

SUNRIGHT SOLAR SCREEN MINERAL FACE SUNSCREEN- zinc oxide lotion
NSE Products, Inc.

Sunright® Solar Screen Mineral Face Sunscreen

Drug Facts

Active Ingredient

Zinc Oxide 18%

Purpose

Sunscreen

Uses

- Helps prevent sunburn. Higher SPF gives more sunburn protection.
- Retains SPF after 80 minutes of activity in the water, sweating, or perspiring.

Warning

- **For external use only.**

Do not use on broken or damaged skin.

Stop use and ask a doctor if rash occurs.

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

- **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 or water exposure.
- Reapply: After 80 minutes of swimming or sweating
 - Immediately after towel drying
 - At least every 2 hours
- Children under 6 months: Ask a doctor

Other Information

- Sun alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun

Inactive Ingredients

Water (Aqua), Dimethicone, Butylene Glycol, PEG-10 Dimethicone, C12-15 Alkyl

Benzoate, Bisabo-lol, Dimethicone/Vinyl Dimethicone Crosspolymer, Tocopheryl Acetate, Tocopherol, Stearalkonium Hectorite, Cetyl PEG/PPG-10/1 Dimethicone, Hexyl Laurate, Polyglyceryl-4 Isostearate, Propylene Carbonate, Sodium Citrate, Sodium Chloride, Hydroxyacetophe-none, Chlorphenesin, Sodium Dehydroacetate.

Questions?

1-801-345-9900

PRINCIPAL DISPLAY PANEL - 50 mL Tube Carton

NU SKIN®

SUNRIGHT®

SOLAR SCREEN

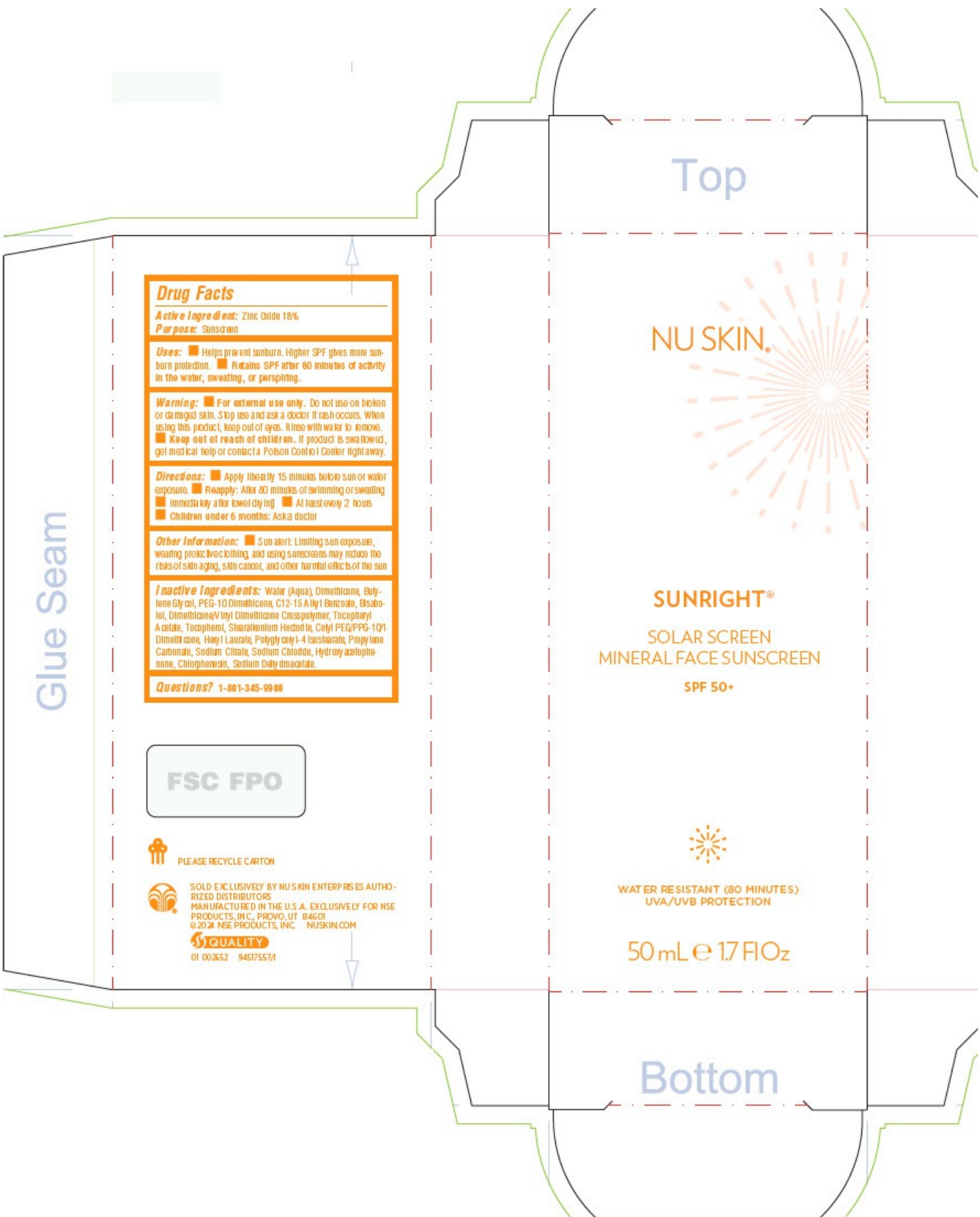
MINERAL FACE SUNSCREEN

SPF 50+

WATER RESISTANT (80 MINUTES)

UVA/UVB PROTECTION

50 mL e 1.7 Fl Oz



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Questions? 1-801-345-9900

FSC FPO

 PLEASE RECYCLE CARTON

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 **QUALITY**
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Top

NU SKIN.

SUNRIGHT®

SOLAR SCREEN
 MINERAL FACE SUNSCREEN

SPF 50+

WATER RESISTANT (80 MINUTES)
 UVA/UVB PROTECTION

50 mL e 1.7 FlOz

Bottom

SUNRIGHT SOLAR SCREEN MINERAL FACE SUNSCREEN
 zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62839-2652
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	180 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Dimethicone, Unspecified (UNII: 92RU3N3Y1O)	
Butylene Glycol (UNII: 3XUS85K0RA)	
Alkyl (C12-15) Benzoate (UNII: A9EJ3J61HQ)	
Levomenol (UNII: 24WE03BX2T)	
.Alpha.-Tocopherol Acetate (UNII: 9E8X80D2L0)	
Stearalkonium Hectorite (UNII: O LX698AH5P)	
Hexyl Laurate (UNII: 4CG9F9W01Q)	
Sodium Chloride (UNII: 451W47IQ8X)	
Polyglyceryl-4 Isostearate (UNII: 820DPX33S7)	
Hydroxyacetophenone (UNII: G1L3HT4CMH)	
Chlorphenesin (UNII: I670DAL4SZ)	
Sodium Citrate, Unspecified Form (UNII: 1Q73Q2JULR)	
Propylene Carbonate (UNII: 8D08K3S51E)	
Tocopherol (UNII: R0ZB2556P8)	
Sodium Dehydroacetate (UNII: 8W46YN971G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-2652-1	1 in 1 CARTON	05/01/2024	
1		50 mL in 1 TUBE; 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	05/01/2024	

Labeler - NSE Products, Inc. (803486393)

