

DR LIFT SPF 50 CREAM- titanium dioxide, zinc oxide cream
Spa de Soleil

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.

Moisturizer Sunscreen SPF 50

Active Ingredients

Zinc Oxide 9.86%

Octinoxate 7.5%

Purpose

Sunscreen

Warnings

Warnings When using this product

- keep out of the eye area
- rinse with water to remove
- stop use and ask a doctor if rash or irritation develops
- for external use only
- keep out of reach of children

KEEP OUT OF REACH OF CHILDREN

Directions

Directions: Apply generously and evenly thirty minutes before sun exposure. Reapply at least every two hours. Use a water-resistant sunscreen if swimming or sweating.

Children under six months of age: Consult a doctor.

Apply generously and evenly thirty minutes before sun exposure. Reapply at least every two hours.

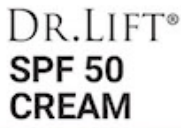


Inactive Ingredients

Inactive Ingredients: Aqua, Glycerin, Cocos Nucifera (Coconut) Oil, Carthamus Tinctorius (Safflower) Seed Oil, Heptyl Undecylenate, Caprylic/Capric Triglyceride, Cetyl Alcohol, Polyglyceryl-6 Distearate, Glyceryl Stearate, PEG-100 Stearate, Titanium Dioxide, Jojoba Esters, Polyhydroxystearic Acid, Polyglyceryl-3 Beeswax, Carbomer, Disodium EDTA, *CO Glycerin, Sodium Hydroxide, Thymus Vulgaris (Thyme) Leaf Extract, Rosa Canina Fruit Extract, Gardenia Florida Fruit Extract, Fucus Vesiculosus Extract, *CO Carum

Petroselinum (Parsley) Extract, *CO Centella Asiatica Leaf Extract, *CO Cucumis Sativus (Cucumber) Seed Extract, *CO Daucus Carota Sativa (Carrot) Root Extract, *CO Euphrasia Officinalis Extract, Ascorbic Acid (Vitamin C), Retinyl Palmitate (Vitamin A), Tocopheryl Acetate (Vitamin E), Phenoxyethanol, Ethylhexylglycerin.

*Certified Organic

Principal Display Panel

 <p>1.75 fl oz (50 mL)</p>		Drug Facts Active Ingredients: Zinc Oxide 9.86% Octinoxate 7.5%	Purpose: Sunscreen Sunscreen	Drug Facts (Continued) Warnings: - For external use only. - Do not use on damaged or broken skin. - When using this product keep out of eyes. - If swallowed, seek medical help or contact a poison control center right away. - Keep out of reach of children. - Jar is glass and may shatter if dropped.	SA11-PL_AMZ Manufactured by Sun Art Co., Inc. Sun Valley, CA MADE IN USA	
		Uses: - Helps prevent sunburn. - Higher SPF gives more sunburn protection. - If used as directed with other sun protection measures (see Sun Alert), decreases the risk of skin cancer and early skin aging caused by the sun.	Directions: Apply generously and evenly thirty minutes before sun exposure. Reapply at least every two hours. Use a water resistant sunscreen if swimming or sweating. Children under six months of age: Consult a doctor. Other Information: Protect the product in this container from excessive heat.	FOR INACTIVE INGREDIENTS PEEL HERE		
		Drug Facts (Continued) SA11-PL_AMZ Inactive Ingredients: Aqua, Glycerin, Cocos Nucifera (Coconut) Oil, Carthamus Tinctorius (Safflower) Seed Oil, Heptyl Undecylenate, Caprylic/Capric Triglyceride, Cetyl Alcohol, Polyglyceryl 6 Distearate, Glyceryl Stearate, PEG-100 Stearate, Titanium Dioxide, Jojoba Esters, Polyhydroxystyrene Acid, Polyglyceryl 3 Beeswax, Carbomer, Disodium EDTA, *CO Glycerin, Sodium Hydroxide, Thymus Vulgaris (Thyme) Leaf Extract, Rosa Canina Fruit Extract, Gardenia Florida Fruit Extract, Fucus Vesiculosus Extract, *CO Carum Petroselinum (Parsley) Extract, *CO Centella Asiatica Leaf Extract, *CO Cucumis Sativus (Cucumber) Seed Extract, *CO Daucus Carota Sativa (Carrot) Root Extract, *CO Euphrasia Officinalis Extract, Ascorbic Acid (Vitamin C), Retinyl Palmitate (Vitamin A), Tocopheryl Acetate (Vitamin E), Phenoxyethanol, Ethylhexylglycerin. *Certified Organic				

<h2>DR LIFT SPF 50 CREAM</h2> <p>titanium dioxide, zinc oxide cream</p>			
<h3>Product Information</h3>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68062-9007
Route of Administration	TOPICAL		
<h3>Active Ingredient/Active Moiety</h3>			
	Ingredient Name	Basis of Strength	Strength
	ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	4.93 mg in 50 mL
	OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	3.75 mg in 50 mL
<h3>Inactive Ingredients</h3>			
	Ingredient Name		Strength
	WATER (UNII: 059QF0KO0R)		
	HEPTYL UNDECYLENATE (UNII: W77QUB6GXO)		
	SAFFLOWER OIL (UNII: 65UEH262IS)		
	GLYCERIN (UNII: PDC6A3C0OX)		
	COCONUT OIL (UNII: Q9L0O73W7L)		
	CAPRYLIC/CAPRIC/LAURIC TRIGLYCERIDE (UNII: FJ1H6M2JG9)		
<h3>Packaging</h3>			
		Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68062-9007-1	50 mL in 1 TUBE; Type 0: Not a Combination Product	08/24/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M020	08/24/2023	

Labeler - Spa de Soleil (874682867)

Establishment

Name	Address	ID/FEI	Business Operations
Spa de Soleil		874682867	manufacture(68062-9007)

Revised: 8/2023

Spa de Soleil