NEUTROGENA SHEER ZINC 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.

Neutrogena ® Sheer Zinc 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, Caprylyl Glycol, Cetyl Dimethicone/Bis-Vinyldimethicone Crosspolymer, 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 - 88 mL Tube Label

Neutrogena ®

DERMATOLOGIST RECOMMENDED BRAND

Sheer Zinc

mineral

sunscreen

BROAD SPECTRUM SPF 50

50

100% mineral active

hypoallergenic

water resistant (80 minutes)

3.0 FL OZ (88 mL)

NEUTROGENA SHEER ZINC MINERAL SUNSCREEN BROAD SPECTRUM SPF 50

zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0613
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

Ingredient Name	Strength
PHENYLETHYL ALCOHOL (UNII: ML9LGA7468)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CETYL DIMETHICONE/BIS-VINYLDIMETHICONE CROSSPOLYMER (UNII: AE7QA6TW0Q)	
FEVERFEW (UNII: Z64FK7P217)	
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: 92RU3N3Y1O)	
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: I670DAL4SZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0613-5	147 mL in 1 TUBE; Type 0: Not a Combination Product	01/31/2020	11/01/2022
2	NDC:69968-0613-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	01/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/31/2020	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 8/2023 Johnson & Johnson Consumer Inc.