

**EVERYRAY SUNCARE SUNSHINE SERUM- everyray suncare sunshine
serum liquid
Guangzhou Fantasy Biotechnology Co.,Ltd**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

83447-001

Active ingredients

Zinc Oxide 25%

Purpose

Sunscreen

Uses

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

Do not use on damaged or broken skin.

When using this product

When using this product, keep out of eyes. Rinse with water to remove.

Stop use

Stop use and ask a doctor if rash occurs.

Keep out of

Keep out of reach of children . If product is swallowed . get medical help or contact a Poison Control Center right away.

Directions

Apply liberally 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 months : Ask a doctor.

Other information

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

Inactive ingredients

Water , Squalane , Shea Butter Ethyl Esters , Isoamyl Laurate , Polyglyceryl-3 Ricinoleate , Butyl Octyl Salicylate Propanediol , Isopropyl Isostearate , Polyglyceryl-3 Diisostearate , Methylpropanediol , Niacinamide , Sodium Chloride , Lecithin Caprylyl Glycol Panthenol (Vitamin B5) , Vitamin C , Silica Aloe Barbadensis Leaf Juice Powder Bisabolol , Hydrolyzed Sodium Hyaluronate , Allantoin , Phenylpropanol Sodium Myristoyl Glutamate , Sodium Hyaluronate , Tetrasodium Glutamate Diacetate , Aluminum Hydroxide Polyhydroxystearic Acid

尺寸: 39*39*123mm+内托





EVERYRAY SUNCARE SUNSHINE SERUM

everyray suncare sunshine serum liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83447-002
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	12 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SILICON DIOXIDE (UNII: ETJ7Z6XB4)	
WATER (UNII: 059QF0KO0R)	
SQUALANE (UNII: GW89575KF9)	
SHEA BUTTER ETHYL ESTERS (UNII: V2CI786FPG)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)
SODIUM MYRISTOYL GLUTAMATE (UNII: AYU7QD893W)
HYALURONIC ACID (UNII: S270N0TRQY)
POLYHYDROXYSTEARIC ACID STEARATE (UNII: 8KQ7I65XZE)
POLYGLYCERYL-3 RICINOLEATE (UNII: MZQ63P0N0W)
PROPANEDIOL (UNII: 5965N8W85T)
NIACINAMIDE (UNII: 25X51I8RD4)
PHENYLPROPANOL (UNII: 0F897O3O4M)
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)
CAPRYLYL GLYCOL (UNII: 00YIU5438U)
LEVOMENOL (UNII: 24WE03BX2T)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
PANTHENOL (UNII: WW9CM0O67Z)
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)
ALLANTOIN (UNII: 344S277G0Z)
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)
METHYLPROPANEDIOL (UNII: N8F53B3R4R)
ASCORBIC ACID (UNII: PQ6CK8PD0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83447-002-01	50 mL in 1 BOX; Type 0: Not a Combination Product	09/25/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	09/25/2023	

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

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Fantasy Biotechnology Co.,Ltd		619047084	manufacture(83447-002)

Revised: 9/2023

Guangzhou Fantasy Biotechnology Co.,Ltd