

GLO SKIN BEAUTY OIL FREE SPF 40- glo skin beauty oil free spf 40 lotion
Hayden Caleel LLC

DRUG FACTS

Active Ingredients

Octinoxate 7.5%

Octisalate 2.5%

Zinc Oxide 7.5%

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 15 minutes before sun exposure
- reapply at least every 2 hours
- use a 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 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

Inactive ingredients

Aqua/Water/Eau, Isopropyl Palmitate, Ethylhexyl Stearate, Ethylhexyl Isononanoate, Cyclopentasiloxane, Cetearyl Glucoside, Dimethicone, Phenoxyethanol, Glycereth-26, Oleth-3 Phosphate, Allantoin, Triethoxycaprylylsilane, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Polyisobutene, Panthenol, Butylene Glycol, PEG-7 Trimethylolpropane Coconut Ether, Sodium Hyaluronate, Tocopheryl Acetate, Ascorbyl Palmitate, Iodopropynyl Butylcarbamate

Other Information

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

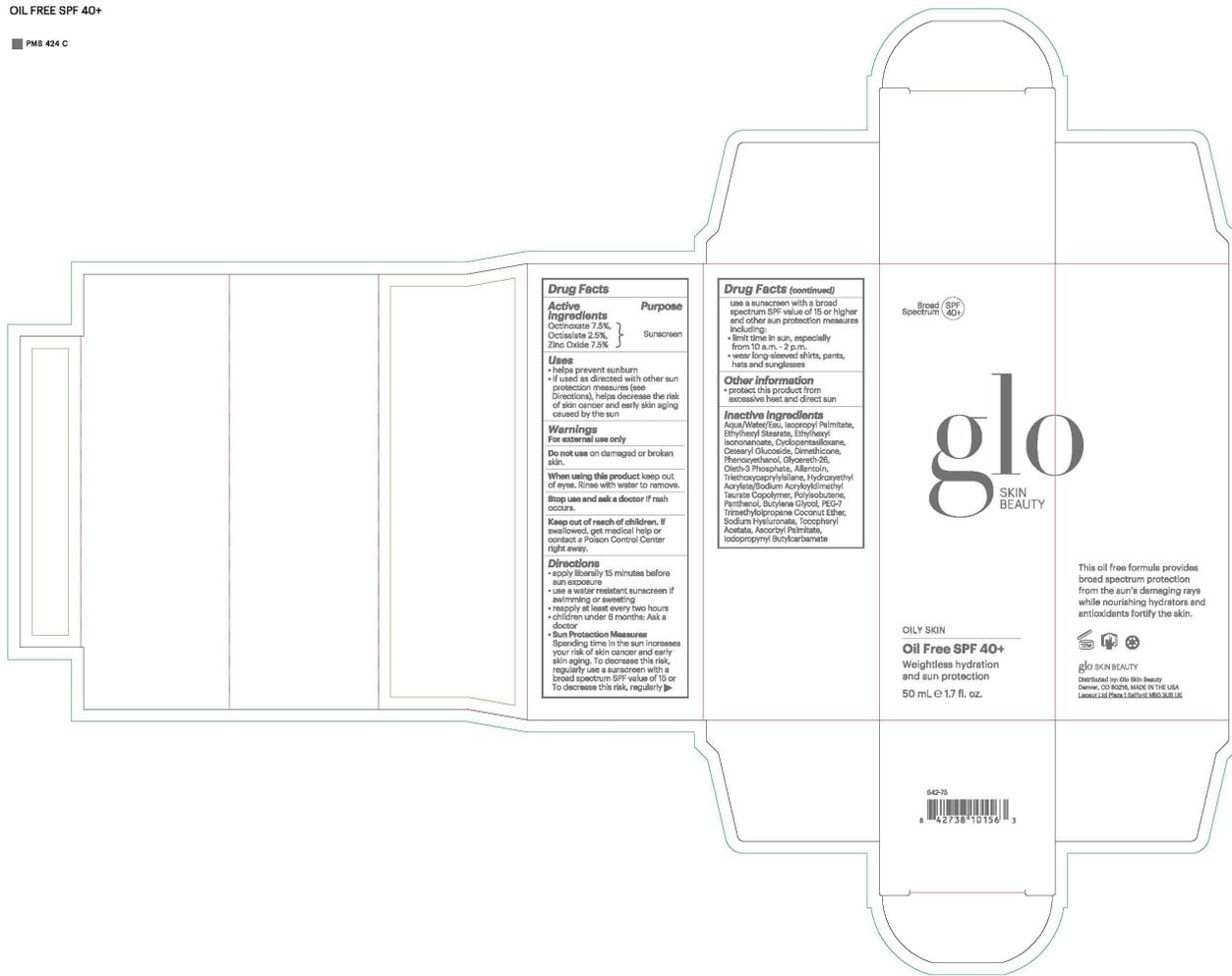
Keep out of reach of children

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Purpose

Sunscreen

Principal Display Panel



GLO SKIN BEAUTY OIL FREE SPF 40

glo skin beauty oil free spf 40 lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60541-1000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	2.5 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	7.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	

ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)
ETHYLHEXYL ISONONANOATE (UNII: I6KB4GE3K4)
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)
DIMETHICONE (UNII: 92RU3N3Y1O)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
GLYCERETH-26 (UNII: NNE56F2N14)
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)
ALLANTOIN (UNII: 344S277G0Z)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
PANTHENOL (UNII: WW9CM0067Z)
ASCORBYL PALMITATE (UNII: QN83US2B0N)
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60541-1000-5	1 in 1 CARTON	12/31/2017	
1		50 g 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	12/31/2017	

Labeler - Hayden Caleel LLC (011367468)

Revised: 1/2025

Hayden Caleel LLC