# NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF70-avobenzone, homosalate, octisalate, octocrylene, and oxybenzone lotion Johnson & Johnson Consumer Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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### Neutrogena® Ultra Sheer® dry-touch sunscreen BROAD SPECTRUM SPF 70 Drug Facts

Active ingredients	Purpose
Avobenzone 3%	Sunscreen
Homosalate 15%	Sunscreen
Octisalate 5%	Sunscreen
Octocrylene 2.8%	Sunscreen
Oxybenzone 6%	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
- protect from freezing

### **Inactive ingredients**

Water, Styrene/Acrylates Copolymer, Silica, Diethylhexyl 2,6-Naphthalate, Beeswax, Caprylyl Methicone, Cetyl Dimethicone, Ethylhexylglycerin, Glyceryl Stearate, PEG-100 Stearate, Sodium Polyacrylate, Acrylates/C12-22 Alkyl Methacrylate Copolymer, Dimethicone, Xanthan Gum, Trimethylsiloxysilicate, Disodium EDTA, Fragrance, Dipotassium Glycyrrhizate, BHT, Methylisothiazolinone, Polyaminopropyl Biguanide

### Questions?

Call toll-free 800-299-4786 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® #1 DERMATOLOGIST RECOMMENDED BRAND Ultra Sheer® dry-touch suns creen

BROAD SPECTRUM SPF 70 70

helioplex<sup>®</sup> broad spectrum uva•uvb

lightweight clean feel fast drying water resistant (80 minutes)

3.0 FL OZ (88mL)

# Neutrogena<sup>®</sup>

#1 DERMATOLOGIST RECOMMENDED BRAND

# Ultra Sheer®

sunscreen

**BROAD SPECTRUM SPF 70** 

70



lightweight clean feel fast drying water resistant (80 minutes)

3.0 FL OZ (88 mL)



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Neutrogena® Ultra Sheer® with Helioplex® provides superior broad-spectrum protection against skin-aging UVA and burning UVB rays with a lightweight, clean feel.

### Drug Facts

### Active ingredients

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## NEUTROGENA ULTRA SHEER DRY TOUCH SUNSCREEN BROAD SPECTRUM SPF70

avobenzone, homosalate, octisalate, octocrylene, and oxybenzone lotion

### Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0168

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)	Ho mo sa la te	150 mg in 1 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	50 mg in 1 mL

	Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	28 mg in 1 mL
ı	Oxybenzone (UNII: 9500S7VE0Y) (Oxybenzone - UNII:9500S7VE0Y)	Oxybenzone	60 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
Yellow Wax (UNII: 2ZA36H0S2V)		
CETYL DIMETHICO NE 25 (UNII: U4AS1BW4ZB)		
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)		
Caprylyl Trisiloxane (UNII: Q95M2P1KJL)		
Diethylhexyl 2,6-Naphthalate (UNII: I0 DQJ7YGXM)		
Dimethicone (UNII: 92RU3N3Y1O)		
Glycyrrhizinate Dipotassium (UNII: CA2Y0FE3FX)		
Edetate Disodium (UNII: 7FLD91C86K)		
Ethylhexylglycerin (UNII: 147D247K3P)		
Glyceryl Monostearate (UNII: 230 O U9 XXE4)		
Methylisothiazolinone (UNII: 229 D0 E1QFA)		
PEG-100 Stearate (UNII: YD01N1999R)		
Polihexanide (UNII: 322U039GMF)		
Silicon Dioxide (UNII: ETJ7Z6XBU4)		
Water (UNII: 059QF0KO0R)		
Xanthan Gum (UNII: TTV12P4NEE)		
Polyaminopropyl Biguanide (UNII: DT9D8Z79ET)		

F	Packaging			
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:69968-0168-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	0 1/0 1/20 0 7	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	0 1/0 1/20 0 7	

### Labeler - Johnson & Johnson Consumer Inc. (002347102)

Revised: 10/2019 Johnson & Johnson Consumer Inc.