

**NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF 55 CLUBTRAY- avobenzone, homosalate, octisalate, and octocrylene**  
Kenvue Brands LLC

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**Neutrogena Ultra Sheer Dry Touch Sunscreen Broad Spectrum SPF 55 - Club**

**Neutrogena® Ultra Sheer® dry-touch sunscreen BROAD SPECTRUM SPF55**

***Drug Facts***

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***Active ingredients***

Avobenzone 3%  
Homosalate 10%  
Octisalate 5%  
Octocrylene 10%

***Purpose***

Sunscreen  
Sunscreen  
Sunscreen  
Sunscreen

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**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?**

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

Distributed by:

### **JOHNSON & JOHNSON CONSUMER INC.**

Skillman, NJ 08558

### **PRINCIPAL DISPLAY PANEL - Kit Package Label**

Neutrogena®

#1 Dermatologist recommended suncare brand

Powerful sun protection,

unbelievably light feel

ACTUAL

SIZES

OXYBENZONE

& OCTINOXATE FREE

helioplex®

broad spectrum uva•uvb

NEUTROGENA® ULTRA SHEER® Sunscreen Lotion Broad Spectrum SPF 55

5.0 FL OZ (147 mL) and 3.0 FL OZ (88 mL), TOTAL 8.0 FL OZ (235 mL)



# NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF 55 CLUBTRAY

avobenzone, homosalate, octisalate, and octocrylene kit

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69968-0847
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## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0847-9	1 in 1 PACKAGE	12/08/2023	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 TUBE	147 mL
Part 2	1 TUBE	88 mL

## Part 1 of 2

# NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF55

avobenzone, homosalate, octisalate, and octocrylene lotion

**Product Information****Item Code (Source)** NDC:69968-0576**Route of Administration** TOPICAL**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>ALPHA-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>EDETATE DISODIUM ANHYDROUS</b> (UNII: 8NLQ36F6MM)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>JOJOBA OIL, RANDOMIZED</b> (UNII: 7F0EV20QYL)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>ALUMINUM STARCH OCTENYLSUCCINATE</b> (UNII: I9PJ0O6294)	
<b>DIMETHICONE/PEG-10/15 CROSSPOLYMER</b> (UNII: 21AS8B1BSS)	
<b>ETHYLHEXYL STEARATE</b> (UNII: EG3PA2K3K5)	
<b>CETYL DIMETHICONE 25</b> (UNII: U4AS1BW4ZB)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>TRIDECETH-6</b> (UNII: 3T5PCR2H0C)	
<b>2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW)</b> (UNII: S7ZA3CCJ4M)	
<b>DOCOSANOL</b> (UNII: 9G1OE216XY)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>YELLOW WAX</b> (UNII: 2ZA36H0S2V)	
<b>HYDROLYZED JOJOBA ESTERS (ACID FORM)</b> (UNII: UDR641JW8W)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>CHLORPHENESIN</b> (UNII: I670DAL4SZ)	
<b>BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER</b> (UNII: V5RS026Q0H)	
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>POTASSIUM CETYL PHOSPHATE</b> (UNII: 03KCY6P7UT)	
<b>CAPRYLYL TRISILOXANE</b> (UNII: Q95M2P1KJL)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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1	NDC:69968-0576-5	147 mL in 1 TUBE; Type 0: Not a Combination Product	
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## Marketing Information

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

## Part 2 of 2

### NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF55

avobenzone, homosalate, octisalate, and octocrylene lotion

## Product Information

Item Code (Source)	NDC:69968-0576
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
<b>2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW)</b> (UNII: S7ZA3CCJ4M)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>EDETATE DISODIUM ANHYDROUS</b> (UNII: 8NLQ36F6MM)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>JOJOBA OIL, RANDOMIZED</b> (UNII: 7F0EV20QYL)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>ALUMINUM STARCH OCTENYLSUCCINATE</b> (UNII: I9PJ006294)	
<b>DIMETHICONE/PEG-10/15 CROSSPOLYMER</b> (UNII: 21AS8B1BSS)	
<b>ETHYLHEXYL STEARATE</b> (UNII: EG3PA2K3K5)	
<b>CETYL DIMETHICONE 25</b> (UNII: U4AS1BW4ZB)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>TRIDECETH-6</b> (UNII: 3T5PCR2H0C)	
<b>DOCOSANOL</b> (UNII: 9G1OE216XY)	

<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)
<b>YELLOW WAX</b> (UNII: 2ZA36H0S2V)
<b>HYDROLYZED JOJOBA ESTERS (ACID FORM)</b> (UNII: UDR641JW8W)
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)
<b>CHLORPHENESIN</b> (UNII: I670DAL4SZ)
<b>BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER</b> (UNII: V5RS026Q0H)
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>POTASSIUM CETYL PHOSPHATE</b> (UNII: 03KCY6P7UT)
<b>CAPRYLYL TRISILOXANE</b> (UNII: Q95M2P1KJL)
<b>ALPHA-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0576-3	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	10/07/2019	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/08/2023	

**Labeler** - Kenvue Brands LLC (118772437)