

**NEOVA DNA DAMAGE CONTROL - EVERYDAY BROAD SPECTRUM SPF 44-
octinoxate, octisalate, zinc oxide emulsion
PHARMA COSMETICS, INC**

Neova DNA Damage Control - Everyday SPF 44 - Drug Facts

Active Ingredients

Octinoxate 6.5%, Octisalate 2.5% Zinc Oxide 8.5%

Purpose

Sunscreen

Uses

- Helps prevent sunburn
- If used as directed and 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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply liberally 15 minutes before sun exposure.
- Use a water resistant sunscreen if swimming or sweating.
- Reapply at least every 2 hours.
- Children under 6 months of age: ask a doctor.
- **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 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-sleeve shirts, pants, hats, and sunglasses.

Inactive Ingredients

Allantoin, Ascorbyl Palmitate, Butylene Glycol, Cetearyl Glucoside, Citric Acid, Cyclopentasiloxane, Dimethicone, Ergothioneine, Ethyl Hexyl Isononanoate, Glycereth-26, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Iodopropynyl Butylcarbamate, Isopropyl Palmitate, Lecithin, Micrococcus Lysate, Octyl Sterate, Oleth-3 Phosphate, Panthenol, PEG-7 Trimethylolpropane Coconut Ether, Phenoxyethanol, Plankton Extract, Polyether-1, Polyisobutene, Purified Water, Retinyl Palmitate, Sodium Hyaluronate, Sodium Hydroxide, Tocopherol Acetate, Triethoxycaprylsilane.

Other Information

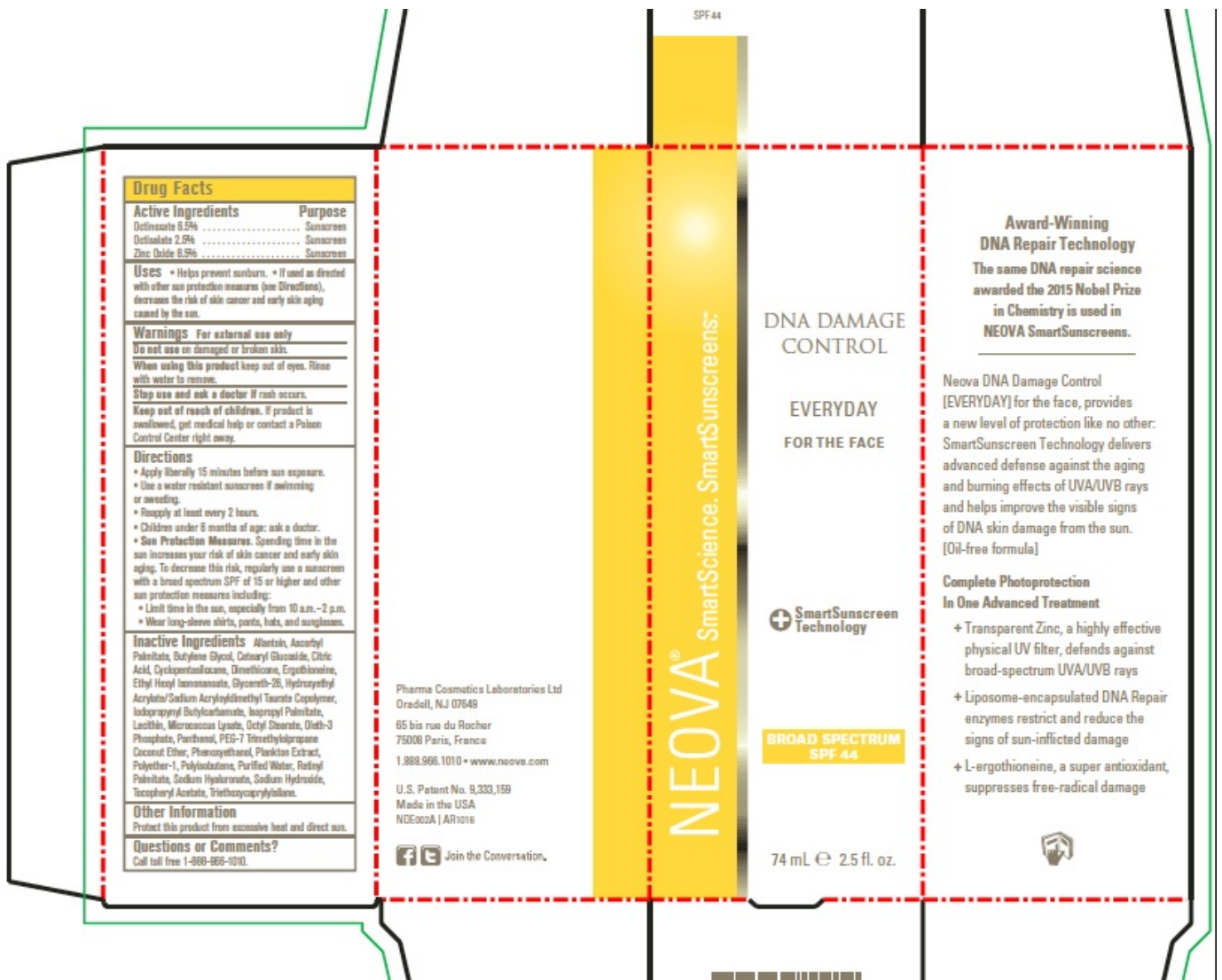
Protect this product from excessive heat and direct sun.

Questions or Comments?

Call toll free 1-888-966-1010.

Product Label

Neova DNA Damage Control
Everyday SPF 44
2.5 fl. oz. (74mL) label and box



NEOVA DNA DAMAGE CONTROL - EVERYDAY BROAD SPECTRUM SPF 44

octinoxate, octisalate, zinc oxide emulsion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72251-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	6.5 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	2.5 g in 100 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	8.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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ALLANTOIN (UNII: 344S277G0Z)
ASCORBYL PALMITATE (UNII: QN83US2B0N)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)
DIMETHICONE (UNII: 92RU3N3Y1O)
ERGOTHIONEINE (UNII: BDZ3DQM98W)
ETHYLHEXYL ISONONANOATE (UNII: I6KB4GE3K4)
GLYCERETH-26 (UNII: NNE56F2N14)
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)
EGG PHOSPHOLIPIDS (UNII: 1Z74184RGV)
OCTYL STEARATE (UNII: 772Y4UFC8B)
OLETH-3 PHOSPHATE (UNII: 8Q0Z18J1VL)
PANTHENOL (UNII: WW9CM0O67Z)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)
DODOXYNOL-5 (UNII: CSH59YN3D0)
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)
POLYISOBUTYLENE (1300 MW) (UNII: 241BN7J12Y)
WATER (UNII: 059QF0K00R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72251-002-74	74 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package	04/19/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/08/2012	

Labeler - PHARMA COSMETICS, INC (080622701)

Establishment

Name	Address	ID/FEI	Business Operations
Swiss-American CDMO, LLC		080170933	manufacture(72251-002)

