RENEWALUXE SPF 50 SUNSCREEN- renewaluxe spf 50 sunscreen cream
Guangzhou Fantesy Biotechnology Co.,Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

83447-005

Zinc Oxide 21.8%

sunscreen

Uses

Uses Helps prevent sunburn. If used as directed with other sun protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun.

Warnings

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For external use only
Do not use on damaged or broken skin.
When using the 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 Poison Control Center right away.

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Directions

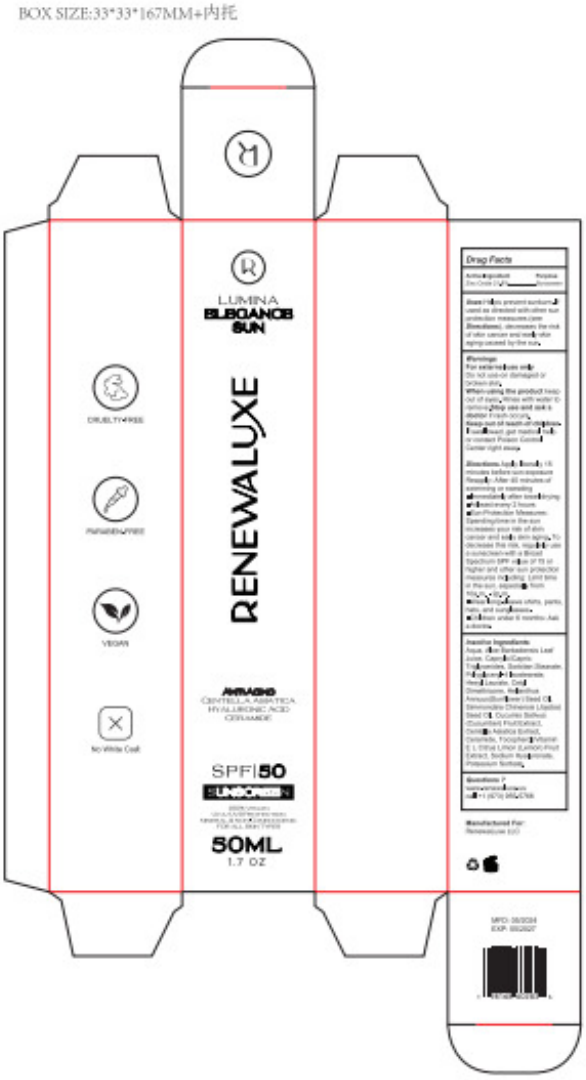
Directions Apply liberally 15 minutes before sun exposure Reapply: After 40 minutes of swimming or sweating
■Immediately after towel drying
■At least every 2 hours
■Sun Protection Measures: Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly 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 10a.m. - 2p.m.
■Wear long-sleeve shirts, pants, hats, and sunglasses.
■Children under 6 months: Ask a doctor.

Other information

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

Inactive Ingredients



No data available

RENEWALUXE SPF 50 SUNSCREEN

renewaluxe spf 50 sunscreen cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83447-005
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	21.8 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	

COCONUT OIL (UNII: Q9L0O73W7L)	
JOJOBA OIL (UNII: 724GKU717M)	
CENTELLA ASIATICA TRITERPENOIDS (UNII: 4YS74Q4G4J)	
CERAMIDE 9 (UNII: 88KCS7120E)	
TOCOPHEROL (UNII: R0ZB2556P8)	
LEMON (UNII: 24RS0A988O)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CETYL DIMETHICONE 45 (UNII: IK315POC44)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83447-005-01	50 mL in 1 BOX; Type 0: Not a Combination Product	06/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/26/2024	

Labeler - Guangzhou Fantesy Biotechnology Co.,Ltd (619047084)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Fantesy Biotechnology Co.,Ltd		619047084	manufacture(83447-005)

Revised: 7/2024 Guangzhou Fantesy Biotechnology Co.,Ltd