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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

Active Ingredients

Octinoxate 7.5%

Octisalate 2.5%

Zinc Oxie 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 docor if rash occurs

Keep out of reach of children. If swalled, 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

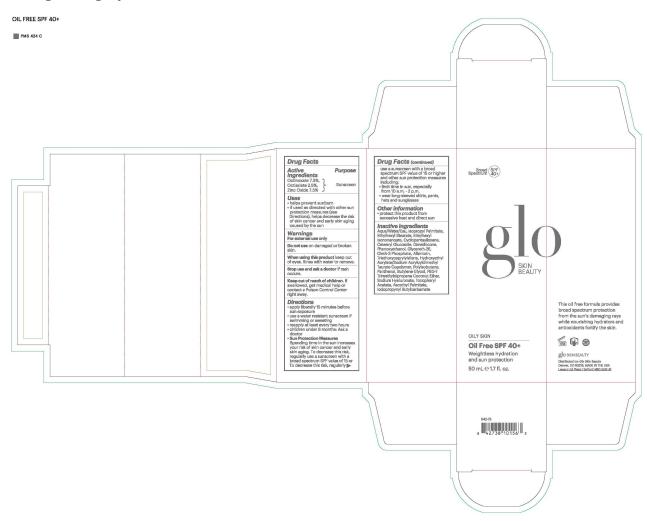
Inactive ingredients

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

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

Principal Display Panel



GLO SKIN BEAUTY OIL FREE SPF 40

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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: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W)	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: 8 CRQ2TH63M)		
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)		
ETHYLHEXYL ISONONANOATE (UNII: 16 KB4GE3K4)		
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)		
DIMETHICO NE (UNII: 92RU3N3Y1O)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
GLYCERETH-26 (UNII: NNE56F2N14)		
OLETH-3 PHO SPHATE (UNII: 8Q0Z18J1VL)		
ALLANTO IN (UNII: 344S277G0Z)		
TRIETHO XYCAPRYLYLSILANE (UNII: LDC331P08E)		
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)		
PANTHENOL (UNII: WV9CM0O67Z)		
ASCORBYL PALMITATE (UNII: QN83US2B0N)		

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 final	part352	12/31/2017		

Labeler - Hayden Caleel LLC (011367468)

Revised: 12/2017 Hayden Caleel LLC