VOLITION PRISMATIC LUMINIZING SHIELD- zinc oxide lotion Volition

Volition Prismatic Luminizing Shield

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Directions Shake well. Apply liberally 15 minutes before sun exposure. Reapply: after 40 minutes of swimming or sweating; immediately after towel drying; at least every 2 hours. 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 this risk, regularly use a sunscreen with broad spectrum SPF value of 15 or higher and other sun protection measures including: limit your time in the sun, especially from 10a.m. - 2 p.m. Wear long-sleeve shirts, pants, hats and sunglasses.

Inactive Ingredients Water, C12-15 Alkyl Benzoate, Jojoba Esters, Caprylic/Capric Triglyceride, Polyglyceryl-4 Diisostearate/Polyhydroxystearate/Sebacate, Propanediol, Tridecyl Salicylate, Niacinamide, Methylpropanediol, Silica, Coco-Caprylate, Sodium Hyaluronate, Allantoin, Helianthus Annuus (Sunflower) Extract, Oryza Sativa (Rice) Bran Extract, Rosmarinus Officinalis (Rosemary) Leaf Extract, Ricinus Communis (Castor) Seed Oil, Caprylyl Glycol, Synthetic Fluorphlogopite, Titanium Dioxide, Tocopherol, Hydrogenated Castor Oil, Bisabolol, Lecithin, Hydrogenated Lecithin, Polyglyceryl-2 Dipolyhydroxystearate, Polyglyceryl-3 Diisostearate, Stearalkonium Hectorite, Polyglyceryl-3 Polyricinoleate, Phenylpropanol, Tetrasodium Glutamate Diacetate, Nylon-12, Sodium Hydroxide, Sodium Chloride, Tin Oxide, Iron Oxides.

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

Volition Prismatic Luminizing Shield

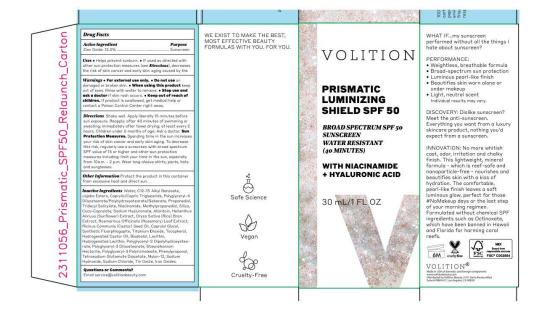
Drug Facts	
Active Ingredient	Purpose
Zinc Oxide: 12.0%	Sunscreen

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Active Ingredient	Purpose	
Zinc Oxide: 12.0%	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 a doctor if skin rash occurs. • Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.



Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72577-624

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
ROSMARINUS OFFICINALIS FLOWERING TOP OIL (UNII: OXN0D3N28L)	
RICINUS COMMUNIS SEED (UNII: 7EK4SFN1TX)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
STEARALKONIUM HECTORITE (UNII: OLX698AH5P)	
JOJOBA OIL (UNII: 724GKU717M)	
ORYZA SATIVA WHOLE (UNII: 84IVV0906Z)	
TRIDECYL SALICYLATE (UNII: AZ Q08K38Z1)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TOCOPHEROL (UNII: R0ZB2556P8)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
NYLON-12 (UNII: 446U8J075B)	
PHENYLPROPANOL (UNII: 0F897O3O4M)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
ALLANTOIN (UNII: 344S277G0Z)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
.ALPHABISABOLOL, (+)- (UNII: 105S6I733Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PROPANEDIOL (UNII: 5965N8W85T)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)	

Product Characteristics

Color	brown	Score
Shape		Size
Flavor		Imprint Code
Contains		

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:72577-624- 02	1 in 1 CARTON	04/22/2024			
1	NDC:72577-624- 01	30 mL in 1 BOTTLE; 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	04/22/2024		

Labeler - Volition (034302621)

Registrant - Nanophase Technologies Corporation (623502044)

Establishment			
Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		050383046	api manufacture(72577-624)

Establishment			
Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		118812921	pack(72577-624), manufacture(72577-624)

Establishment			
Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		623502044	api manufacture(72577-624) , manufacture(72577-624)

Revised: 9/2024 Volition