

**NU SKIN SUNRIGHT 35 BROAD SPECTRUM SPF 35- titanium dioxide and zinc oxide lotion**  
**NSE Products, 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.*

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**Nu Skin® Sunright® 35 Broad Spectrum SPF 35**

**Drug Facts**

**Active Ingredient**

Titanium Dioxide 4.2%, Zinc Oxide 4.5%.

**Purpose**

Sunscreen

**Use**

- Helps prevent sunburn. Higher SPF gives more sunburn protection.
- If used as directed with other sun protection measures (see **Directions**), decreases the risk of skin cancer and early skin aging caused by the sun.

**Warnings**

- **For external use only.**
- **Do not use** on damaged or broken 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 exposure.
- Reapply:
  - After 40 minutes of swimming or sweating
  - Immediately after towel drying
  - At least every 2 hours
- Children under 6 months: Ask a doctor
- **Sunprotection 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 10 a.m.–2p.m.
- Wear long-sleeved shirts, pants, hats, and sunglasses.

### **Inactive Ingredients**

Acrylates/Dimethicone Copolymer, C12-15 Alkyl Benzoate, Allantoin, Aloe Barbadensis Leaf Juice, Bisabolol, Caprylic/Capric Triglyceride, Caprylyl Methicone, Carthamus Tinctorius (Safflower) Seed Oil, Chlorphenesin, Cyclopentasiloxane, Di-C12-13 Alkyl Tartrate, Dicaprylyl Carbonate, Diglycerin, Dimethicone/PEG-10/15 Crosspolymer, Dimethicone/Vinyl Dimethicone Crosspolymer, Fragrance (Parfum), Haematococcus Pluvialis Extract, Lauryl PEG-9 Polydimethylsiloxyethyl Dimethicone, Panthenol, Phenoxyethanol, Physalis Angulata Extract, Polyglyceryl-3 Polydimethylsiloxyethyl Dimethicone, Propylene Carbonate, Propylheptyl Caprylate, Silica, Sodium Chloride, Sodium Citrate, Stearalkonium Hectorite, Tocopherol, Triethoxysilylethyl Polydimethylsiloxyethyl Hexyl Dimethicone, Vinyl Dimethicone/Methicone Silsesquioxane Crosspolymer, Water (Aqua).

### **Other Information**

- Protect this product from excessive heat and direct sun.

