

**ANUA ZERO-CAST MOISTURIZING FINISH SUNSCREEN SPF 50- avobenzone, homosalate, octisalate, octocrylene lotion
ENGLEWOOD LAB, INC.**

Anua Zero-Cast Moisturizing Finish Sunscreen SPF 50

Drug Facts

Active Ingredients

Avobenzone 3.0%
Homosalate 7.0%
Octisalate 5.0%
Octocrylene 5.0%

Purpose

Sunscreen

Uses

- 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

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

Keep out of reach of children.

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

Directions

- For sunscreen use:
- Apply liberally 15 minutes before sun exposure
- Reapply:
- after 80 minutes of swimming or sweating

- immediately after towel drying
- at least every 2 hours
- Children under 6 months: Ask a doctor
- **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

Other information

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

Inactive ingredients

Water (Aqua), Butyloctyl Salicylate, Glycerin, Dimethicone, Poly C10-30 Alkyl Acrylate, VP/Eicosene Copolymer, Cetearyl Alcohol, Niacinamide, Sodium Stearoyl Glutamate, Ammonium Polyacryloyldimethyl Taurate, Hydroxyacetophenone, Phenoxyethanol, Tocopheryl Acetate, Panthenol, Pentaerythrityl Tetra-Di-T-Butyl Hydroxyhydrocinnamate, Xanthan Gum, 1,2-Hexanediol, Sodium Hyaluronate

Questions?

+1-888-899-9083 Monday-Friday (9 a.m.-5 p.m. EST) www.anua.us

Package Labeling:

Anua

Anua

Zero-cast™

ANUA ZERO-CAST MOISTURIZING
FINISH SUNSCREEN
ANUA ZERO-CAST ÉCRAN
SOLAIRE FINI VELOUTÉ

SPF 50

Broad Spectrum SPF 50
Water Resistant (80 minutes)

50 ML / 1.69 FL. OZ.

ANUA ZERO-CAST
MOISTURIZING FINISH
SUNSCREEN

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Drug Facts (continued)

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Questions?

+1-888-899-9063
Monday-Friday (9 a.m.-5 p.m. EST)
www.anualus



Manufactured by
ENGLEWOOD LAB, INC.

Distributed by The Founders Inc.,
operating as ANUA INC. in the U.S.
Headquarters: 8F, 10F, Parnas Tower,
521, Teheran-ro, Gangnam-gu, Seoul,
06164, Republic of Korea
US Branch: 222 Pacific Coast Hwy,
#10-145, El Segundo, CA 90245,
United States

Made in USA



X004KJHHP9
Anua Zero-Cast Moistur...screen
(50ml/1.69fl.oz) New



ANUA ZERO-CAST MOISTURIZING FINISH SUNSCREEN SPF 50

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:14268-169
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) (ETHYLHEXYL SALICYLATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	50 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
NIACINAMIDE (UNII: 25X51I8RD4)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (UNII: F01RIY4371)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	

PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PANTHENOL (UNII: WW9CM0O67Z)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
XANTHAN GUM (UNII: TTV12P4NEE)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:14268-169-50	50 mL in 1 TUBE; Type 0: Not a Combination Product	04/10/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M	04/10/2025	

Labeler - ENGLEWOOD LAB, INC. (172198223)

Establishment

Name	Address	ID/FEI	Business Operations
ENGLEWOOD LAB, INC.		172198223	manufacture(14268-169) , label(14268-169)

Revised: 4/2025

ENGLEWOOD LAB, INC.