

# Photoprotection

## Part I. Photoprotection by naturally occurring, physical, and systemic agents

Rebecca Jansen, MD,<sup>a</sup> Steven Q. Wang, MD,<sup>b</sup> Mark Burnett, MD,<sup>b</sup> Uli Osterwalder, MS,<sup>c</sup> and Henry W. Lim, MD<sup>a</sup>  
*Detroit, Michigan; New York, New York; and Monheim, Germany*

### CME INSTRUCTIONS

The following is a journal-based CME activity presented by the American Academy of Dermatology and is made up of four phases:

1. Reading of the CME Information (delineated below)
2. Reading of the Source Article
3. Achievement of a 70% or higher on the online Case-based Post Test
4. Completion of the Journal CME Evaluation

#### CME INFORMATION AND DISCLOSURES

##### Statement of Need:

The American Academy of Dermatology bases its CME activities on the Academy's core curriculum, identified professional practice gaps, the educational needs which underlie these gaps, and emerging clinical research findings. Learners should reflect upon clinical and scientific information presented in the article and determine the need for further study.

##### Target Audience:

Dermatologists and others involved in the delivery of dermatologic care.

##### Accreditation

The American Academy of Dermatology is accredited by the Accreditation Council for Continuing Medical Education to provide continuing medical education for physicians.

##### AMA PRA Credit Designation

The American Academy of Dermatology designates this journal-based CME activity for a maximum of 1 *AMA PRA Category 1 Credits*<sup>™</sup>. Physicians should claim only the credit commensurate with the extent of their participation in the activity.

##### AAD Recognized Credit

This journal-based CME activity is recognized by the American Academy of Dermatology for 1 AAD Credit and may be used toward the American Academy of Dermatology's Continuing Medical Education Award.

##### Disclaimer:

The American Academy of Dermatology is not responsible for statements made by the author(s). Statements or opinions expressed in this activity reflect the views of the author(s) and do not reflect the official policy of the American Academy of Dermatology. The information provided in this CME activity is for continuing education purposes only and is not meant to substitute for the independent medical judgment of a healthcare provider relative to the diagnostic, management and treatment options of a specific patient's medical condition.

##### Disclosures

##### Editors

The editors involved with this CME activity and all content validation/peer reviewers of this journal-based CME activity have reported no relevant financial relationships with commercial interest(s).

##### Authors

Dr Lim has served as consultant for Ferndale, La Roche-Posay, Pierre Fabre, Uriage, and Palatin. He has received research grants from Clinuvel and Estee Lauder. Mr Osterwalder is a full time employee of BASF. Dr Wang has served on the advisory board of L'Oreal. Drs Burnett and Jansen have no conflicts of interest to declare.

##### Planners

The planners involved with this journal-based CME activity have reported no relevant financial relationships with commercial interest(s). The editorial and education staff involved with this journal-based CME activity have reported no relevant financial relationships with commercial interest(s).

##### Resolution of Conflicts of Interest

In accordance with the ACCME Standards for Commercial Support of CME, the American Academy of Dermatology has implemented mechanisms, prior to the planning and implementation of this Journal-based CME activity, to identify and mitigate conflicts of interest for all individuals in a position to control the content of this Journal-based CME activity.

##### Learning Objectives

After completing this learning activity, participants should be able to provide an overview of natural, physical, and systemic photoprotective agents.

**Date of release:** December 2013

**Expiration date:** December 2016

© 2013 by the American Academy of Dermatology, Inc.

<http://dx.doi.org/10.1016/j.jaad.2013.08.021>

##### Technical requirements:

##### American Academy of Dermatology:

- Supported browsers: FireFox (3 and higher), Google Chrome (5 and higher), Internet Explorer (7 and higher), Safari (5 and higher), Opera (10 and higher).
- JavaScript needs to be enabled.

##### Elsevier:

##### Technical Requirements

This website can be viewed on a PC or Mac. We recommend a minimum of:

- PC: Windows NT, Windows 2000, Windows ME, or Windows XP
- Mac: OS X
- 128MB RAM
- Processor speed of 500MHz or higher
- 800x600 color monitor
- Video or graphics card
- Sound card and speakers

##### Provider Contact Information:

American Academy of Dermatology

Phone: Toll-free: (866) 503-SKIN (7546); International: (847) 240-1280

Fax: (847) 240-1859

Mail: P.O. Box 4014; Schaumburg, IL 60168

##### Confidentiality Statement:

##### American Academy of Dermatology: POLICY ON PRIVACY AND CONFIDENTIALITY

**Privacy Policy** - The American Academy of Dermatology (the Academy) is committed to maintaining the privacy of the personal information of visitors to its sites. Our policies are designed to disclose the information collected and how it will be used. This policy applies solely to the information provided while visiting this website. The terms of the privacy policy do not govern personal information furnished through any means other than this website (such as by telephone or mail).

**E-mail Addresses and Other Personal Information** - Personal information such as postal and e-mail address may be used internally for maintaining member records, marketing purposes, and alerting customers or members of additional services available. Phone numbers may also be used by the Academy when questions about products or services ordered arise. The Academy will not reveal any information about an individual user to third parties except to comply with applicable laws or valid legal processes.

