NEUTROGENA HEALTHY DEFENSE DAILY MOISTURIZER SUNSCREEN BROAD SPECTRUM SPF 50- 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.

Neutrogena ® HEALTHY DEFENSE ® daily moisturizer with sunscreen Broad Spectrum SPF50

Drug Facts

Active ingredients	***Purpose***
Avobenzone 3%	Sunscreen
Homosalate 12%	Sunscreen
Octisalate 5%	Sunscreen
Octocrylene 2.35%	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

For Sunscreen Use:
- apply generously 15 minutes before sun exposure
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- **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

Inactive ingredients

Water, Dimethicone, Diethylhexyl 2,6-Naphthalate, Glycerin, Trisiloxane, PEG-100 Stearate, Glyceryl Stearate, Cetearyl Alcohol, Behenyl Alcohol, Potassium Cetyl Phosphate, Caprylyl Methicone, Styrene/Acrylates Copolymer, Hydrogenated Palm Glycerides, Benzyl Alcohol, Ethylhexylglycerin, Methylparaben, Cetearyl Glucoside, Xanthan Gum, Propylparaben, Tocopherol, Disodium EDTA, Polymethyl Methacrylate, BHT, Methylisothiazolinone, Tocopheryl Acetate, Ascorbic Acid, Retinyl Palmitate, Pantothenic Acid

Questions?

Call toll-free **866-638-4636** or **215-273-8755** (collect). www.neutrogena.com

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 50 mL Tube Carton

HEALTHY
DEFENSE ®

daily
moisturizer
with
sunscreen

Broad Spectrum
SPF 50

helioplex ®
broad spectrum uva•uvb

Lightweight, non-greasy moisturizer

Neutrogena ®

#1 DERMATOLOGIST RECOMMENDED
FACIAL MOISTURE

1.7 FL OZ (50mL)

NEUTROGENA HEALTHY DEFENSE DAILY MOISTURIZER SUNSCREEN BROAD SPECTRUM SPF 50

avobenzone, homosalate, octisalate, octocrylene, and oxybenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0627
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	120 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	23.5 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	60 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TRISILOXANE (UNII: 9G1ZW13R0G)	
DIETHYLHEXYL 2,6-NAPHTHALATE (UNII: I0DQJ7YGXM)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
DOCOSANOL (UNII: 9G1OE216XY)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PANTOTHENIC ACID (UNII: 19F5HK2737)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TOCOPHEROL (UNII: R0ZB2556P8)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
POLY(METHYL METHACRYLATE; 450000 MW) (UNII: Z47NNT4J11)	
HYDROGENATED PALM GLYCERIDES (UNII: YCZ8EM144Q)	
BUTYL METHACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID/STYRENE CROSSPOLYMER (UNII: V5RS026Q0H)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0627-1	1 in 1 CARTON	01/31/2020	08/02/2022
1		14 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69968-0627-2	1 in 1 CARTON	01/31/2020	07/20/2024
2		50 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 not final	part352	01/31/2020	07/20/2024

Labeler - Johnson & Johnson Consumer Inc. (118772437)

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