WOODSLEEP TALLOW SUNSCREEN- titanium dioxide cream
Shantou Woodsleep Biotechnology Co., Ltd.

---

1. Clean skin and keep it dry.

2. 20-30 minutes before going out, take an appropriate amount of sunscreen and apply it evenly on the face, neck, arms and other parts.

3. Gently massage with fingertips until completely absorbed.

1. Effectively block ultraviolet rays, build a natural protective barrier for the skin, and protect against the sun with peace of mind.

2. The unique formula deeply moisturizes and lasts for a long time, preventing the skin from drying out due to sun exposure, and keeping the skin moisturized and tender.

3. The soothing formula reduces skin irritation, relieves discomfort after sun exposure, and gives gentle care to the skin, suitable for all skin types.

Discontinue use if signs of irritation or rash occur.

Discontinue use if signs of irritation or rash occur.

Please keep out of reach of children. Do not swallow.

Please keep out of reach of children. Do not swallow.Please clean your hands before use to ensure the best results from the product. Discontinue use if signs of irritation or rash occur. Store in a cool and dry place.

Store in a cool and dry place.

AQUA□TITANIUM DIOXIDE□ETHYLHEXYL METHOXYCINNAMATE□ISONONYL ISONONANOATE□GLYCERYL STEARATE□TOCOPHEROL

TALLOW

纸盒
规格尺寸：长5.1X宽5.1X高5.1CM

WOODSLEEP TALLOW SUNSCREEN

titanium dioxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85053-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TALLOW (UNII: 98HPY76U4W) (TALLOW - UNII:98HPY76U4W)	TALLOW	0.012 mg in 60 mg

Inactive Ingredients

Ingredient Name	Strength
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	2.4 mg in 60 mg
TOCOPHEROL (UNII: R0ZB2556P8)	0.012 mg in 60 mg
GLYCERYL STEARATE (UNII: 230OU9XXE4)	1.2 mg in 60 mg
AQUA (UNII: 059QF0KO0R)	35.976 mg in 60 mg
ETHYLHEXYL METHOXYCINNAMATE (UNII: 4Y5P7MUD51)	5.4 mg in 60 mg
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	15 mg in 60 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85053-003-01	60 mg in 1 BOX; Type 0: Not a Combination Product	04/28/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/28/2025	

Labeler - Shantou Woodsleep Biotechnology Co., Ltd. (699296106)

Registrant - Shantou Woodsleep Biotechnology Co., Ltd. (699296106)

Establishment

Name	Address	ID/FEI	Business Operations
Shantou Woodsleep Biotechnology Co., Ltd.		699296106	manufacture(85053-003)

Revised: 4/2025 Shantou Woodsleep Biotechnology Co., Ltd.