

SUGARGIRL SET SCREEN SPF 30 SETTING POWDER SUNSCREEN LIGHT- zinc oxide, titanium dioxide powder
I World LLC

Sugargirl Set Screen SPF 30 Setting Powder Sunscreen Light

Drug Facts

Active ingredients

Zinc Oxide 15.18% Titanium Dioxide 9.84%

Purpose

Sunscreen

Uses

Helps prevent sunburn.

Warnings

For external use only.

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.

Do not use

- on damaged or broken skin.
- in the eye area.

When using this product

- keep out of eyes. rinse with water to remove.

Stop use and ask doctor if

- rash occurs.

Keep out of reach of children.

- If product is swallowed get medical help or contact a poison control center right away.

Directions

- Apply generously and evenly 15 minutes before sun exposure.
- Reapply at least every 2 hours.

- Use a water resistant sunscreen if swimming or sweating.
- Children under 6 month: ask a doctor.

Other information

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

Inactive Ingredients

CALCIUM SODIUM BOROSILICATE, SILICA, VINYL DIMETHICONE/METHICONE SILSESQUIOXANE CROSSPOLYMER, LAUROYL LYSINE, TRIETHOXYCAPRYLYLSILANE, ETHYLHEXYLGLYCERIN, CAPRYLYL GLYCOL, POLYHYDROXYSTEARIC ACID, LECITHIN, ALUMINUM HYDROXIDE, ISOPROPYL MYRISTATE, ETHYLHEXYL PALMITATE, ISOSTEARIC ACID, POLYGLYCERYL-3 POLYRICINOLEATE, IRON OXIDES (CI 77492), IRON OXIDES (CI 77491), YELLOW 5 LAKE (CI 19140), IRON OXIDES (CI 77499), YELLOW 6 LAKE (CI 15985).

Questions or comments?

1-212- 244-5170 info@amora.beauty

Package labeling:

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85179-033
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)			ZINC CATION	151.8 mg in 1 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (CI 77891 - UNII:15FIX9V2JP)			TITANIUM DIOXIDE	98.4 mg in 1 g
Inactive Ingredients				
Ingredient Name				Strength
CALCIUM SODIUM BOROSILICATE (UNII: 4MM76N4WMY)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
VINYL DIMETHICONE/METHICONE SILSESQUIOXANE CROSSPOLYMER (UNII: 9NH1UDD2RR)				
LAUROYL LYSINE (UNII: 113171Q70B)				
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
CAPRYLYL GLYCOL (UNII: 00YIU5438U)				
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)				
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)				
ETHYLHEXYL PALMITATE (UNII: 2865993309)				
ISOSTEARIC ACID (UNII: X33R8U0062)				
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)				
FERRIC OXIDE RED (UNII: 1K09F3G675)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
FERROSOFERRIC OXIDE (UNII: XM0M87F357)				
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85179-033-00	4 g in 1 TUBE; Type 0: Not a Combination Product	12/06/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M020	12/06/2025	

Labeler - I World LLC (830590126)

Registrant - I World LLC (830590126)

Establishment			
Name	Address	ID/FEI	Business Operations
Shanghai Chuanyue Biotechnology Co., Ltd.		841882128	manufacture(85179-033)

