

HALO GLOW SKIN TINT SPF 50 84504- zinc oxide lotion
e.l.f. Cosmetics, Inc.

Halo Glow Skin Tint SPF 50

Drug Facts

Active Ingredients

Zinc Oxide 12%

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

- Apply generously and evenly 15 minutes before sun exposure.
- Reapply:
 - after 80 minutes of swimming or sweating.
 - immediately after towel drying.
 - 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 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: Ask a doctor.

Other Information

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

Inactive ingredients

Water, C12-15 Alkyl Benzoate, Isododecane, Lauryl PEG-8 Dimethicone, Tridecyl Salicylate, Calcium Sodium Borosilicate, Propanediol, Butyloctyl Salicylate, Caprylyl Methicone, Dimethicone, Niacinamide, C13-15 Alkane, Silica, Octyldodecyl Neopentanoate, Polyglyceryl-2 Dipolyhydroxystearate, Polyglyceryl-3 Polyricinoleate, Sodium Chloride, Distearidimonium Hectorite, Phenoxyethanol, Bisabolol, Lecithin, Caprylyl Glycol, Allantoin, Aluminum Hydroxide, Polyglyceryl-4 Diisostearate/Polyhydroxystearate/Sebacate, Ethylhexylglycerin, Hexylene Glycol, Hydrogenated Lecithin, Tetrasodium Glutamate Diacetate, Sodium Hydroxide

May Contain: Titanium Dioxide (CI 77891), Iron Oxides (CI 77491, CI 77492, CI 77499)

Questions or comments?

1-888-315-9814

Packaging Label

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76354-550
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)		ZINC OXIDE	120 mg in 1 mL
Inactive Ingredients			
Ingredient Name			Strength
ISODODECANE (UNII: A8289P68Y2)			
CALCIUM SODIUM BOROSILICATE (UNII: 4MM76N4VMY)			
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)			
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)			
LAURYL PEG-8 DIMETHICONE (300 CPS) (UNII: ELL2U7K8T8)			
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)			
C13-15 ALKANE (UNII: 114P5I43UJ)			
.BETA.-BISABOLOL (UNII: LP618AV2EA)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
CAPRYLYL TRIMETHICONE (UNII: H6HK6E4EB1)			
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)			
WATER (UNII: 059QF0KO0R)			
PROPANEDIOL (UNII: 5965N8W85T)			
FERROSOFERRIC OXIDE (UNII: XM0M87F357)			
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)			
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)			
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)			
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
HEXYLENE GLYCOL (UNII: KEH0A3F75J)			
FERRIC OXIDE RED (UNII: 1K09F3G675)			
TRIDECYL SALICYLATE (UNII: AZQ08K38Z1)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
DIMETHICONE (UNII: 92RU3N3Y1O)			
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
NIACINAMIDE (UNII: 25X51I8RD4)			
POLYGLYCERYL-3 RICINOLEATE (UNII: MZQ63P0N0W)			
CAPRYLYL GLYCOL (UNII: 00YIU5438U)			
ALLANTOIN (UNII: 344S277G0Z)			
HYDROGENATED SOYBEAN LECITHIN (UNII: H1109Z9J4N)			
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)			
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)			
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)			
Packaging			
		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-550-01	1 in 1 CARTON	05/05/2025	
1		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	05/05/2025	

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