OMNILUX MINERAL FACIAL SUNSCREEN- zinc oxide lotion
The Lotus Global Group, Inc, dba GlobalMed Technologies

Omnilux Mineral Facial Sunscreen

Omnilux Mineral Facial Sunscreen

Directions

- Apply liberally 15 minutes before sun exposure
- Reapply at least every 2 hours
- Use a water resistant sunscreen if swimming or sweating
- Children under 6 months of age: Ask a doctor
- **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease the risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including: ● Limit your time in the sun, especially from 10 a.m. - 2 p.m. ● Wear long-sleeved shirts, pants, hats, and sunglasses

Omnilux Mineral Facial Sunscreen

Inactive ingredients

Water, Caprylic/Capric Triglyceride, Isoamyl Laurate, Squalane, Butyloctyl Salicylate, Jojoba Esters, Glycerin, Methylpropanediol, Glyceryl Stearate, Cetyl Alcohol, Silica, Polyglyceryl-4 Diisostearate/Polyhydroxystearate/ Sebacate, PEG-75 Stearate, Caprylyl Glycol, Pyrus Malus (Apple) Fruit Extract, Allantoin, Lecithin, Lysolecithin, Sclerotium Gum, Rutin, Xanthan Gum, Pullulan, Phenylpropanol, Ceteth-20, Steareth-20, Aloe Barbadensis Leaf Juice Powder, Bisabolol, Polyglyceryl-2 Dipolyhydroxystearate, Tetrasodium Glutamate Diacetate, Phenoxyethanol, C12-15 Alkyl Benzoate, Ethylhexylglycerin, Hexylene Glycol, Tocopherol, Sodium Hydroxide, Hydrogenated Lecithin, Iron Oxides.

Omnilux Mineral Facial Sunscreen

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

Omnilux Mineral Facial Sunscreen

Drug Facts

Active ingredient	Purpose
Zinc Oxide 12%	Sunscreen

Omnilux Mineral Facial Sunscreen

Warnings

For external use only

Do not use • on damaged or broken skin

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.

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Other information

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

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Questions? Call toll-free **707-309-7024** or visit www.omniluxled.com

OMNILUX MINERAL FACIAL SUNSCREEN

zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83866-6202
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	132 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALLANTOIN (UNII: 344S277G0Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
HYDROLYZED JOJOBA ESTERS (ACID FORM) (UNII: UDR641JW8W)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PHENYLPROPANOL (UNII: 0F897O3O4M)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
CETETH-20 (UNII: I835H2IHHX)	
PULLULAN (UNII: 8ZQ0AYU1TT)	
SQUALANE (UNII: GW89575KF9)	
STEARETH-20 (UNII: L0Q8IK9E08)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
PEG-75 STEARATE (UNII: OT38R0N74H)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
RUTIN (UNII: 5G06TVY3R7)	

Product Characteristics			
Color	yellow	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83866-6202-2	1 in 1 CARTON	07/26/2024	
1	NDC:83866-6202-1	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	07/26/2024	

Labeler - The Lotus Global Group, Inc, dba GlobalMed Technologies (969797406)

Registrant - Nanophase Technologies Corporation (623502044)

Revised: 8/2024 The Lotus Global Group, Inc, dba GlobalMed Technologies