NEOVA DNA DAMAGE CONTROL - EVERYDAY SPF 43- zinc oxide, octinoxate, octisalate emulsion
PhotoMedex, 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.

Neova DNA Damage Control - Everyday SPF 43 - Drug Facts

Active Ingredients

Zinc Oxide 7.5%, Octinoxate 7.5%, Octisalate 2.5%

Sunscreen

• Helps prevent sunburn. • Higher SPF gives more sunburn protection. • Retains SPF after 80 minutes of activity in the water or perspiring.

Warnings

- For external use only.
- When using this product keep out of eyes. Rinse with water to remove.
- Stop use and ask a doctor if rash or irritation develops and lasts.
- Keep out of reach of children. If swallowed, contact a Poison Control Center immediately or get medical help right away.

When using this product

Keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if

Rash or irritation develops and lasts.

Keep out of reach of children

If swallowed, contact a Poison Control Center immediately or get medical help right away.

Directions

• Apply every morning to face, neck and décolletage. • Reapply as needed or after towel drying, swimming or perspiring.
• Children under 6 months of age: ask a doctor.

Other Information

High protection sun product.
Sun Alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

Serious side effects associated with use of this product may be reported to this number: 888-966-1010.

Inactive Ingredients

Water (Aqua), Isopropyl Palmitate, Octyl Stearate, Ethyl Hexyl Isononanoate, Cyclopentasiloxane, Cetearyl Glucoside, Micrococcus Lysate, Plankton Extract, L-ergothioneine, Dimethicone, Glycereth-26, Sodium Hyaluronate, Panthenol, Allantoin, Tocopheryl Acetate, Ascorbyl Palmitate (Vitamin C), Oleth-3 Phosphate, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Polyisobutene, PEG-7 Trimethylolpropane Coconut Ether, Lecithin, Polyether-1, Phenoxyethanol, Butylene Glycol, Citric Acid, Iodopropynyl Butylcarbamate, Triethoxycaprylylsilane.

Image of box and label

DNADamageControlEday.jpg

EVERYDAY
SPF 43

NEOVA®

DNA DAMAGE CONTROL™

EVERYDAY | SPF 43

FOR THE FACE

DNA REPAIR ⊕
TRANSPARENT ZINC

PROTECT+

74 mL ℮ 2.5 fl. oz.

Defends Against and Repairs the signs of UV-Inflicted Damage.

- DNA Repair enzymes inhibit and correct the consequences of photodamage
- High-performance UVA/UVB protection
- Hyper antioxidant defense + oil-free hydration

Directions: Apply every morning to face, neck , décolletage and hands. Reapply as needed. For optimal results, use daily with Neova [DNA + Copper] products.

WARNINGS: For external use only. Avoid contact with eyes. Discontinue use if signs of irritation or rash appear. If irritation persists, consult a doctor. Keep out of reach of children.

Active Ingredients: Zinc Oxide 7.5%, Octinoxate 7.5%, Octisalate 2.5%.

Key Performance Ingredients: DNA Repair Enzymes: Photolysomes, Endosomes; Antioxidant: L-ergothioneine.

PhotoMedex, Inc., Montgomeryville, PA 18936
Stamford New Road, Altrincham, Cheshire WA14 1EP UK
1.888.966.1010 • www.neova.com
Patent Pending
Reorder# NDE002 Made in the USA REV 6/11

