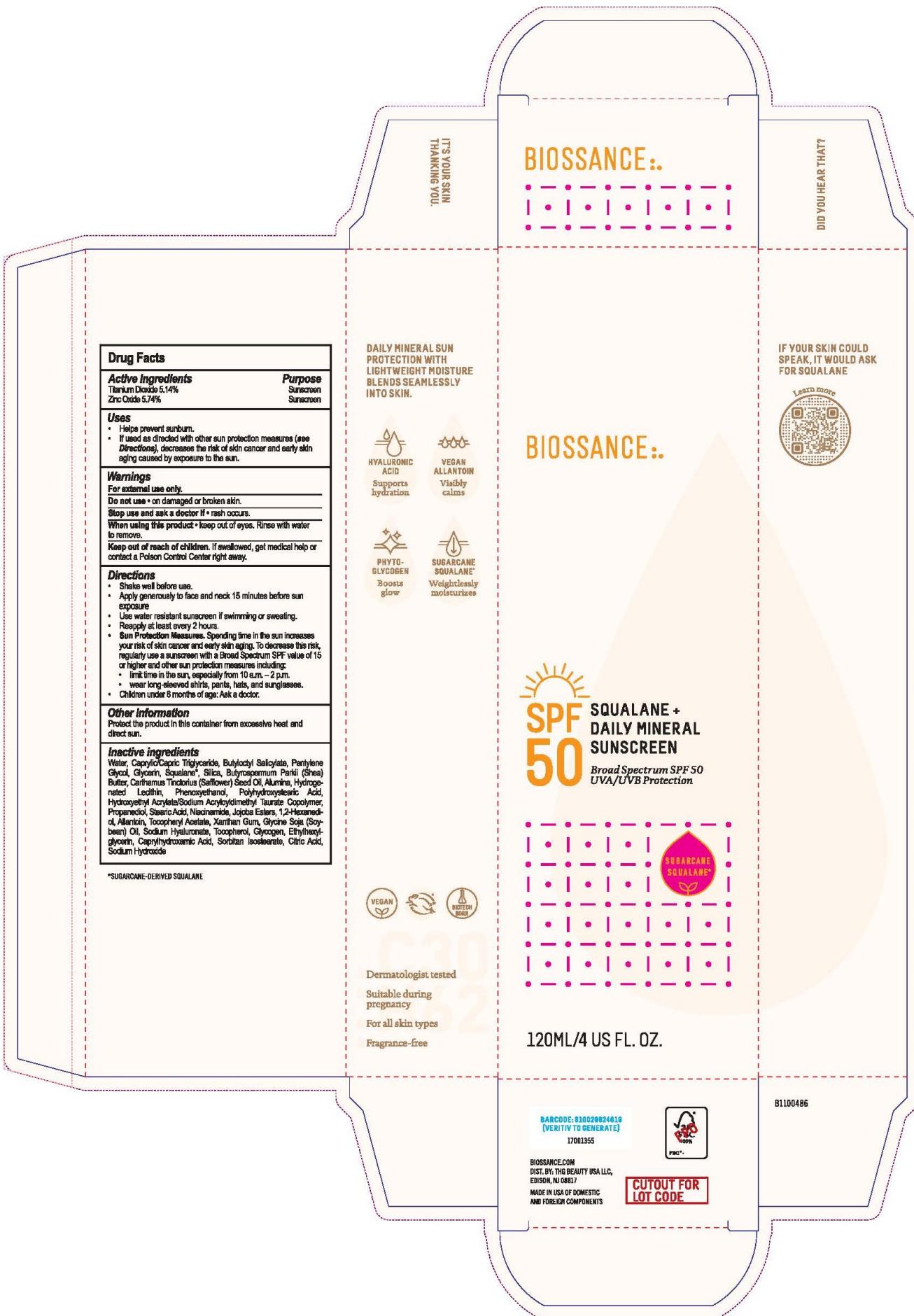
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Centre right away.

- Shake well before use.
- Apply generously to face and neck 15 minutes before sun exposure.
- Use water resistant sunscreen if swimming or sweating.
- Reapply 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 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.

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

WATER, CAPRYLIC/CAPRIC TRIGLYCERIDE, BUTYLOCTYL SALICYLATE, PENTYLENE GLYCOL, GLYCERIN, SQUALANE, SILICA, BUTYROSPERMUM PARKII (SHEA) BUTTER, CARTHAMUS TINCTORIUS (SAFFLOWER) SEED OIL, ALUMINA, HYDROGENATED LECITHIN, PHENOXYETHANOL, POLYHYDROXYSTEARIC ACID, HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER, PROPANEDIOL, STEARIC ACID, NIACINAMIDE, JOJOBA ESTERS, 1,2-HEXANEDIOL, ALLANTOIN, TOCOPHERYL ACETATE, XANTHAN GUM, GLYCINE SOJA (SOYBEAN) OIL, SODIUM HYALURONATE, TOCOPHEROL, GLYCOGEN, ETHYLHEXYLGLYCERIN, CAPRYLHYDROXAMIC ACID, SORBITAN ISOSTEARATE, CITRIC ACID, SODIUM HYDROXIDE







BIOSSANCE SQUALANE AND DAILY MINERAL SUNSCREEN BROAD SPECTRUM SPF 50

titanium dioxide, zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84448-801
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	5.74 g in 100 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	5.1359 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PROPANEDIOL (UNII: 5965N8W85T)	
NIACINAMIDE (UNII: 25X51I8RD4)	
HYDROGENATED SOYBEAN LECITHIN (UNII: H1109Z9J4N)	
ALUMINA (UNII: LMI26O6933)	
SILICA (UNII: ETJ7Z6XBU4)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155WL9Y)	
GLYCERIN (UNII: PDC6A3C00X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	
JOJOBA OIL, RANDOMIZED (UNII: 7F0EV20QYL)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
SQUALANE (UNII: GW89575KF9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CARTHAMUS TINCTORIUS (SAFFLOWER) SEED OIL (UNII: 65UEH262IS)	
ALLANTOIN (UNII: 344S277G0Z)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SOYBEAN OIL (UNII: 241ATL177A)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
TOCOPHEROL (UNII: R0ZB2556P8)	
GLYCOGEN (UNII: 309GSC92U1)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
CITRIC ACID (UNII: 2968PHW8QP)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84448-801-12	1 in 1 CARTON	01/01/2026	
1		120 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:84448-801-60	1 in 1 CARTON	01/01/2026	
2		60 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:84448-801-30	1 in 1 CARTON	01/01/2026	
3		30 mL in 1 TUBE; 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	01/01/2026	

Labeler - THG Beauty USA LLC (015792160)

Revised: 11/2025

THG Beauty USA LLC