

NEUTROGENA CLEAR FACE BREAKOUT FREE OIL FREE SUNSCREEN BROAD SPECTRUM SPF 50- avobenzone, homosalate, octisalate, and octocrylene lotion
Johnson & Johnson Consumer Inc.

Neutrogena® Clear Face BREAKOUT FREE oil-free sunscreen BROAD SPECTRUM SPF 50

Drug Facts

<i>Active Ingredients</i>	<i>Purpose</i>
Avobenzone 3%	Sunscreen
Homosalate 10%	Sunscreen
Octisalate 5%	Sunscreen
Octocrylene 10%	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 generously and evenly 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, Silica, Cetyl Dimethicone, Styrene/Acrylates Copolymer, C12-15 Alkyl Benzoate, Steareth-100, Ethylhexylglycerin, Aluminum Starch Octenylsuccinate, Phenoxyethanol, Caprylyl Glycol, Sodium Polyacrylate, Dimethicone, Steareth-2, Polyester-7, Chlorphenesin, Ethylhexyl Stearate, Disodium EDTA, Propylene Glycol, Neopentyl Glycol Diheptanoate, Bisabolol, Acrylates/Dimethicone Copolymer, Butylene Glycol, Mannan, Xanthan Gum, BHT, Capryloyl Glycine, Trideceth-6, Sarcosine, Cedrus Atlantica Bark Extract, Cinnamomum Zeylanicum Bark Extract, Portulaca Oleracea Extract

Questions?

Call toll-free **800-582-4048** 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

Clear Face

BREAKOUT FREE

oil-free

sunscreen

BROAD SPECTRUM SPF 50

50

helioplex®

broad spectrum uva.uvb

won't cause breakouts

oxybenzone free

water resistant (80 minutes)

3.0 FL OZ (88 mL)

lacquer free area

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DERMATOLOGIST RECOMMENDED BRAND

Clear Face

BREAKOUT FREE

oil-free
sunscreen

BROAD SPECTRUM SPF 50

50

helioplex®
broad spectrum uva-uvb

won't cause breakouts
oxybenzone free
water resistant (80 minutes)

3.0 FL OZ (88 mL)



- Fragrance-free
- Oil-free
- Ultra-light
- Oxybenzone Free

30050882

Drug Facts

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non-printable area

NEUTROGENA CLEAR FACE BREAKOUT FREE OIL FREE SUNSCREEN BROAD SPECTRUM SPF 50

avobenzone, homosalate, octisalate, and octocrylene lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0662
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL

OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
WATER (UNII: 059QF0KO0R)	
STEARETH-100 (UNII: 4OH5W9UM87)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
STEARETH-2 (UNII: V56DFE46J5)	
POLYESTER-7 (UNII: 0841698D2F)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
LEVOMENOL (UNII: 24WE03BX2T)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
YEAST MANNAN (UNII: 91R887N59P)	
XANTHAN GUM (UNII: TTV12P4NEE)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CAPRYLOYL GLYCINE (UNII: 8TY5YO42NJ)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
CEDRUS ATLANTICA BARK (UNII: ITP1Q41UPF)	
SARCOSINE (UNII: Z711V88R5F)	
CINNAMON BARK OIL (UNII: XE54U569EC)	
PURSLANE (UNII: M6S840WYG5)	
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0662-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	10/05/2020	
2	NDC:69968-0662-2	2 in 1 CARTON	10/05/2020	
2		88 mL in 1 TUBE; 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	10/05/2020	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2024

Johnson & Johnson Consumer Inc.