**Cookies** - A cookie is a small file stored on the site user's computer or Web server and is used to aid Web navigation. Session cookies are temporary files created when a user signs in on the website or uses the personalized features (such as keeping track of items in the shopping cart). Session cookies are removed when a user logs off or when the browser is closed. Persistent cookies are permanent files and must be deleted manually. Tracking or other information collected from persistent cookies or any session cookie is used strictly for the user's efficient navigation of the site.

**Links** - This site may contain links to other sites. The Academy is not responsible for the privacy practices or the content of such websites.

**Children** - This website is not designed or intended to attract children under the age of 13. The Academy does not collect personal information from anyone it knows is under the age of 13.

**Elsevier:** [http://www.elsevier.com/wps/find/privacypolicy.cws\\_home/privacypolicy](http://www.elsevier.com/wps/find/privacypolicy.cws_home/privacypolicy)

The acute and chronic consequences of ultraviolet radiation on human skin are reviewed. An awareness of variations in naturally occurring photoprotective agents and the use of glass, sunglasses, and fabric can lead to effective protection from the deleterious effects of ultraviolet radiation. New systemic agents, including *Polypodium leucotomos*, afamelanotide, and antioxidants have potential as photoprotective agents. (J Am Acad Dermatol 2013;69:853.e1-12.)

**Key words:** afamelanotide; antioxidants; glass; photoprotection; photoprotective agents; physical systemic photoprotective agents; ultraviolet radiation.

Effective protection from the harmful effects of ultraviolet radiation (UVR) requires the regular practice of photoprotective strategies, which include seeking shade, the use of clothing, wide-brimmed hats, sunglasses, and the application of sunscreens. Geographic and environmental variations affecting UVR transmission, protective cutaneous chromophores, glass, sunglasses, and fabric will be discussed in this article, along with the multiple systemic agents that show potential as photoprotective agents. Part II of our review will focus on sunscreen development, efficacy, and controversies.

## ULTRAVIOLET RADIATION

### Key points

- **Acute effects of UVR include erythema, pigment darkening, delayed tanning, epidermal hyperplasia, free radical formation, and vitamin D synthesis.**
- **Chronic effects of UVR include photoaging, immunosuppression, photocarcinogenesis, and exacerbation of photodermatoses.**

### Acute effects of ultraviolet radiation exposure on human skin

The sun emits UVR with wavelengths shorter than visible light (VL; 400-760 nm) but longer than x-rays. As agreed in the Second International Congress on

### CAPSULE SUMMARY

- Ultraviolet radiation exposure leads to harmful acute and chronic effects on human skin.
- Variations in naturally occurring photoprotective agents affect the degree of ultraviolet radiation exposure.
- Physical photoprotective agents, especially glass, are often underused in photoprotection strategies.
- Limitations of topical photoprotection have driven the search for systemic alternatives.

Light in August 1932, UVR is divided into ultraviolet C (UVC; 270-290 nm), ultraviolet B (UVB; 290-315 nm), and ultraviolet A (UVA; 315-400 nm) wavelengths.<sup>1</sup> More recently, because of the recognition that the biologic effects of shorter spectrum UVA are close to that of UVB, UVA is further subdivided into UVA2 (315-340 nm) and UVA1 (340-400 nm). Because ozone in the stratosphere filters UVC radiation, the cutaneous effects of UVR exposure are attrib-

uted to UVA and UVB.

The initial response of human skin exposure to UVR includes erythema, immediate pigment darkening (IPD), persistent pigment darkening (PPD), delayed tanning, epidermal hyperplasia, free radical formation, and vitamin D synthesis.<sup>2,3</sup>

**Erythema.** Sunburn, primarily caused by UVB exposure (and to a lesser extent UVA2 exposure), is the best recognized acute effect of ultraviolet (UV) light exposure on human skin. UVB-induced erythema reaches its peak between 6 and 24 hours after exposure. An immediate erythema reaction, which lasts for 48 to 72 hours after exposure, characterizes skin reaction to high-dose UVA.

**Immediate and persistent pigment darkening.** Exposure to UVR also results in pigmentary alteration. IPD, predominantly caused by UVA light, is often gray in color. It develops within minutes of irradiation and fades within hours. At UVA

From the Department of Dermatology,<sup>a</sup> Henry Ford Hospital, Detroit; Division of Dermatology,<sup>b</sup> Memorial Sloan Kettering Cancer Center, New York; and BASF Personal Care and Nutrition GmbH,<sup>c</sup> Monheim.

Funding sources: None.

Dr Lim has served as consultant for Ferndale, La Roche-Posay, Pierre Fabre, Uriage, and Palatin. He has received research grants from Clinuvel and Estee Lauder. Mr Osterwalder is a full

time employee of BASF. Dr Wang has served on the advisory board of L'Oreal. Drs Burnett and Jansen have no conflicts of interest to declare.

Reprint requests: Henry W. Lim, MD, Department of Dermatology, Henry Ford Medical Center – New Center One, 3031 W Grand Blvd, Ste 800, Detroit, MI 48202. E-mail: [hlim1@hfhs.org](mailto:hlim1@hfhs.org). 0190-9622/\$36.00

Download English Version:

<https://daneshyari.com/en/article/3205823>

Download Persian Version:

<https://daneshyari.com/article/3205823>

[Daneshyari.com](https://daneshyari.com)