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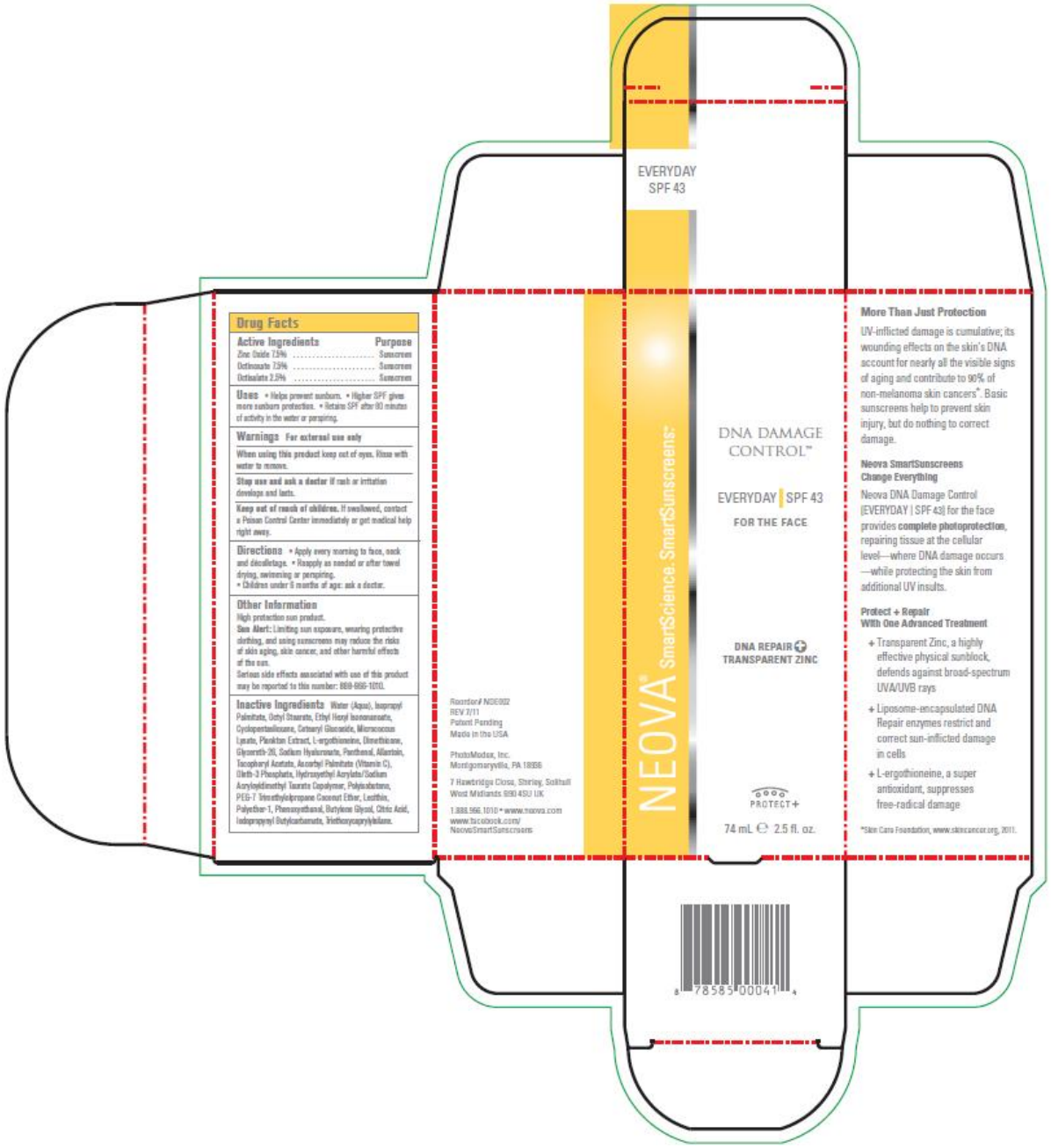

NEOVA DNA DAMAGE CONTROL - EVERYDAY SPF 43

zinc oxide, octinoxate, octisalate emulsion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62362-139
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	7.5 mL in 100 mL
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	7.5 mL in 100 mL

Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	2.5 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Isopropyl Palmitate (UNII: 8CRQ2TH63M)	
Octyl Stearate (UNII: 772Y4UFC8B)	
Ethylhexyl Isononanoate (UNII: I6KB4GE3K4)	
Cyclomethicone 5 (UNII: 0THT5PCI0R)	
Cetearyl Glucoside (UNII: 09FUA47KNA)	
Ergothioneine (UNII: BDZ3DQM98W)	
Dimethicone (UNII: 92RU3N3Y1O)	
Glycereth-26 (UNII: NNE56F2N14)	
Hyaluronate Sodium (UNII: YSE9PPT4TH)	
Panthenol (UNII: WV9CM0O67Z)	
Allantoin (UNII: 344S277G0Z)	
Alpha-Tocopherol Acetate (UNII: 9E8X80D2L0)	
Ascorbyl Palmitate (UNII: QN83US2B0N)	
Oleth-3 Phosphate (UNII: 8Q0Z18J1VL)	
Polyisobutylene (1300 MW) (UNII: 241BN7J12Y)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Butylene Glycol (UNII: 3XUS85K0RA)	
Citric Acid (UNII: 2968PHW8QP)	
Iodopropynyl Butylcarbamate (UNII: 603P14DHEB)	
Triethoxycaprylylsilane (UNII: LDC331P08E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62362-139-01	1 in 1 BOX		
1	NDC:62362-139-74	74 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/06/2012	

Labeler - PhotoMedex, Inc. (054503875)

Establishment

Name	Address	ID/FEI	Business Operations
PhotoMedex, Inc.		054503875	manufacture

Revised: 1/2012 PhotoMedex, Inc.