

OUHOE SUNSCREEN- titanium dioxide cream
Shantou Ouhoe Technology Co., Ltd.

CENTELLA ASIATICA LEAF EXTRACT□GLYCYRRHIZA URALENSIS (LICORICE) ROOT EXTRACT

1. Efficient sun protection and isolation, comprehensive protection against ultraviolet rays, allowing you to enjoy the sun with peace of mind.
 2. Deeply nourish, the skin will be hydrated and translucent, giving it a natural luster.
 3. Improve skin elasticity, even skin tone, significantly brighten and rejuvenate.
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.

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.

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

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

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纸盒

规格尺寸：长6.6X宽3.2X高11.3CM



OUHOE SUNSCREEN

titanium dioxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85163-010
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
CENTELLA ASIATICA LEAF (UNII: 6810070TYD) (CENTELLA ASIATICA LEAF -		CENTELLA ASIATICA	0.01 mg

UNII:6810070TYD)	LEAF	in 50 mg
GLYCYRRHIZA URALENSIS ROOT (UNII: 42B5YD8F0K) (GLYCYRRHIZA URALENSIS ROOT - UNII:42B5YD8F0K)	GLYCYRRHIZA URALENSIS ROOT	0.01 mg in 50 mg

Inactive Ingredients

Ingredient Name	Strength
NIACINAMIDE (UNII: 25X51I8RD4)	0.01 mg in 50 mg
AQUA (UNII: 059QF0KO0R)	29.97 mg in 50 mg
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	12.5 mg in 50 mg
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	2 mg in 50 mg
ETHYLHEXYL METHOXYCINNAMATE (UNII: 4Y5P7MUD51)	4.5 mg in 50 mg
GLYCERYL STEARATE (UNII: 230OU9XXE4)	1 mg in 50 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85163-010-01	50 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/07/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/07/2025	

Labeler - Shantou Ouhoe Technology Co., Ltd. (617531875)

Registrant - Shantou Ouhoe Technology Co., Ltd. (617531875)

Establishment

Name	Address	ID/FEI	Business Operations
Shantou Ouhoe Technology Co., Ltd.		617531875	manufacture(85163-010)

Revised: 11/2025

Shantou Ouhoe Technology Co., Ltd.