### **Questions?**

**1-888-742-7626**

### **PRINCIPAL DISPLAY PANEL - 100 mL Bottle Carton**

NU SKIN®

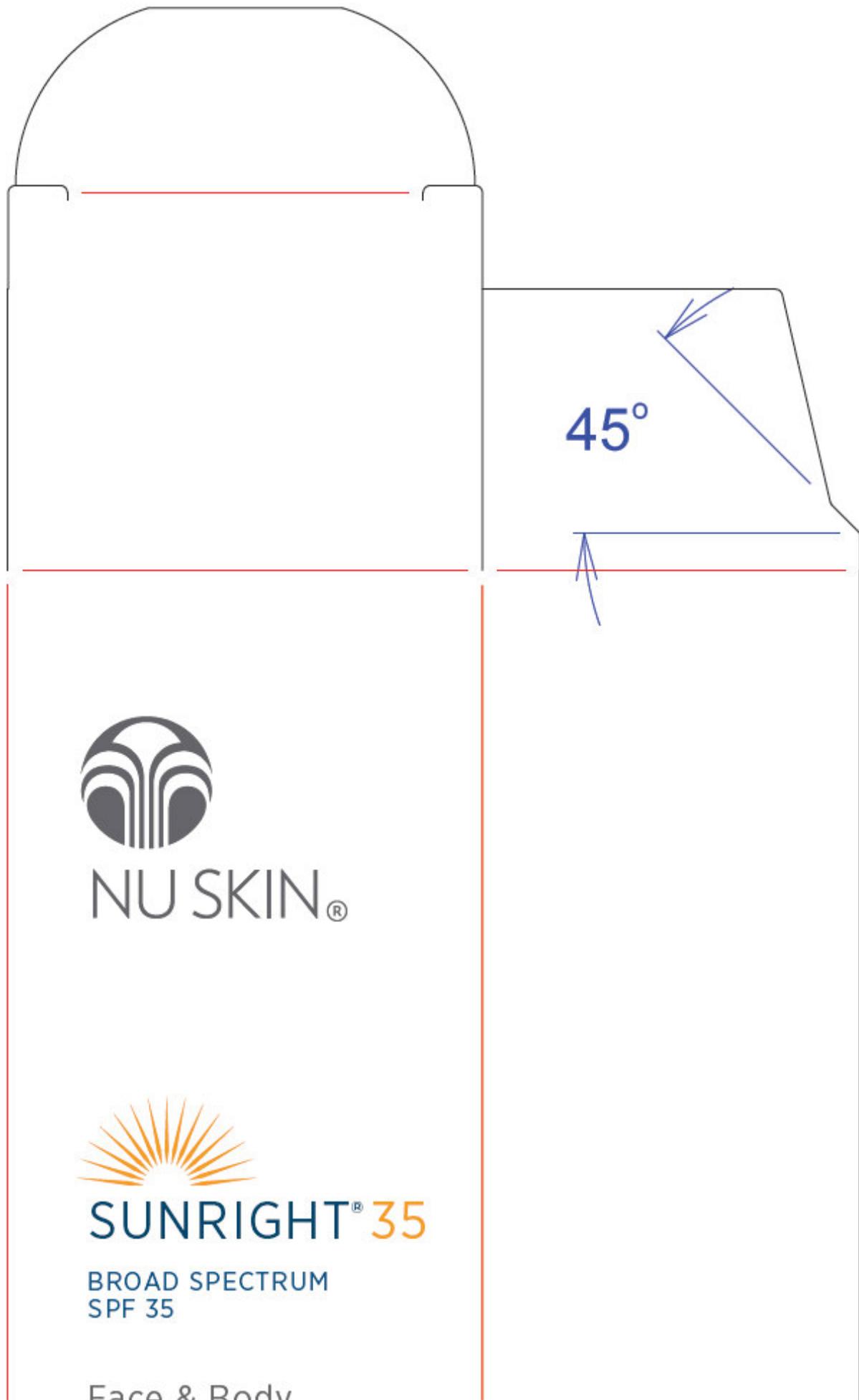
SUNRIGHT® 35

BROAD SPECTRUM  
SPF 35

Face & Body  
Sunscreen

Water Resistant  
(80 minutes)

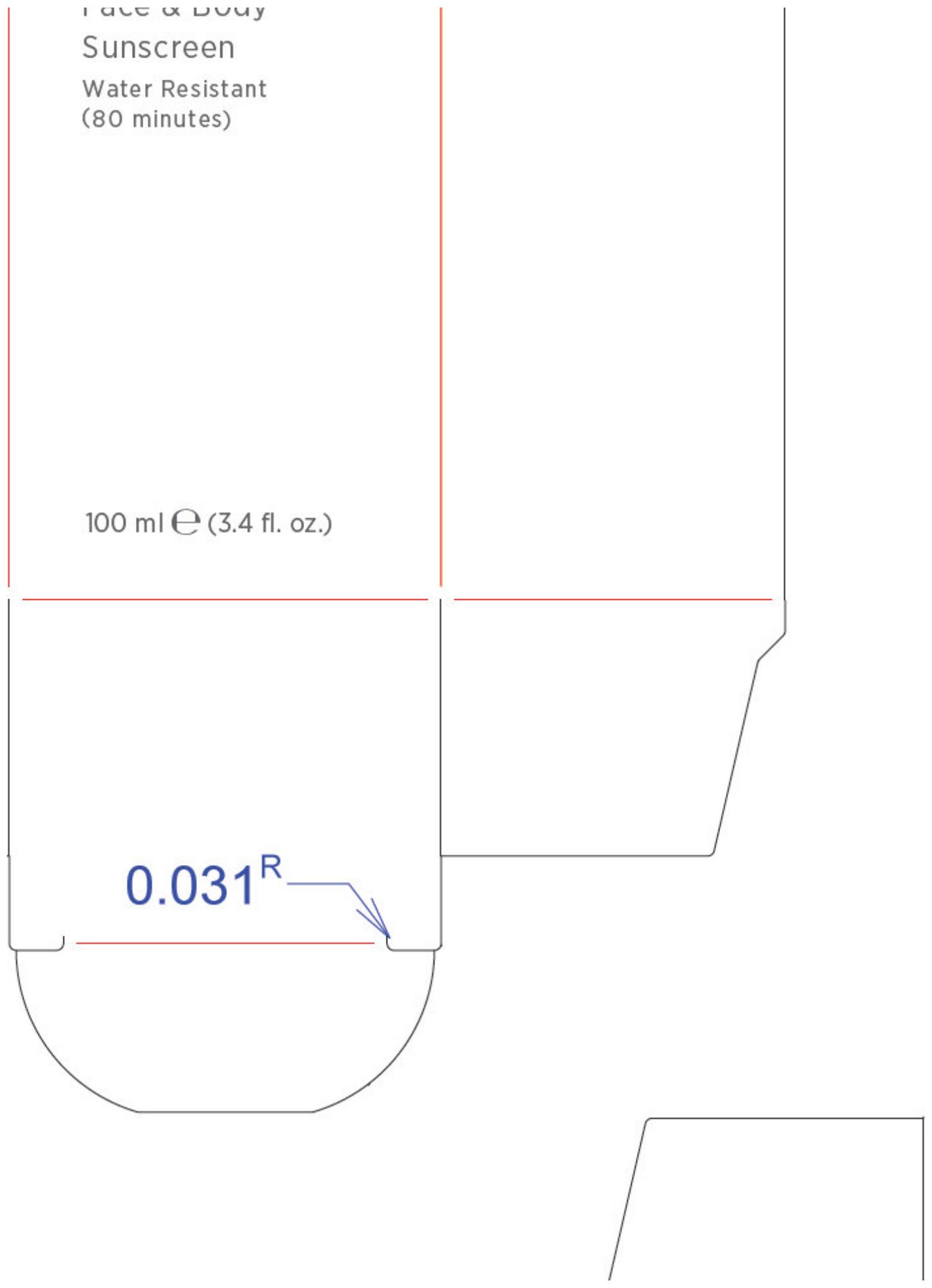
100 ml e (3.4 fl. oz.)



