NEUTROGENA MINERAL UV TINT FACE MEDIUM DEEP SPF 30- titanium dioxide, zinc oxide lotion

NEUTROGENA MINERAL UV TINT FACE DEEP SPF 30- titanium dioxide, zinc oxide lotion

NEUTROGENA MINERAL UV TINT FACE LIGHT SPF 30- titanium dioxide, zinc oxide lotion

NEUTROGENA MINERAL UV TINT FACE MEDIUM SPF 30- titanium dioxide, zinc oxide lotion

Johnson & Johnson Consumer Inc.

Neutrogena MINERAL UV TINT FACE LIQUID SPF 30

Drug Facts

Active ingredients

Titanium Dioxide 3.2%, 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

- Shake well before use
- apply generously 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, Dicaprylyl Carbonate, Dimethicone, Isopropyl Palmitate, Isononyl Isononanoate, Cetyl PEG/PPG-10/1 Dimethicone, C12-15 Alkyl Benzoate, Sodium Chloride, Polyhydroxystearic Acid, Tocopheryl Acetate, Triethoxycaprylylsilane, Sorbitan Sesquioleate, Phenoxyethanol, Ethylhexylglycerin, Dimethiconol, Aluminum Hydroxide, Dimethicone Crosspolymer, Stearic Acid, Xanthan Gum, Iron Oxides

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 - 32 mL Carton Label - LIGHT

NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

PURESCREEN+ TM

MINERAL

UV • TINT

FACE LIQUID

with

Vitamin E

LIGHT

30

TINTED SUNSCREEN

BROAD SPECTRUM SPF 30

UVA/UVB PROTECTION

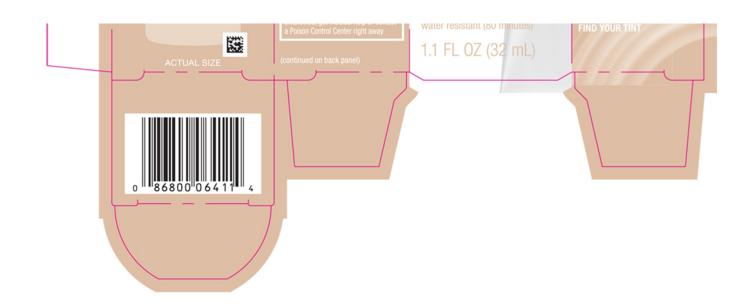
neutral undertone

for flexible coverage

water resistant (80 minutes)

1.1 FL OZ (32mL)





PRINCIPAL DISPLAY PANEL - 32 mL Carton Label - DEEP

NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

PURESCREEN+ TM

MINERAL

UV ● TINT

FACE LIQUID

with

Vitamin E

DEEP

30

TINTED SUNSCREEN

BROAD SPECTRUM SPF 30

UVA/UVB PROTECTION

neutral undertone

for flexible coverage

water resistant (80 minutes)

1.1 FL OZ (32mL)





NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

PURESCREEN+ TM

MINERAL

UV ● TINT

FACE LIQUID

with

Vitamin E

MEDIUM

30

TINTED SUNSCREEN

BROAD SPECTRUM SPF 30

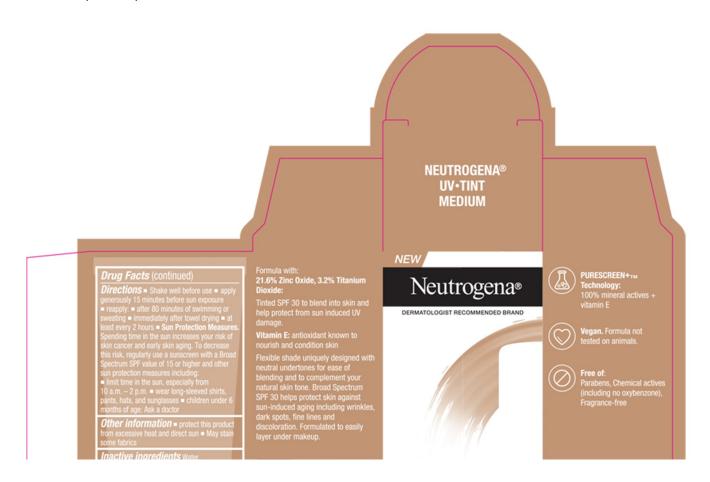
UVA/UVB PROTECTION

neutral undertone

for flexible coverage

water resistant (80 minutes)

