# NEUTROGENA SHEER ZINC FACE MINERAL SUNSCREEN BROAD SPECTRUM SPF 50- zinc oxide lotion Johnson & Johnson Consumer 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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# Neutrogena <sup>®</sup> Sheer Zinc Face mineral sunscreen BROAD SPECTRUM SPF 50 Drug Facts

## **Active ingredient**

Zinc Oxide (21.6%)

## **Purpose**

Sunscreen

#### Uses

- helps prevent sunburn
- 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
- When using this product keep out of eyes. Rinse with water to remove
- Stop use and ask a doctor if rash occurs
- **Keep out of reach of children.** If 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
  - immediately after towel drying
  - at least 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 value of 15 or higher and other sun protection measures

## including:

- limit time in the sun, especially from 10 a.m. 2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses
- children under 6 months of age: Ask a doctor

#### Other information

- protect this product from excessive heat and direct sun
- may stain some fabrics

## **Inactive ingredients**

Water, C12-15 Alkyl Benzoate, Styrene/Acrylates Copolymer, Octyldodecyl Citrate Crosspolymer, Phenyl Trimethicone, Cetyl PEG/PPG-10/1 Dimethicone, Dimethicone, Glycerin, Polyhydroxystearic Acid, Ethyl Methicone, Silica, Cetyl Dimethicone, Triethoxycaprylylsilane, Phenoxyethanol, Glyceryl Behenate, Sodium Chloride, Acrylates/Dimethicone Copolymer, Chlorphenesin, Phenethyl Alcohol, Avena Sativa (Oat) Kernel Flour, Caprylyl Glycol, Cetyl Dimethicone/Bis-Vinyldimethicone Crosspolymer, Tocopheryl Acetate, Chrysanthemum Parthenium (Feverfew) Flower/Leaf/Stem Juice

# **Questions?**

Call toll-free **800-299-4786** or **215-273-8755** (collect). www.neutrogena.com

Distributed by:

JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 59 mL Tube Carton

Neutrogena ®

**DERMATOLOGIST RECOMMENDED BRAND** 

**Sheer Zinc** 

Face

mineral

sunscreen

**BROAD SPECTRUM SPF 50** 

**50** 

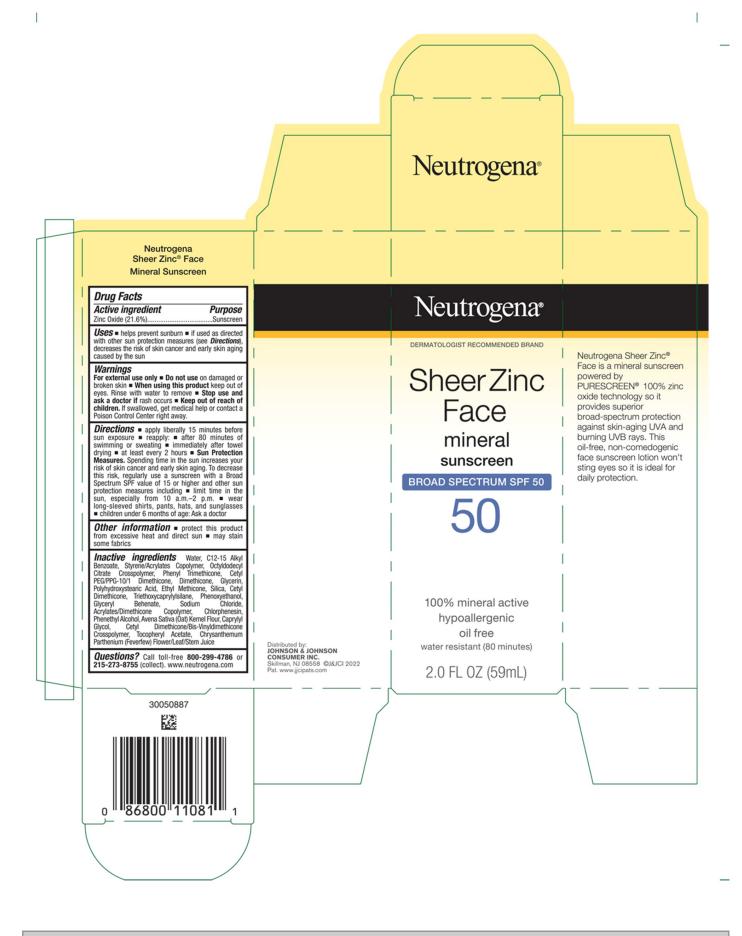
100% mineral active

hypoallergenic

oil free

water resistant (80 minutes)

2.0 FL OZ (59mL)



#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0623

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name Basis of Strength Strength

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) ZINC OXIDE 216 mg in 1 mL

#### **Inactive Ingredients**

ı	ingredient Name	Strength
I	WATER (UNII: 059QF0KO0R)	

ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)

BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER

(UNII: V5RS026Q0H)

OCTYLDODECYL CITRATE CROSSPOLYMER (UNII: X323T6QO4M)

PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)

CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)

DIMETHICONE (UNII: 92RU3N3Y10)

GLYCERIN (UNII: PDC6A3C0OX)

POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)

ETHYL METHICONE (8 MPA.S) (UNII: 3YWG8XYT8H)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)

TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)

PHENOXYETHANOL (UNII: HIE492ZZ3T)

GLYCERYL DIBEHENATE (UNII: R8WTH25YS2)

SODIUM CHLORIDE (UNII: 451W47IQ8X)

CHLORPHENESIN (UNII: 1670DAL4SZ)

PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)

OATMEAL (UNII: 8PI54V663Y)

CAPRYLYL GLYCOL (UNII: 00YIU5438U)

CETYL DIMETHICONE/BIS-VINYLDIMETHICONE CROSSPOLYMER (UNII: AE7QA6TW0Q)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

FEVERFEW (UNII: Z64FK7P217)

#### **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968- 0623-2	1 in 1 CARTON	01/31/2020	
1		59 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information				
Marketing End Date				

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023 Johnson & Johnson Consumer Inc.