NEUTROGENA WET SKIN SWIM HUMIDITY SWEAT SUNSCREEN BROAD SPECTRUM SPF50- avobenzone, homosalate, octisalate, octocrylene, and oxybenzone aerosol, spray
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® wet skin SWIM.HUMIDITY.SWEAT SUNSCREEN Broad Spectrum SPF 50

Drug Facts

Active ingredients

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

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 away from face to avoid breathing it
  - 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.
- **Danger: Flammable. Contents under pressure.** Do not use near fire, heat or while smoking. Do not puncture or incinerate. Store at temperature below 120°F (48°C). Use only as directed. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal.

Directions

- shake well before and frequently during use
- spray liberally and spread evenly by hand 15 minutes before sun exposure
- apply to all skin exposed to the sun
- hold container 4-6 inches from the skin to apply. Rub in.
- do not spray directly into face. Spray on hands and then apply to face. Rub in.

- do not apply in windy conditions
- use in a well-ventilated area
- 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

alcohol denat., dimethyl ether, octyldodecyl citrate crosspolymer, ethyl methicone, acrylates/octylacrylamide copolymer, dimethicone, acrylates/dimethicone copolymer, fragrance, tocopheryl acetate, nelumbo nucifera flower wax, diethylhexyl 2,6-naphthalate, cetyl dimethicone/bis-vinyldimethicone crosspolymer, ascorbyl palmitate, retinyl palmitate, tocopherol

Questions?

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

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

PRINCIPAL DISPLAY PANEL - 141 g Can Label

Neutrogena®
wet skin
SWIM-HUMIDITY-SWEAT
SUNSCREEN
Broad
Spectrum
SPF 50
helioplex®
broad spectrum uva-uvb

full strength protection
on wet or dry skin
water resistant (80 minutes)

#1 DERMATOLOGIST
RECOMMENDED SUNCARE

NET WT 5.0 OZ (141 g)

NEUTROGENA WET SKIN SWIM HUMIDITY SWEAT SUNSCREEN BROAD SPECTRUM SPF50

avobenzone, homosalate, octisalate, octocrylene, and oxybenzone aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0177
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 g

Homosalate (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Homosalate	100 mg in 1 g
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	50 mg in 1 g
Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	100 mg in 1 g
Oxybenzone (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
Alcohol (UNII: 3K9958V90M)	
Ascorbyl Palmitate (UNII: QN83US2B0N)	
Diethylhexyl 2,6-Naphthalate (UNII: I0DQJ7YGXM)	
Dimethicone (UNII: 92RU3N3Y1O)	
Dimethyl Ether (UNII: AM13FS69BX)	
Ethyl Methicone (8 Mpa.S) (UNII: 3YWG8XYT8H)	
Nelumbo Nucifera Flower Wax (UNII: U01S6C427I)	
Vitamin A Palmitate (UNII: 1D1K0N0VVC)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0177-5	141 g in 1 CAN; Type 0: Not a Combination Product	10/01/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	10/01/2010	

Labeler - Johnson & Johnson Consumer Inc. (002347102)

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