SHEGLAM DEW AND DONE SKIN TINT WITH SPF 20 SANDALWOOD- zinc oxide cream EVER FAME PTE. LTD.

Sheglam Dew & Done Skin Tint with SPF 20, Sandalwood

Drug Facts

Active ingredient

Zinc Oxide 14.625%

Purpose

Sunscreen

Use

helps prevent sunburn

Warnings

Skin Cancer/Skin Aging Alert: Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to help prevent sunburn, **not skin cancer or early skin aging.**

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 a water resistant sunscreen if swimming or sweating
- children under 6 months of age: Ask a doctor

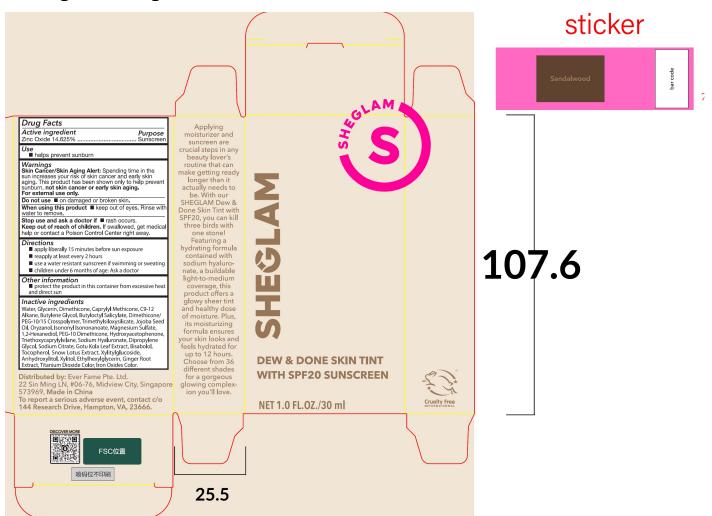
Other information

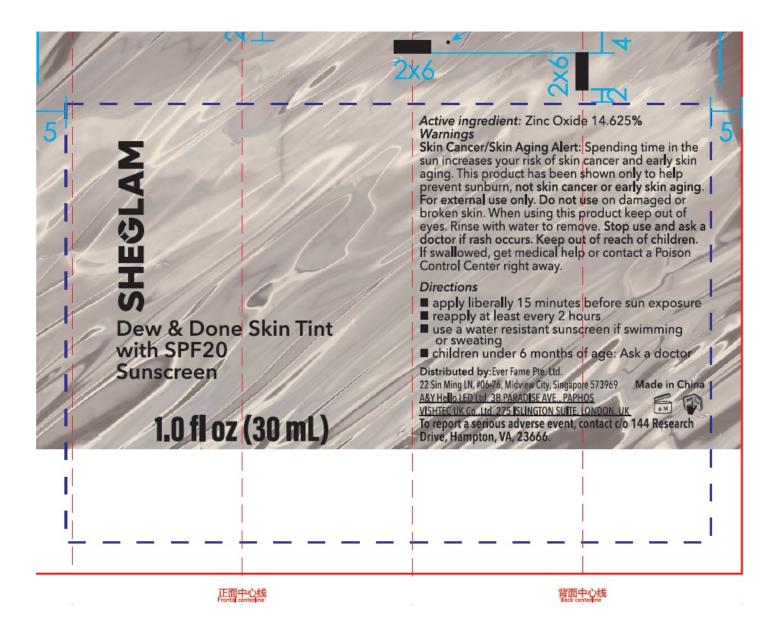
protect the product in this container from excessive heat and direct sun

Inactive ingredients

Water, Glycerin, Dimethicone, Caprylyl Methicone, C9-12 Alkane, Butylene Glycol, Butyloctyl Salicylate, Dimethicone/PEG-10/15 Crosspolymer, Trimethylsiloxysilicate, Jojoba Seed Oil, Oryzanol, Isononyl Isononanoate, Magnesium Sulfate, 1,2-Hexanediol, PEG-10 Dimethicone, Hydroxyacetophenone, Triethoxycaprylylsilane, Sodium Hyaluronate, Dipropylene Glycol, Sodium Citrate, Gotu Kola Leaf Extract, Bisabolol, Tocopherol, Snow Lotus Extract, Xylitylglucoside, Anhydroxylitol, Xylitol, Ethylhexylglycerin, Ginger Root Extract, Titanium Dioxide Color, Iron Oxides Color.

Package Labeling:





SHEGLAM DEW AND DONE SKIN TINT WITH SPF 20 SANDALWOOD

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85068-029
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	146.25 mg in 1 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

DIMETHICONE (UNII: 92RU3N3Y10)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	
C9-12 ALKANE (UNII: 7J5R5W72QM)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
DIMETHICONE/PEG-10/15 CROSSPOLYMER (UNII: 21AS8B1BSS)	
JOJOBA OIL (UNII: 724GKU717M)	
ORYZANOL (UNII: SST9XCL51M)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
MAGNESIUM SULFATE (UNII: DE08037SAB)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
BISABOLOL (UNII: 24WE03BX2T)	
TOCOPHEROL (UNII: R0ZB2556P8)	
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	
XYLITOL (UNII: VCQ006KQ1E)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GINGER (UNII: C5529G5JPQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85068-029- 01	1 in 1 CARTON	05/30/2025	
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	05/30/2025	

Labeler - EVER FAME PTE. LTD. (659974769)

Revised: 1/2025 EVER FAME PTE. LTD.