

REJUVASKIN MINERAL FACIAL SUNSCREEN SPF 32 SHEER- rejuvaskin mineral facial sunscreen spf 32 sheer lotion Atlantic Medical Products , LLC

Rejuvaskin Mineral Facial Sunscreen Broad Spectrum SPF32 Sheer lotion for daily use 4% Niacinamide (IS0122-7)

Active ingredients

Zinc Oxide 20%.....Sunscreen

When using this Product keep out of eyes. Rise with water to remove.

For external use only.

Stop use and ask a physician if rash occurs.

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

Do not use section on damaged or broken skin

Inactive Ingredients

Benzyl Alcohol

Butyloctyl Salicylate

Caprylic/Capric Triglyceride

Citric Acid

Coco-Caprylate

Dehydroacetic Acid

Glycerin

Glyceryl Stearate

Niacinamide

Panthenol

Polyhydroxystearic Acid

Sodium Phytate

Saccharide Isomerate

Sodium Stearoyl Glutamate

Sucrose palmitate-stearate

Sucrose Laurate

Tocopheryl Acetate

Water

Xanthan Gum

Active Ingredients

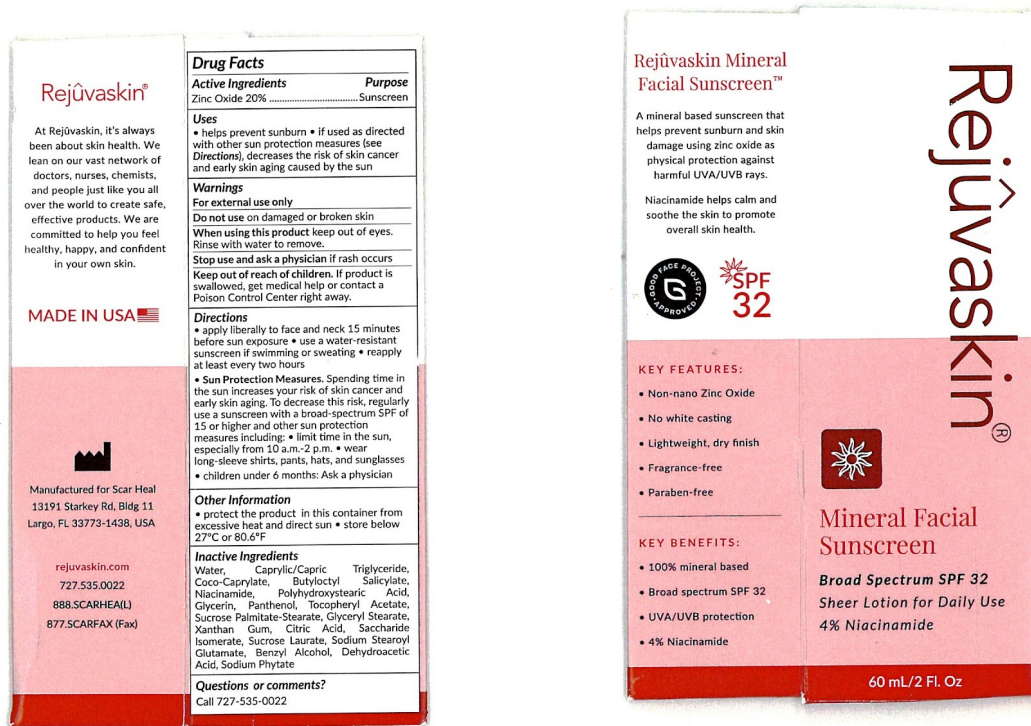
Zinc Oxide 20%.....Sunscreen

Uses helps prevent sunburn

if used directed with other sun protection measures(see Directions),decreases the risk of skin cancer and early agung caused by the sun.

Directions

- Apply liberally to face and neck 15 minutes before sun exposure
- use a water-resistant sunscreen if swimming or sweating
- reapply at least every two hours
- Sun Protection measures Spending time in the sun increase 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 10 am- 2 pm
- wear long-sleeve shirts, pants, hats, and sunglasses
- children under 6 months: Ask a physician



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84375-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	20 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
NIACINAMIDE (UNII: 25X518RD4)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
SUCROSE STEARATE (UNII: 274KW0050M)	
XANTHAN GUM (UNII: TTV12P4NEE)	
WATER (UNII: 059QF0KO0R)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PANTHENOL (UNII: WW9CM0067Z)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
PHYTATE SODIUM (UNII: 88496G1ERL)	
SACCHARIDE ISOMERATE (UNII: W8K377W98I)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
SUCROSE STEARATE/PALMITATE ESTER (75% MONO ESTER) (UNII: L98X941W2B)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SUCROSE LAURATE (UNII: 05Q7CD0E49)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84375-101-01	60 g in 1 TUBE; Type 0: Not a Combination Product	06/04/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/04/2024	

Labeler - Atlantic Medical Products , LLC (014302733)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions		081030372	manufacture(84375-101)

Revised: 6/2024

Atlantic Medical Products , LLC