

**NEUTROGENA ULTRA SHEER FACE SUNSCREEN SPF 60- avobenzone, homosalate, octisalate, octocrylene stick
KDC/One Development Corporation, Inc.**

Neutrogena Ultra Sheer Face Stick Sunscreen SPF 60

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

- apply liberally and evenly 15 minutes before sun exposure
- ensure complete coverage to the area above the lip, nose and tops of ears
- 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
- package contains small parts, which could cause a choking hazard
- Protect from freezing

Inactive ingredients

Neopentyl Glycol Diethylhexanoate, Octyldodecyl Neopentanoate, Synthetic Wax, Isohexadecane, Butyloctyl Salicylate, Ozokerite, Paraffin, Synthetic Beeswax, C12-15 Alkyl Benzoate, VP/Eicosene Copolymer, Neopentyl Glycol Diisostearate, Dimethicone, Diethylhexyl 2,6-Naphthalate, Styrene/Acrylates Copolymer, Tocopheryl Acetate, Ascorbyl Palmitate

Questions or comments?

Call toll-free **800-299-4786** or **215-273-8755**(collect) or visit www.neutrogena.com

Distributed by:

JOHNSON & JOHNSON

CONSUMER INC.

Skillman, NJ 08558

Neutrogena Ultra Sheer Face Stick Sunscreen SPF 60 - 13 g Canister in Blister Pack Label

New

Neutrogena ®

DERMATOLOGIST RECOMMENDED BRAND

ULTRA SHEER ®

Face Stick

Sunscreen

60

BROAD

SPECTRUM

SPF 60

Antioxidants

Vitamin C & E

Invisible,

non-greasy

Shields skin

6 layers deep

with

HELIOPLEX®

Fragrance Free

For sensitive skin

Water resistant

(80 minutes)

NET WT

0.47 OZ (13g)

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Face Stick

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BROAD SPECTRUM SPF 60

Antioxidants Vitamin C & E

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Shields skin 6 layers deep

with HELIOPLEX®

Fragrance Free
For sensitive skin
Water resistant (80 minutes)

NET WT
0.47 OZ (13 g)

Neutrogena® Ultra Sheer® with Helioplex® provides superior broad-spectrum protection against skin-aging UVA and burning UVB rays. This gentle formula is ideal for sensitive skin.

- Won't transfer or smudge • Dries instantly
- Spreads on easily • Mess-free, lightweight

Drug Facts

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Other information • protect this product from excessive heat and direct sun • may stain some fabrics • package contains small parts, which could cause a choking hazard • Protect from freezing

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Free of: Fragrance, oxybenzone, parabens and phthalates

Dermatologist tested

HELIOPLEX®: UVA/UVB protection + antioxidants vitamin C & E

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From Curd
Check Company
PAPER CARD
PLASTIC TRAY
PLASTIC STICK
No commingling at all

NEUTROGENA ULTRA SHEER FACE SUNSCREEN SPF 60

avobenzone, homosalate, octisalate, octocrylene stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54565-1645
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZ ONE	30 mg in 1 g

HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)

HOMOSALATE

150 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)	
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)	
CERESIN (UNII: Q1LS2UJO3A)	
SYNTHETIC BEESWAX (UNII: 08MNR5YE2R)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
PARAFFIN (UNII: I9O0E3H2ZE)	
NEOPENTYL GLYCOL DIISOSTEARATE (UNII: 4M6OQ34JWW)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
DIETHYLHEXYL 2,6-NAPHTHALATE (UNII: I0DQJ7YGXM)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54565-1645-2	1 in 1 BLISTER PACK	10/04/2024	
1	NDC:54565-1645-1	13 g in 1 CANISTER; 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	10/04/2024	

Labeler - KDC/One Development Corporation, Inc. (204006464)**Establishment**

Name	Address	ID/FEI	Business Operations
KDC/One Development Corporation, Inc		204006464	analysis(54565-1645) , manufacture(54565-1645) , pack(54565-1645) , label(54565-1645)

Revised: 1/2026

KDC/One Development Corporation, Inc.