Face & Body  
Sunscreen  
Water Resistant  
(80 minutes)

100 ml e (3.4 fl. oz.)

0.031<sup>R</sup>



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### Other Information

SAFETY-ALLERGY-DERMATOLOGIST TESTED  
Sunright® Face & Body Sunscreen Water  
Resistant with a sun protection factor (SPF) of  
35 shields your skin from UVA/UVB rays.  
Formulated with Avobenzone to provide protec-  
tion from UVA rays, which may contribute to skin

Protect this product from excessive heat and direct sun.

**Questions?**  
1-888-742-7626

damage and premature aging.

SOLD EXCLUSIVELY BY NU SKIN ENTERPRISES  
AUTHORIZED DISTRIBUTORS  
MANUFACTURED IN THE U.S.A. EXCLUSIVELY  
FOR NSE PRODUCTS, INC., PROVO, UT 84601  
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## NU SKIN SUNRIGHT 35 BROAD SPECTRUM SPF 35

titanium dioxide and zinc oxide lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:62839-1344
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Titanium Dioxide</b> (UNII: 15FIX9V2JP) (Titanium Dioxide - UNII:15FIX9V2JP)	Titanium Dioxide	42 mg in 1 mL
<b>Zinc Oxide</b> (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	45 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>Alkyl (C12-15) Benzoate</b> (UNII: A9EJ3J61HQ)	
<b>Allantoin</b> (UNII: 344S277G0Z)	
<b>Aloe Vera Leaf</b> (UNII: ZY81Z83H0X)	
<b>Levomenol</b> (UNII: 24WE03BX2T)	
<b>Medium-Chain Triglycerides</b> (UNII: C9H2L21V7U)	
<b>Caprylyl Trisiloxane</b> (UNII: Q95M2P1KJL)	
<b>Safflower Oil</b> (UNII: 65UEH262IS)	
<b>Chlorphenesin</b> (UNII: I670DAL4SZ)	
<b>Cyclomethicone 5</b> (UNII: 0THT5PCI0R)	
<b>Di-C12-13 Alkyl Tartrate</b> (UNII: 603USX4K0G)	
<b>Dicaprylyl Carbonate</b> (UNII: 609A3V1SUA)	
<b>Diglycerin</b> (UNII: 3YC120743U)	

<b>Dimethicone</b> (UNII: 92RU3N3Y1O)	
<b>Haematococcus Pluvialis</b> (UNII: 31T0FF0472)	
<b>Lauryl PEG-9 Polydimethylsiloxylethyl Dimethicone</b> (UNII: 25G622K2RA)	
<b>Panthenol</b> (UNII: WW9CM0O67Z)	
<b>Phenoxyethanol</b> (UNII: HIE492ZZ3T)	
<b>Physalis Angulata whole</b> (UNII: W4TKW9D5GG)	
<b>Propylene Carbonate</b> (UNII: 8D08K3S51E)	
<b>Propylheptyl Caprylate</b> (UNII: 991Z19V2OD)	
<b>Silicon Dioxide</b> (UNII: ETJ7Z6XBU4)	
<b>Sodium Chloride</b> (UNII: 451W47IQ8X)	
<b>Sodium Citrate, Unspecified Form</b> (UNII: 1Q73Q2JULR)	
<b>Stearalkonium Hectorite</b> (UNII: O LX698AH5P)	
<b>Tocopherol</b> (UNII: R0ZB2556P8)	
<b>Water</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-1344-1	1 in 1 CARTON	04/23/2014	
1		100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part352	04/23/2014	

**Labeler** - NSE Products, Inc. (803486393)

Revised: 1/2022

NSE Products, Inc.