

**NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM  
SPF45- avobenzene, homosalate, octisalate, and octocrylene lotion  
Kenvue Brands LLC**

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**Neutrogena<sup>®</sup> Ultra Sheer<sup>®</sup> dry-touch sunscreen BROAD SPECTRUM SPF45**

***Drug Facts***

<b><i>Active ingredients</i></b>	<b><i>Purpose</i></b>
Avobenzene 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 liberally 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, Styrene/Acrylates Copolymer, Silica, Dimethicone, Potassium Cetyl Phosphate, Benzyl Alcohol, Beeswax, Caprylyl Methicone, Glyceryl Stearate, PEG-100 Stearate, Cetyl Dimethicone, Caprylyl Glycol, Ethylhexylglycerin, Aluminum Starch Octenylsuccinate, Behenyl Alcohol, Acrylates/Dimethicone Copolymer, Xanthan Gum, Sodium Polyacrylate, Chlorphenesin, Dimethicone PEG - 10/15 Crosspolymer, Hydrolyzed Jojoba Esters, Fragrance, Disodium EDTA, Ethylhexyl Stearate, Tocopheryl Acetate, BHT, Trideceth-6, Jojoba Esters

## **Questions or Comments?**

Call toll-free **800-299-4786** or **215-273-8755** (collect). [www.neutrogena.com](http://www.neutrogena.com)

Distributed by:

**Kenvue Brands LLC**

Summit, NJ 07901

## **PRINCIPAL DISPLAY PANEL - 88 mL Tube Label**

*NEW LOOK*

**Neutrogena**®

**DERMATOLOGIST RECOMMENDED BRAND**

**ULTRA**

**SHEER**®

**Dry-Touch**

**Sunscreen**

**45**

**BROAD SPECTRUM SPF 45**

**Antioxidant Vitamin E**

Lightweight clean feel

Shields skin 6 layers deep

**with helioplex**®

Water resistant (80 minutes)

3.0 FL OZ (88 mL)

**Neutrogena®  
Ultra Sheer®  
with Helioplex®**



30059537

provides superior broad-spectrum protection against skin-aging UVA and burning UVB rays with a non-greasy, invisible finish.

**Drug Facts**

**Active ingredients** **Purpose**  
Avobenzone (3%), Homosalate (10%),  
Octisalate (5%), 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

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**Other information** ■ protect this product from excessive heat and direct sun ■ may stain some fabrics

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



Dermatologist tested



HELIOPLEX®: UVA/UVB protection + antioxidant vitamin E



**NEW LOOK**

**Neutrogena®**

DERMATOLOGIST RECOMMENDED BRAND

**ULTRA  
SHEER®  
Dry-Touch  
Sunscreen**

**45**

**BROAD SPECTRUM SPF 45**

**Antioxidant Vitamin E**

Lightweight clean feel  
Shields skin 6 layers deep

with **HELIOPLEX®**  
Water resistant (80 minutes)

3.0 FL OZ (88 mL)

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**NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF45**

avobenzone, homosalate, octisalate, and octocrylene lotion

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69968-0553
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 mL
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER</b> (UNII: V5RS026Q0H)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>POTASSIUM CETYL PHOSPHATE</b> (UNII: 03KCY6P7UT)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>YELLOW WAX</b> (UNII: 2ZA36H0S2V)	
<b>CAPRYLYL TRISILOXANE</b> (UNII: Q95M2P1KJL)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>ALUMINUM STARCH OCTENYLSUCCINATE</b> (UNII: I9PJ006294)	
<b>DOCOSANOL</b> (UNII: 9G1OE216XY)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>CHLORPHENESIN</b> (UNII: I670DAL4SZ)	
<b>HYDROLYZED JOJOBA ESTERS (ACID FORM)</b> (UNII: UDR641JW8W)	
<b>EDETATE DISODIUM ANHYDROUS</b> (UNII: 8NLQ36F6MM)	
<b>ETHYLHEXYL STEARATE</b> (UNII: EG3PA2K3K5)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)	
<b>TRIDECETH-6</b> (UNII: 3T5PCR2H0C)	
<b>JOJOBA OIL, RANDOMIZED</b> (UNII: 7F0EV20QYL)	
<b>CETYL DIMETHICONE 25</b> (UNII: U4AS1BW4ZB)	
<b>DIMETHICONE/PEG-10/15 CROSSPOLYMER</b> (UNII: 21AS8B1BSS)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW)</b> (UNII: S7ZA3CCJ4M)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0553-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	10/07/2019	
2	NDC:69968-0553-2	2 in 1 PACKAGE	10/07/2019	
3		88 mL in 1 TUBE; Type 0: Not a Combination		

### 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: 3/2025

Kenvue Brands LLC