

**NEUTROGENA INVISIBLE DAILY DEFENSE BROAD SPECTRUM SPF 60 PLUS-
avobenzene, homosalate, octisalate, and octocrylene lotion
Kenvue Brands LLC**

Neutrogena Invisible Daily Defense Lotion Broad Spectrum SPF 60 Plus

Drug Facts

<i>Active ingredients</i>	<i>Purpose</i>
Avobenzene 3%	Sunscreen
Homosalate 13.5%	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, butyloctyl salicylate, glycerin, alcohol denat., silica, dimethicone, aluminum starch octenylsuccinate, polyurethane-62, phenoxyethanol, pentylene glycol, sodium acryloyldimethyltaurate/VP crosspolymer, acrylates/dimethicone copolymer, glyceryl stearate, chlorphenesin, styrene/acrylates copolymer, fragrance, menthyl lactate, tocopheryl acetate, disodium EDTA, trideceth-6, chrysanthemum parthenium (feverfew) flower/leaf/stem juice, sodium hydroxide, yellow 5, red 40

Questions?

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

PRINCIPAL DISPLAY PANEL - 88 mL Tube Label

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

Invisible Daily

defense

lotion

sunscreen

BROAD SPECTRUM SPF 60+

60+

sun + environment

oxybenzone free

water resistant (80 minutes)

3.0 FL OZ (88 mL)

lacquer free area



30050914

Neutrogena Invisible Daily™ defense with Helioplex® + technology provides superior broad spectrum UVA/UVB protection. Antioxidants help fight environmental aggressors such as cold, ozone and pollution. Rubs in invisible on all skin tones.

Neutrogena®

DERMATOLOGIST RECOMMENDED BRAND

Invisible Daily defense

lotion

sunscreen

BROAD SPECTRUM SPF 60+

60+

sun + environment
oxybenzone free
water resistant (80 minutes)

3.0 FL OZ (88 mL)

non-printable area

Distributed by: JOHNSON & JOHNSON CONSUMER INC., Skillman, NJ 08558 © J&J 2022 Made in Canada Pat. www.jjcpats.com

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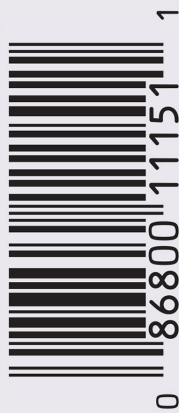
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0841
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)	HOMOSALATE	135 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
METHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
FEVERFEW (UNII: Z64FK7P217)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
POLYURETHANE-62 (UNII: TBK645J3J8)	
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0841-3	88 mL in 1 TUBE; Type 0: Not a Combination Product	01/31/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/31/2024	

Labeler - Kenvue Brands LLC (118772437)

