

NEUTROGENA CLEAR FACE BREAKOUT FREE OIL FREE SUNSCREEN BROAD SPECTRUM SPF 30- avobenzone, homosalate, octisalate, and octocrylene lotion
Kenvue Brands LLC

Neutrogena® Clear Face BREAKOUT FREE oil-free sunscreen BROAD SPECTRUM SPF 30

Drug Facts

<i>Active ingredients</i>	<i>Purpose</i>
Avobenzone 2.5%	Sunscreen
Homosalate 8%	Sunscreen
Octisalate 5%	Sunscreen
Octocrylene 8%	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, Ethylhexylglycerin, Steareth-100, Aluminum Starch Octenylsuccinate, Phenoxyethanol, Caprylyl Glycol, Sodium Polyacrylate, Dimethicone, Polyester-7, Chlorphenesin, Steareth-2, Ethylhexyl Stearate, Disodium EDTA, Propylene Glycol, Neopentyl Glycol Diheptanoate, Bisabolol, Acrylates/Dimethicone Copolymer, Butylene Glycol, BHT, Mannan, Xanthan Gum, 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:

Kenvue Brands LLC

Summit, NJ 07901

PRINCIPAL DISPLAY PANEL - 88 mL Tube Label

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

Clear Face

BREAKOUT FREE

oil-free

sunscreen

BROAD SPECTRUM SPF 30

30

helioplex®

broad spectrum uva•uvb

won't cause breakouts

oxybenzone free

water resistant (80 minutes)

3.0 FL OZ (88 mL)

lacquer free area

Neutrogena[®]

DERMATOLOGIST RECOMMENDED BRAND

Clear Face

BREAKOUT FREE

oil-free
sunscreen

BROAD SPECTRUM SPF 30

30


helioplex[®]
broad spectrum uva-uvb

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

3.0 FL OZ (88 mL)

non-printable area



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Drug Facts

Active ingredients	Purpose
Avobenzone (2.5%), Homosalate (8%), Octisalate (5%), Octocrylene (8%)	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

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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

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Pat. www.kenvuepats.com

NEUTROGENA CLEAR FACE BREAKOUT FREE OIL FREE SUNSCREEN BROAD SPECTRUM SPF 30

avobenzone, homosalate, octisalate, and octocrylene lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0571
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	25 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	80 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	80 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
STEARETH-100 (UNII: 4OH5W9UM87)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
POLYESTER-7 (UNII: 0841698D2F)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
STEARETH-2 (UNII: V56DFE46J5)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
LEVOMENOL (UNII: 24WE03BX2T)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
YEAST MANNAN (UNII: 91R887N59P)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CAPRYLOYL GLYCINE (UNII: 8TY5YO42NJ)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
SARCOSINE (UNII: Z711V88R5F)	
CEDRUS ATLANTICA BARK (UNII: ITP1Q41UPF)	
CINNAMON BARK OIL (UNII: XE54U569EC)	
PURSLANE (UNII: M6S840WYG5)	
CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0571-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	10/07/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/07/2019	

Labeler - Kenvue Brands LLC (118772437)

Revised: 8/2025

Kenvue Brands LLC