E L F WHOA GLOW BROAD SPECTRUM SPF 30 SUNSCREEN SHEER BRONZE SHIMMER SUNBURST- avobenzone, homosalate, octisalate, octocrylene cream
e.l.f. Cosmetics, Inc

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e.l.f. Whoa Glow Broad Spectrum SPF 30 Sunscreen, Sheer Bronze Shimmer Sunburst

Drug Facts

Active Ingredients

Avobenzone 3.0%

Homosalate 7.3%

Octisalate 4.0%

Octocrylene 8.5%

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

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

- Apply liberally 15 minutes before sun exposure. Reapply at least every 2 hours.
- Use 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 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

Other Information

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

Inactive Ingredients

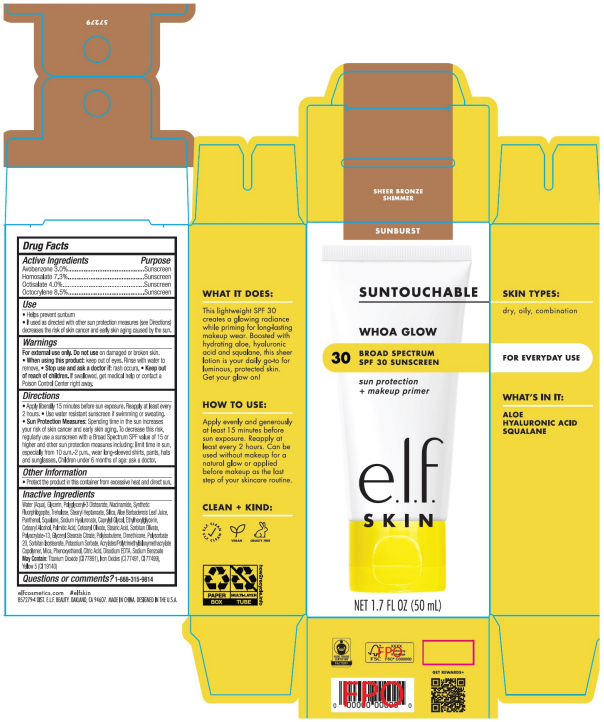
Water (Aqua), Glycerin, Polyglyceryl-3 Distearate, Niacinamide, Synthetic Fluorphogopite, Trehalose, Stearyl Heptanoate, Silica, Aloe Barbadensis Leaf Juice, Panthenol, Squalane, Sodium Hyaluronate, Caprylyl Glycol, Ethylhexylglycerin, Cetearyl Alcohol, Palmitic Acid, Cetearyl Olivate, Stearic Acid, Sorbitan Olivate, Polyacrylate-13, Glyceryl Stearate Citrate, Polyisobutene, Dimethicone, Polysorbate 20, Sorbitan Isostearate, Potassium Sorbate, Acrylates/Polytrimethylsiloxymethacrylate Copolymer, Mica, Phenoxyethanol, Citric Acid, Disodium EDTA, Sodium Benzoate **May Contain:**Titanium Dioxide (CI 77891), Iron Oxides (CI 77491, CI 77499), Yellow 5 (CI 19140)

Questions or comments?

1-888-315-9814

Package labelling:

REFERENCE



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avobenzone, homosalate, octisalate, octocrylene cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76354-451
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	73 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	40 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	85 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYGLYCERYL-3 DISTEARATE (UNII: ZI1LK470XV)	
NIACINAMIDE (UNII: 25X51I8RD4)	
MAGNESIUM POTASSIUM ALUMINOSILICATE FLUORIDE (UNII: YK3DC63Y5M)	
TREHALOSE (UNII: B8WCK70T7I)	

STEARYL HEPTANOATE (UNII: 2M4UGL1NCN)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PANTHENOL (UNII: WV9CM0O67Z)	
SQUALANE (UNII: GW89575KF9)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PALMITIC ACID (UNII: 2V16EO95H1)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
POLYACRYLATE-13 (UNII: FS2D4T67EA)	
GLYCERYL STEARATE CITRATE (UNII: WH8T92A065)	
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y1O)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
MICA (UNII: V8A1AW0880)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-451-01	50 mL in 1 TUBE; Type 0: Not a Combination Product	07/20/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	07/20/2024	

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

Revised: 7/2024 e.l.f. Cosmetics, Inc