NEUTROGENA MINERAL INVISIBLE DAILY DEFENSE FACE SUNSCREEN BROAD SPECTRUM SPF 30- titanium dioxide, zinc oxide lotion
Johnson & Johnson Consumer Inc.

Neutrogena Mineral Invisible Daily Defense Face Sunscreen Broad Spectrum SPF 30

Drug Facts

Active ingredients	***Purpose***
Titanium Dioxide (4.9%)	Sunscreen
Zinc Oxide (21.6%)	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 bottle well before use.

• **For Sunscreen 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 the product from excessive heat and direct sun
- may stain some fabrics

Inactive ingredients

Water, Isononyl Isononanoate, Dicaprylyl Ether, C12-15 Alkyl Benzoate, Polyglyceryl-2 Dipolyhydroxystearate, Propylene Glycol, Dimethicone, Zingiber Officinale (Ginger) Root Extract, Tocopheryl Acetate, Chrysanthemum Parthenium (Feverfew) Flower/Leaf/Stem Juice, Calcium Sodium Borosilicate, Polyhydroxystearic Acid, Triethoxycaprylylsilane, Sorbitan Sesquioleate, Sodium Chloride, Aluminum Hydroxide, Phenoxyethanol, Triacontanyl PVP, Ethylhexylglycerin, Stearic Acid, 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 - 40 mL Bottle Label

NEW

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

PURESCREEN+TM

MINERAL

INVISIBLE

DAILY

DEFENSE

FACE LIQUID

30

SUNSCREEN

BROAD SPECTRUM SPF 30

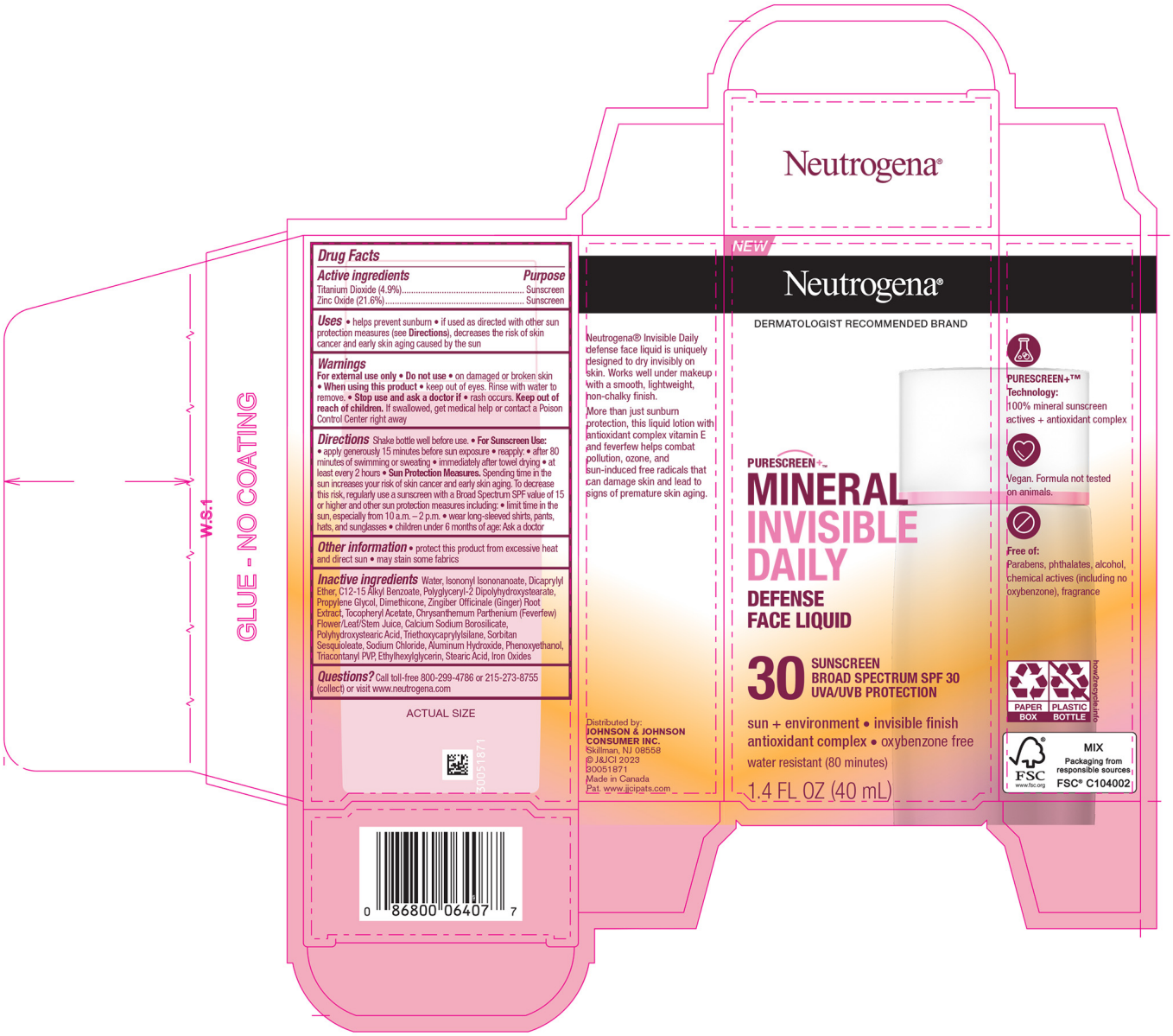
UVA/UVB PROTECTION

sun + environment • invisible finish

antioxidant complex • oxybenzone free

water resistant (80 minutes)

1.4 FL OZ (40 mL)



NEUTROGENA MINERAL INVISIBLE DAILY DEFENSE FACE SUNSCREEN BROAD SPECTRUM SPF 30

titanium dioxide, zinc oxide lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0815
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	49 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	216 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GINGER (UNII: C5529G5JPQ)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIACONTANYL PVP (WP-660) (UNII: N0SS3Q238D)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FERROUS OXIDE (UNII: G7036X8B5H)	
FEVERFEW (UNII: Z64FK7P217)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
DICAPRYLYL ETHER (UNII: 77JZM5516Z)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0815-1	1 in 1 CARTON	08/21/2023	
1		40 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:69968-0815-2	24 in 1 TRAY	01/01/2024	
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	08/21/2023	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 12/2023 Johnson & Johnson Consumer Inc.