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

ACTIVE INGREDIENT

Zinc Oxide 20%.....Sunscreen

WHEN USING

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

WARNINGS

For external use only.

STOP USE

Stop use and ask a physician if rash occurs.

KEEP OUT OF REACH OF CHILDREN

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

DO NOT USE

Do not use on damaged or broken skin.

INACTIVE INGREDIENTS

Inactive Ingredients

Benzyl Alcohol

Butyloctyl Salicylate

Caprylic/Capric Triglyceride

Citric Acid

Coco-Caprylate

Dehydroacetic Acid

Glycerin

Glyceryl Stearate

Niacinamide

Pathenol

Polyhydroxystearic Acid

Sodium Phytate

Saccharide Isomerate

Sodium Stearoyl Glutamate

Sucrose palmitate-stearate

Sucrose Laurate

Tocopheryl Acetate

Water

Xanthan Gum

PURPOSE

Active Ingredients

Zinc Oxide 20%.....Sunscreen

INDICATIONS & USAGE

Uses helps prevent sunburn.

If used as directed with other sun protection measures (see Directions) decreases the risk of skin cancer and early aging caused by the sun.

DOSAGE & ADMINISTRATION

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 increases your risk of skin cancer and early

skin aging. To decrease this risk, regularly use a sunscreen with 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 physician

PRINCIPAL DISPLAY PANEL

REJUVASKIN MINERAL FACIAL SUNSCREEN SPF 32 SHEER

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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)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PHYTATE SODIUM (UNII: 88496G1ERL)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
SUCROSE STEARATE/PALMITATE ESTER (75% MONO ESTER) (UNII: L98X941W2B)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
PANTHENOL (UNII: WV9CM0O67Z)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
XANTHAN GUM (UNII: TTV12P4NEE)	
NIACINAMIDE (UNII: 25X51I8RD4)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
SUCROSE STEARATE (UNII: 274KW0O50M)	
WATER (UNII: 059QF0KO0R)	

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SACCHARIDE ISOMERATE (UNII: W8K377W98I)	
SUCROSE LAURATE (UNII: 05Q7CD0E49)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84375-101-02	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: 1/2026 Atlantic Medical Products, LLC