

**NEUTROGENA ULTRA SHEER DRY-TOUCH SUNSCREEN MAX BROAD SPECTRUM
SPF 100 PLUS- avobenzone, homosalate, octisalate, octocrylene lotion
Kenvue Brands LLC**

Neutrogena Ultra Sheer Dry-Touch Sunscreen MAX Broad Spectrum SPF 100 Plus

Drug Facts

Active ingredients

Avobenzone (3%), Homosalate (15%),
Octisalate (5%), Octocrylene (10%)

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

For Sunscreen Use:

- apply generously 15 minutes before sun exposure
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- **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
- protect from freezing

Inactive ingredients

Water, Ethylhexyl Methoxycrylene, Styrene/Acrylates Copolymer, Copernicia Cerifera (Carnauba) Wax, Hydrogenated Castor Oil, Silica, Triaccontanyl PVP, Potassium Cetyl Phosphate, Caprylyl Methicone, Aluminum Starch Octenylsuccinate, Glyceryl Stearate, PEG-100 Stearate, Cetyl Dimethicone, Dimethicone, Phenoxyethanol, Caprylyl Glycol, Ethylhexylglycerin, Acrylates/Dimethicone Copolymer, Xanthan Gum, Chlorphenesin, Sodium Polyacrylate, Hydrolyzed Jojoba Esters, Disodium EDTA, Ethylhexyl Stearate, Tocopheryl Acetate, Jojoba Esters, Trideceth-6, Chrysanthemum Parthenium (Feverfew) Flower/Leaf/Stem Juice

Questions?

Call toll-free **800-299-4786** 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

ULTRA

SHEER®

Dry-Touch

Sunscreen

max

100+

BROAD SPECTRUM SPF 100+

With Antioxidants

Lightweight clean feel

Blocks up to 99% of UVB rays

with HELIOPLEX360™

3.0 FL OZ (88 mL)

**Neutrogena®
Ultra Sheer®
with HELIOPLEX360™**



30058711

advanced formula combines superior UVA/UVB protection with antioxidants to help defend against free radicals from blue light exposure. Highest SPF level within Ultra Sheer® line

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Other information • protect this product from excessive heat and direct sun • may stain some fabrics • protect from freezing

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Ethylhexyl Methoxycrylene, Styrene/Acrylates
Copolymer, Copernicia Cerifera (Carnaubá) Wax,
Hydrogenated Castor Oil, Silica, Triacentauryl PVP,
Potassium Cetyl Phosphate, Caprylyl Methicone,
Aluminum Starch Octenylsuccinate, Glyceryl
Stearate, PEG-100 Stearate, Cetyl Dimethicone,
Dimethicone, Phenoxethanol, Caprylyl Glycol,
Ethylhexylglycerin, Acrylates/Dimethicone
Copolymer, Xanthan Gum, Chlorphenesin,
Sodium Polycarboxylate, Hydrolyzed Jojoba Esters,
Disodium EDTA, Ethylhexyl Stearate, Tocopheryl
Acetate, Jojoba Esters, Trideceth-6,
Chrysanthemum Parthenium (Feverfew)
Flower/leaf/Stem Juice

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Free of: Oxybenzone, PABA, parabens, phthalates and dyes



**Dermatologist
Tested**



HELIOPLEX360™: UVA/UVB
protection + antioxidants



Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

**ULTRA
SHEER®
Dry-Touch
Sunscreen**

max 100+
BROAD SPECTRUM SPF 100+

With Antioxidants

Lightweight clean feel
Blocks up to 99% of UVB rays

with HELIOPLEX360™

3.0 FL OZ (88 mL)

NEUTROGENA ULTRA SHEER DRY-TOUCH SUNSCREEN MAX BROAD SPECTRUM SPF 100 PLUS

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0936
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
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	150 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
WATER (UNII: 059QF0KO0R)	
TRIACONTANYL PVP (UNII: N0SS3Q238D)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
XANTHAN GUM (UNII: TTV12P4NEE)	
JOJOBA OIL, RANDOMIZED (UNII: 7F0EV20QYL)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
CETYL DIMETHICONE 25 (UNII: U4AS1BW4ZB)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TANACETUM PARTHENIUM WHOLE (UNII: 6GE7Z0761K)	
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
SILICA (UNII: ETJ7Z6XBU4)	
HYDROLYZED JOJOBA ESTERS (UNII: UDR641JW8W)	
COPERNICIA CERIFERA (CARNAUBA) WAX (UNII: R12CBM0EIZ)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0936-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	08/18/2025	
2	NDC:69968-0936-1	12 in 1 TRAY	08/18/2025	

2	14 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	08/18/2025	

Labeler - Kenvue Brands LLC (118772437)

Revised: 8/2025

Kenvue Brands LLC