

**SUN GLAZED SPF 30 SUNSCREEN GLOW OIL- avobenzone, homosalate,
octocrylene oil
EVERYDAY GROUP LIMITED**

Sun Glazed SPF 30 Sunscreen Glow Oil

Drug Facts

Active ingredients

Avobenzone 3% Homosalate 10% Octocrylene 8%

Purpose

Sunscreen

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see) decreases the risk of skin cancer and early skin aging caused by the sun ***Directions***

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

- use at least 3 to 4 pumps
- apply liberally 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 the 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. to 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months of age: Ask a doctor

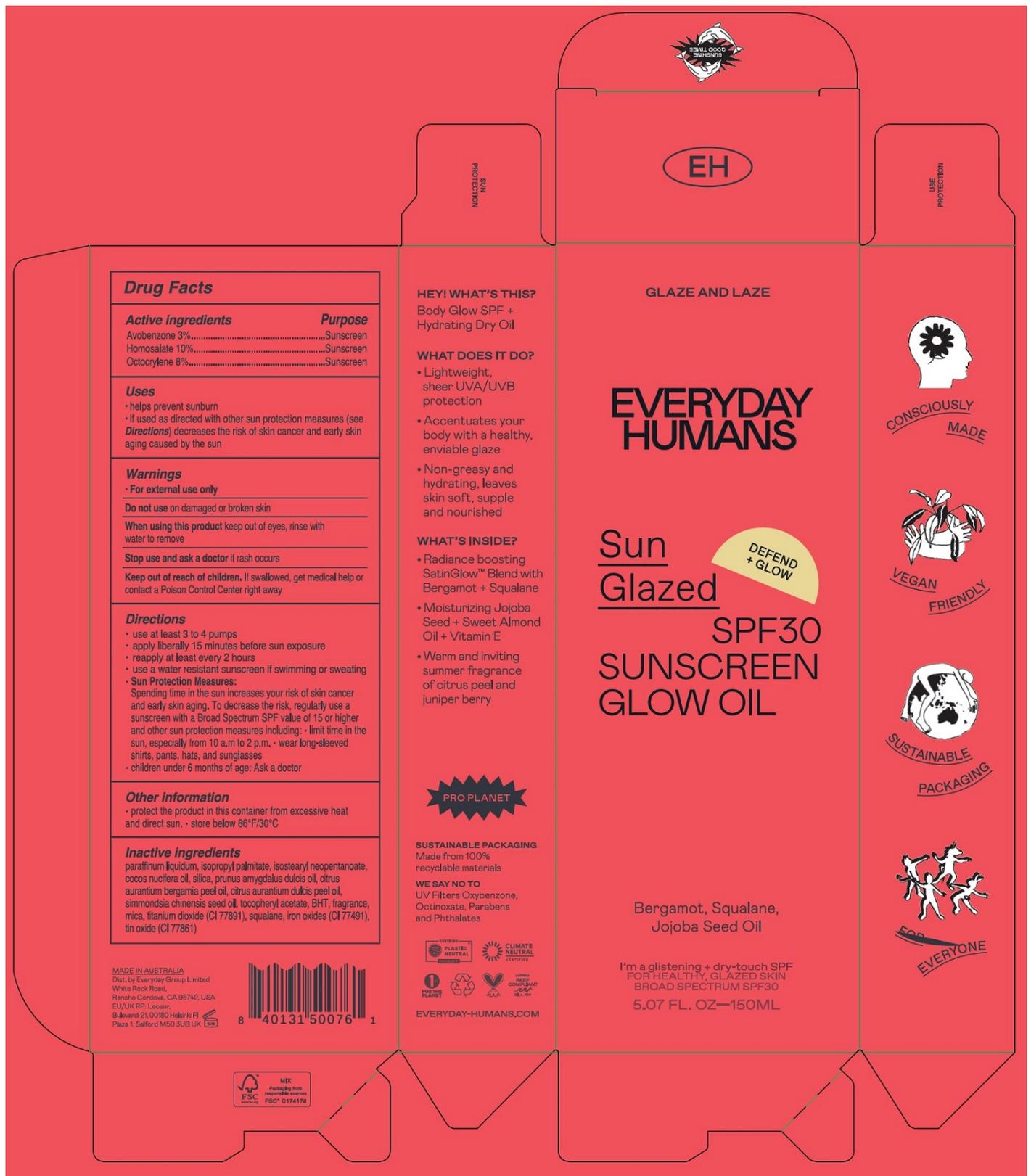
Other information

- protect the product in this container from excessive heat and direct sun.
- store below 86°F/30° C

Inactive ingredients

paraffinum liquidum, isopropyl palmitate, isosterayl neopentanoate, cocos nucifera oil, silica, prunus amygdalus dulcis oil, citrus aurantium bergamia peel oil, citrus aurantium dulcis peel oil, simmondsia chinensis seed oil, tocopheryl acetate, BHT, fragrance, mica, titanium dioxide (CI 77891), squalane, iron oxides (CI 77491), tin oxide (CI 77861)

Package Labeling:



SUN GLAZED SPF 30 SUNSCREEN GLOW OIL
 avobenzone, homosalate, octocrylene oil

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72098-007
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	100 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	80 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ISOSTEARYL NEOPENTANOATE (UNII: 411THY156Q)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
JOJOBA OIL (UNII: 724GKU717M)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
MICA (UNII: V8A1AW0880)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SQUALANE (UNII: GW89575KF9)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STANNIC OXIDE (UNII: KM7N50LOS6)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72098-007-01	1 in 1 BOX	04/01/2023	
1		150 mL in 1 BOTTLE; 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	04/01/2023	

Labeler - EVERYDAY GROUP LIMITED (664624682)