DESERT SPIRIT SUNSCREEN- zinc oxide, octinoxate, octisalate, and oxybenzone lotion
R & R Lotion, Inc

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.

Desert Spirit Sunscreen

Drug Facts

Active Ingredients	Purpose
Octinoxate 5.9%	Sunscreen
Octisalate 3.8%	Sunscreen
Oxybenzone 4.7%	Sunscreen
Zinc Oxide 6.4%	Sunscreen

Uses

- Helps prevent sunburn
- If used as directed with other sun protection measures (see directions), decreases the risk of skin cancer & early skin aging caused by the sun

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 a rash occurs

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

Directions

- apply liberally 15 minutes before sun exposure reapply:
  - after 80 minutes of swimming or sweating
  - immediatly after towel drying
  - 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 of 15 or higher and other sun protection measures including:

  - limit time in the sun, especially from 10am-2pm
  - wear long-sleeve shirts, pants, hats, and sunglasses

- children under 6 months: Ask a doctor

Inactive Ingredients

Purified DI Water, Octyl Stearate/Octyl Palmitate/Dioctyl Adipate, Propylene Glycol, Glyceryl Stearate & Peg 100, Ceteareth Alcohol & Ceteareth 20, Xanthan Gum, Imidazolidinyl Urea, Methyl Paraben,

Propyl Paraben, Crodafos N3N, and Vitamin E.

Other Information

- protect this product from excessive heat and direct sun

PRINCIPAL DISPLAY PANEL - 50 ML Bottle Label

DESERT SPIRIT

Broad Spectrum
Sunscreen

SPF 30

80 Minute Water
Resistance

1.7OZ. (50ML) e

Pump to Prime

A Natural Zinc Oxide Sunscreen formulated to protect individuals who are exposed to the Sun's Ultraviolet Rays.

- Dermatologist Recommended
- Physical Zinc Technology
- Fragrance Free
- Anti-Aging
- UVA
- Natural Healing Properties
- Hypoallergenic

SUN ALERT
UV Rays of the Sun are made of UVA & UVB. It is important to protect against both UVA and UVB Rays. Zinc Oxide provides the highest UVA-UVB protection by creating a Physical Barrier and reflects both UVA & UVB Rays.

FDA Registered

Made by Americans for American Workers

Desert Spirit
15547 N. 77th Street
Scottsdale, AZ 85260
480-443-9255
Desert-Spirit.com

No Animal Testing
36M
DS-50ML-SS
6 54441 34770 7

DESERT SPIRIT

Broad Spectrum Sunscreen

SPF 30

80 Minute Water Resistance

1.7OZ. (50ML) e

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DESERT SPIRIT SUNSCREEN

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59555-103
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	64 mg in 1 mL
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	59 mg in 1 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	38 mg in 1 mL
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	47 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Xanthan Gum (UNII: TTV12P4NEE)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Methylparaben (UNII: A2I8C7HI9T)	
Diethylhexyl Adipate (UNII: MBY1SL921L)	
Ethylhexyl Palmitate (UNII: 2865993309)	
Ethylhexyl Stearate (UNII: EG3PA2K3K5)	
Propylparaben (UNII: Z8IX2SC1OH)	
.Alpha.-Tocopherol (UNII: H4N855PNZ1)	
Imidurea (UNII: M629807ATL)	
Cetostearyl Alcohol (UNII: 2DMT128M1S)	
Polyoxyl 20 Cetostearyl Ether (UNII: YRC528SWUY)	
Diethanolamine Oleth-3 Phosphate (UNII: Y67NX5905E)	
Glyceryl Monostearate (UNII: 230OU9XXE4)	
Peg-100 Stearate (UNII: YD01N1999R)	

Product Characteristics

Color	WHITE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59555-103-03	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/21/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	12/21/2015	

Labeler - R & R Lotion, Inc (062979000)