NEUTROGENA MINERAL ULTRA SHEER DRY TOUCH SPF 30- titanium dioxide, 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 Mineral Ultra Sheer Dry Touch Lotion SPF 30

Drug Facts

Active ingredients	Purpose	
Titanium Dioxide (6.6%) Zinc	Sunscreen	
Oxide (18.0%)	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, Isohexadecane, Butyloctyl Salicylate, C12-15 Alkyl Benzoate, Dimethicone, Ethylhexyl methoxycrylene, Caprylyl Methicone, Glyceryl Stearate, PEG-100 Stearate, Calcium Aluminum Borosilicate, VP/Hexadecene Copolymer, Styrene/Acrylates Copolymer, Triethoxycaprylylsilane, Aluminum Hydroxide, Xanthan Gum, Stearic Acid, Trisiloxane, Polyester-7, Phenoxyethanol, Polyhydroxystearic Acid, Caprylyl Glycol, Ethylhexylglycerin, Neopentyl Glycol Diheptanoate, Behenyl Alcohol, Tocopheryl Acetate, Disodium EDTA, Silica

Questions?

Call toll-free **800-299-4786** or **215-273-8755** (collect) or visit www.neutrogena.com Distributed by: **JOHNSON & JOHNSON CONSUMER INC.** Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 88 mL Tube Label

NEW

Neutrogena®

DERMATOLOGIST RECOMMENED BRAND

PURESCREEN+ TM

MINERAL

ULTRA

SHEER®

DRY-TOUCH LOTION

30

SUNSCREEN

BROAD SPECTRUM SPF 30

UVA/UVB PROTECTION

lightweight clean feel

100% mineral actives + vitamin e

water resistant (80 minutes)

3.0 FL OZ (88 mL)

NEUTROGENA® MINERAL ULTRA SHEER® SPF 30

Broad spectrum UVA/UVB protection

- Fragrance-free Sensitive skin Non-chalky
- · Shine-free · Lightweight clean feel
- Skin nourishing formula

Formulated with antioxidant vitamin e and a dry-touch finish.





Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

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Sunscreen

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J&JCI 2022

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Skillman.

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of reach of children. If swallowed, get medical help or contact a
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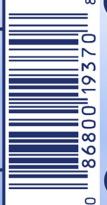
Other information = protect

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

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Free of: Chemical actives (including no oxybenzone)

Vegan. Formula not tested on animals.



PURESCREEN+TM:100% mineral actives + vitamin e





SUNSCREEN
BROAD SPECTRUM SPF 30
UVA/UVB PROTECTION

lightweight clean feel 100% mineral actives + vitamin e water resistant (80 minutes)

3.0 FL OZ (88 mL)

NEUTROGENA MINERAL ULTRA SHEER DRY TOUCH SPF 30

titanium dioxide, zinc oxide lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0776

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
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	66 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ISOHEXADECANE (UNII: 918X1OUF1E)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)	
VINYLPYRROLIDONE/HEXADECENE COPOLYMER (UNII: KFR5QEN0N9)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
DOCOSANOL (UNII: 9G10E216XY)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
WATER (UNII: 059QF0KO0R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
DIMETHICONE (UNII: 92RU3N3Y10)	
TRISILOXANE (UNII: 9G1ZW13R0G)	
CALCIUM ALUMINUM BOROSILICATE (UNII: 3JRB8A35M0)	
PEG-100 STEARATE (UNII: YD01N1999R)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POLYESTER-7 (UNII: 0841698D2F)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968- 0776-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	12/06/2022	
2	NDC:69968- 0776-1	12 in 1 TRAY	12/06/2022	
2		14 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

OTC monograph not final	part352	12/06/2022	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023 Johnson & Johnson Consumer Inc.