

**NEUTROGENA WET SKIN KIDS BEACH AND POOL SUNBLOCK SPF 70PLUS  
HELIOPLEX- 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 Kids Beach & Pool Sunblock Spray 70+**

**Directions**

For best results, apply 15-30 minutes before sun exposure. Hold Can 4-6 inches away from body and apply liberally, spraying slowly and evenly until product is visible on skin. Can be applied directly to wet skin. Reapply after swimming, excessive perspiration, towel drying or extended sun exposure. Do not apply in windy conditions. Do not spray into face. Spray into hand and apply to the face. Use in well ventilated areas.

**Warnings**

**For external use only.**

Not to be swallowed. Avoid contact with eyes.

Discontinue use if irritation or rash appear.

Use of children under 6 months of age only with advice of a physician.

**Keep out of reach of children.** If case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**Caution**

**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.

**Questions or comments?**

**1-800-299-4789** or [www.neutrogena.com](http://www.neutrogena.com)

**Active Ingredients**

Avobenzone 3%, Homosalate 15%, Octisalate 5%, Octocrylene 10%, Oxybenzone 6%

**Inactive Ingredients**

Alcohol Denat.

Dimethyl Ether

Octyldodecyl Citrate Crosspolymer

Acrylates/Octylacrylamide Copolymer

Ethyl Methicone

Cetyl Dimethicone/Bis-Vinyl Dimethicone Crosspolymer

Dimethicone

Acrylates/Dimethicone Copolymer  
Fragrance  
Tocopheryl Acetate  
Nelumbo Nucifera Flower Wax  
Diethylhexyl 2,6-Naphthalate  
Ascorbyl Palmitate  
Retinyl Palmitate

**PRINCIPAL DISPLAY PANEL - 141 g Can Label**

***NEW***

**Neutrogena®**

**wet skin**

**kids**

**BEACH & POOL**

sunblock spray

**SPF 70+**

**helioplex®**

**broad spectrum uva•uvb**

**full strength protection**

**even on wet skin**

applies to wet or dry skin

hypoallergenic, waterproof

**#1 DERMATOLOGIST**

**RECOMMENDED SUNCARE**

NET WT 5.0 OZ (141 g)

**NEW**

**Neutrogena®**

wet skin  
**kids**

**BEACH & POOL**

sunblock spray



**helloplex®**  
broad spectrum UVA-UVB

full strength protection  
even on wet skin  
applies to wet or dry skin  
hypoallergenic, waterproof

**#1 DERMATOLOGIST  
RECOMMENDED SUNCARE**

NET WT 5.0 OZ (141 g)

# NEUTROGENA WET SKIN KIDS BEACH AND POOL SUNBLOCK SPF 70PLUS HELIOPLEX

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

## Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10812-966
Route of Administration	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>Avobenzone</b> (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	30 mg in 1 g
<b>Homosalate</b> (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Homosalate	150 mg in 1 g
<b>Octisalate</b> (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	50 mg in 1 g
<b>Octocrylene</b> (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	100 mg in 1 g
<b>Oxybenzone</b> (UNII: 95OOS7VE0Y) (Oxybenzone - UNII:95OOS7VE0Y)	Oxybenzone	60 mg in 1 g

## Inactive Ingredients

Ingredient Name	Strength
<b>Alcohol</b> (UNII: 3K9958V90M)	
<b>Dimethyl Ether</b> (UNII: AM13FS69BX)	
<b>Citric Acid Monohydrate</b> (UNII: 2968PHW8QP)	
<b>Dimethicone</b> (UNII: 92RU3N3Y1O)	
<b>.alpha.-tocopherol acetate, dl-</b> (UNII: WR1WPI7EW8)	
<b>Ascorbyl Palmitate</b> (UNII: QN83US2B0N)	
<b>Vitamin A Palmitate</b> (UNII: 1D1K0N0VVC)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10812-966-01	141 g in 1 CAN; Type 0: Not a Combination Product	12/01/2010	12/01/2023

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part352	12/01/2010	12/01/2023

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

Revised: 1/2019

Johnson & Johnson Consumer Inc.