

NATURIUM DEW GLOW TINTED MOISTURIZER SPF50-MEDIUM- homosalate, octisalate, avobenzone lotion
e.l.f. Cosmetics, Inc.

Naturium Dew-Glow Tinted Moisturizer SPF50-medium

Drug Facts

Avobenzone 3.0%

Homosalate 10.0%

Octisalate 5.0%

Sunscreen

Uses

Helps prevent sunburn.

If used as directed with other sun protection measures (see **Directions**), decreases the risk of skin cancer and early signs of aging caused by the sun.

For external use only.

on damaged or broken skin.

keep out of eyes. Rinse with water to remove.

if product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

For sunscreen use:

Apply generously and evenly 15 minutes before sun exposure.

Apply to all skin exposed to the sun.

Use a water-resistant sunscreen if swimming or sweating.

Reapply at least every 2 hours.

Sun Protection Measures: Spending time in the sun increases risk of skin cancer and early 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 10:00 a.m. - 2:00 p.m.

Wear long sleeved shirts, pants, hats, and sunglasses.

Children under 6 months of age: Ask a doctor.

Other Information

Protect this product from excessive heat and direct sun.

Aqua/Water/Eau, Butyloctyl Salicylate, C12-15 Alkyl Benzoate, Calcium Sodium Borosilicate, Glycerin, Glyceryl Stearate, Ethylhexyl Methoxycrylene, VP/Eicosene Copolymer, Stearyl Dimethicone, Argania Spinosa Kernel Oil, Tocopheryl Acetate, 1,2-Hexandiol, Niacinamide, Sodium Stearoyl Glutamate, Tocopherol, Cetearyl Olivatate, Bisabolol, 3-O-Ethyl Ascorbic Acid, Sodium Hyaluronate, Squalane, Ammonium Acryloyldimethyltaurate/VP Copolymer, Hydroxyacetophnone, Silica, Octadecene, Sorbitan Olivatate, Sorbitan Oleate, Caprylyl/Capryl Glucoside, Xanthan Gum, Sodium Acrylate/Sodium Acryloyldimethyltaurate/VP Copolymer, Hydroxyacetophenone, Silica, Octadecene, Sorbitan Olivatate, Sorbitan Oleate, Caprylyl/Capryl Glucoside, Xanthan Gum, Sodium Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Trisodium Ethylenediamine Disuccinate, Polyisobutene, Citric Acid, Sodium Hydroxide

May Contain: Iron Oxide (CI 77491), Iron Oxide (CI 77492), Iron Oxide (CI 77499)

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homosalate, octisalate, avobenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76354-132
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 50 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 50 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 50 mL

Inactive Ingredients

Ingredient Name	Strength
NIACINAMIDE (UNII: 25X51I8RD4)	
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
TOCOPHEROL (UNII: R0ZB2556P8)	
3-O-ETHYL ASCORBIC ACID (UNII: 6MW60CB71P)	
OCTADECENE (UNII: H5ZUQ6V4AK)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYISOBUTYLENE (400000 MW) (UNII: X9N69O5R5X)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
BROWN IRON OXIDE (UNII: 1N032N7MFO)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
HYALURONIC ACID (UNII: S270N0TRQY)	
SODIUM ACRYLATE/SODIUM ACRYLOYLDIMETHYLTAURATE COPOLYMER (4000000 MW) (UNII: 1DXE3F3OZX)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)	
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)	
BOROSILICATE GLASS (UNII: BOJ6T9AR90)	
ARGAN OIL (UNII: 4V59G5UW9X)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
SQUALANE (UNII: GW89575KF9)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	

TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-132-01	1 in 1 CARTON	12/27/2023	
1		50 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	12/27/2023	

Labeler - e.l.f. Cosmetics, Inc. (093902816)

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