1.1 FL OZ (32mL)





PRINCIPAL DISPLAY PANEL - 32 mL Carton Label - MEDIUM DEEP

NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

PURESCREEN+ TM

MINERAL

UV ● TINT

FACE LIQUID

with

Vitamin E

MEDIUM

DEEP

30

TINTED SUNSCREEN
BROAD SPECTRUM SPF 30
UVA/UVB PROTECTION
neutral undertone
for flexible coverage
water resistant (80 minutes)
1.1 FL OZ (32mL)

NEUTROGENA® UV-TINT MEDIUM DEEP

Drug Facts (continued)

Drug Facts (continued)

Directions = Shake well before use = apply generously 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

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

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ACTUAL SIZE

Formula with: 21.6% Zinc Oxide, 3.2% Titanium

Tinted SPF 30 to blend into skin and help protect from sun-induced UV damage. Vitamin E: antioxidant known to nourish and condition skin

nourish and condition skin
Flexible shade uniquely designed with
neutral undertones for ease of
blending and to complement your
natural skin tone. Broad Spectrum
SPF 30 helps protect skin against
sun-induced aging including wrinkles,
dark spots, fine lines and
discoloration. Formulated to easily
layer under makeup.





Drug Facts

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

NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND



PURESCREEN+TM
Technology:
100% mineral actives +



Vegan. Formula not tested on animals.



Free of: Parabens, Chemical actives (including no oxybenzone), Fragrance-free



Vitamin E

30

TINTED SUNSCREEN BROAD SPECTRUM SPF 30 **UVA/UVB PROTECTION**

neutral undertone for flexible coverage

water resistant (80 minutes)

1.1 FL OZ (32 mL)

TINT RANGE DESIGNED TO BE INCLUSIVE OF ALL SKIN TONES WITH 4 FLEXIBLE SHADES.



FIND YOUR TINT



NEUTROGENA MINERAL UV TINT FACE MEDIUM DEEP SPF 30

titanium dioxide, zinc oxide lotion

Product	Intorm	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0784

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP) TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP) TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
XANTHAN GUM (UNII: TTV12P4NEE)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968- 0784-1	1 in 1 CARTON	12/05/2022	
1		32 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69968- 0784-2	24 in 1 TRAY	03/04/2024	
_		1.4 mL in 1 POUCH; Type 0: Not a Combination		

Marketing	Information
Markating	Annlication N

Marketing Application Number or Monograph
Category Citation

Marketing Start Date Marketing End Date

OTC Monograph Drug M020 12/05/2022

NEUTROGENA MINERAL UV TINT FACE DEEP SPF 30

titanium dioxide, zinc oxide lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0783

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
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	32 mg in 1 mL

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
XANTHAN GUM (UNII: TTV12P4NEE)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69968- 0783-1	1 in 1 CARTON	12/05/2022		
1		32 mL in 1 TUBE; Type 0: Not a Combination Product			
2	NDC:69968- 0783-2	24 in 1 TRAY	03/04/2024		
2		1.4 mL in 1 POUCH; 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	12/05/2022	

NEUTROGENA MINERAL UV TINT FACE LIGHT SPF 30

titanium dioxide, zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0781
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	32 mg in 1 mL		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	216 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ISOHEXADECANE (UNII: 918X1OUF1E)		
DIMETHICONE (UNII: 92RU3N3Y10)		
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)		
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)		
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)		
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)		
XANTHAN GUM (UNII: TTV12P4NEE)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
STEARIC ACID (UNII: 4ELV7Z65AP)		
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)		

.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69968- 0781-1	1 in 1 CARTON	12/05/2022		
1		32 mL in 1 TUBE; Type 0: Not a Combination Product			
2	NDC:69968- 0781-2	24 in 1 TRAY	03/04/2024		
2		1.4 mL in 1 POUCH; 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	12/05/2022	

NEUTROGENA MINERAL UV TINT FACE MEDIUM SPF 30

titanium dioxide, zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0782
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	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	32 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ISOHEXADECANE (UNII: 918X1OUF1E)			
DIMETHICONE (UNII: 92RU3N3Y10)			
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)			
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)			

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

SODIUM CHLORIDE (UNII: 451W47IQ8X)

PHENOXYETHANOL (UNII: HIE492ZZ3T)

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)

ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)

XANTHAN GUM (UNII: TTV12P4NEE)

FERRIC OXIDE RED (UNII: 1K09F3G675)

STEARIC ACID (UNII: 4ELV7Z65AP)

TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)

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

DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)

ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)

SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)

DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:69968- 0782-1	1 in 1 CARTON	12/05/2022		
1		32 mL in 1 TUBE; Type 0: Not a Combination Product			
2	NDC:69968- 0782-2	24 in 1 TRAY	03/04/2024		
2		1.4 mL in 1 POUCH; 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	12/05/2022	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 12/2023 Johnson & Johnson Consumer Inc.