

Sunless Tanning: A Review

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Despite rising numbers of melanoma and nonmelanoma skin cancers, many Americans continue to tan their skin and believe that a tan enhances their appearance. It is well documented that ultraviolet (UV) radiation from the sun or indoor tanning beds is linked to skin cancer and accelerated aging of the skin. In an effort to reduce exposure to UV radiation, and subsequently decrease the risk of skin cancer, some dermatologists have advocated the use of sunless tanning products.

Most sunless tanning products contain the active ingredient dihydroxyacetone (DHA). DHA tans the skin by binding to amino acids in the stratum corneum, producing covalently bound chromophores called melanoids through a process known as the Maillard browning reaction. To decrease the adverse effects associated with UV-radiation-induced tanning, physicians must continue to advocate a safe alternative. Sunless tanning with DHA-based formulations should be recommended to patients desiring a tanned appearance.

Despite rising numbers of melanoma and nonmelanoma skin cancers, many Americans continue to tan their skin believing that they are enhancing their appearance. Throughout time, skin color has played a crucial role in individual identity, self-esteem, and character. Although tanned skin is currently considered by many to be attractive, this has not always been the case. Many ancient cultures, such as the Romans and Greeks, valued pale skin. During the 1800s, high society looked down upon people with tanned skin because a tan implied that they were outdoor laborers and of a lower class. To convey an image of upper-class status, wealthy women carried parasols and spent much of their time inside to prevent darkening of the skin.

The Industrial Revolution created more jobs in sun-protected environments. Instead of connoting lower-class status, tanned skin became a sign of financial strength, an indication that one had enough money to vacation in the sun. The popularity of a fashionable tan was epitomized by the French fashion designer Coco

Chanel in 1923, when she began using tanned women in her advertisements and consequently initiated a trend that still persists today.^{1,2}

As sun exposure increased during the 1920s, physicians began to voice concerns about possible risks. The US Public Health Service issued warnings about the potential dangers of excessive sunbathing. In the 1930s, ultraviolet (UV) radiation was recognized as a carcinogen; a 1935 article in the *Journal of the American Medical Association* listed sunlight as a possible cause of cancer. Further studies confirmed this hypothesis, implicating sunlight as a cause of cutaneous malignancy and accelerated photoaging.³ However, despite the recent increase in public awareness of this association, sunbathing outdoors and indoor tanning under UV lights continue at a high rate.⁴

It is well documented that UV radiation from the sun and indoor tanning beds is linked to skin cancer and accelerated aging of the skin.⁵⁻¹⁴ In an effort to decrease the practice of UV-radiation-induced tanning and subsequently decrease the risk of skin cancer, some dermatologists have advocated the use of sunless tanning products.^{4-8,14,15} Most sunless tanning products contain the active ingredient dihydroxyacetone (DHA), a compound observed serendipitously in the 1950s to pigment the skin on contact. At that time, DHA was being administered orally in studies on glycogen storage disease.¹⁶⁻¹⁸

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The authors report no actual or potential conflicts of interest in relation to this article.

BASIC SCIENCE OF SUNLESS TANNING

DHA is a 3-carbon sugar involved in carbohydrate metabolism in higher plants and animals through processes such as glycolysis and photosynthesis.^{17,19-21} DHA tans the skin by binding to amino acids in the stratum corneum, producing covalently bound chromophores called melanoins through a process known as the Maillard browning reaction.^{1,8,18,20,22-24} This tanning effect is independent of light but is enhanced by UV radiation.²⁵

To produce a sunless tanning product, DHA is usually added to a lotion, cream, liquid, or mousse base in concentrations ranging from 3% to 5%.^{1,4,8} The intensity of the color is dependent on the concentration, thus allowing for a wide range of shades to accommodate individual preference.⁸ Increasing the protein content of the stratum corneum may enhance coloring; thus, the shade achieved is dependent on the thickness of keratin as well.^{1,20,26} Rougher, hyperkeratotic skin, as well as freckled and mottled skin, takes up the color at an increased intensity, resulting in coloring irregularities.⁸ The brown color produced from DHA is resistant to soap and water but may be removed by sloughing of the stratum corneum, as with vigorous scrubbing.^{19,20,22,24} While the pigment from DHA takes several hours to form, many products now contain water-soluble dyes that impart immediate color upon application.

CLINICAL APPLICATIONS

Dermatologists encourage sunless tanning as an alternative to UV radiation, and fortunately, sunless tanning products have advanced in cosmetic appeal over the years. More recent products create a natural golden-brown hue as opposed to the orange color that often resulted from the use of older products. In addition to DHA-based topical tanning products, sunless tanning booths using a spray-on application have increased the ease and accessibility of UV-radiation-free tanning.^{4,8} In a recent study by Sheehan and Leshner,⁷ the majority of individuals who have used UV-radiation indoor tanning beds in the past reported doing so less frequently as a result of sunless tanning. Another recent study associated the use of sunless tanning products with increased sun protection behavior.²⁷

Other studies have explored the possibility of continued indoor tanning despite known health risks as a psychosocial phenomenon or an addictive behavior learned in adolescence.^{5,6,14,28} Demko et al¹⁴ reported that 36.8% of white female adolescents and 11.2% of white male adolescents in their study population had used indoor tanning beds at least once. Feldman et al⁵ showed that UV exposure by indoor tanning beds in adolescents is a reinforcing stimulus. In addition, Zeller et al⁶ reported that teenagers who begin tanning at a young age and do

so frequently are more likely to have difficulty quitting. These findings are consistent with other addictive behaviors that can arise during adolescence.

Although DHA has been popularized as the browning agent in topical sunless tanning formulations, it is widely known for other uses as well. Initially, before the browning properties of DHA were discovered, this compound was used to treat diabetic coma, was administered to patients with diabetes as a glucose alternative, and was used as a tool to test for glycogen storage diseases.¹⁷ More recently, DHA has been investigated for its potential use in skin camouflaging for patients with vitiligo. Studies by Fesq et al²⁶ and Suga et al²⁹ showed it to be both a practical and well-accepted cosmetic treatment for this form of skin depigmentation. DHA has also been used to enhance photochemotherapy in psoriasis treatment.²⁴

DHA-based formulations have been found to protect the skin of patients with ultraviolet-A (UVA) sensitivity such as hereditary polymorphic light eruption, photosensitive porphyria, drug photoallergy, and actinic reticuloid.^{20,30-34} Many traditional sunscreens do not protect from UV radiation in the longer UVA and visible light range.^{32,35,36} DHA has a proven benefit in blocking longer UVA (320–340 nm) and visible (400–750 nm) light.^{31,34} This photoprotection has been demonstrated in the laboratory as well. In 1975, Fusaro and Johnson³³ demonstrated that a combination of DHA and lawsone prevented photo-induced edema in the paws of photosensitized rats.

SAFETY

With the increasing use of DHA, physicians must consider not only its efficacy but safety issues as well. This 3-carbon sugar is a physiologic product of the body and is presumed to be nontoxic.^{16,23} DHA has been used as diet supplementation in rats to investigate postprandial glycogen metabolism; no adverse effects were noted.³⁷ Two case reports published by Morren et al³⁸ documented contact allergy to DHA. However, in a reply to these case reports, Johnson and Fusaro²² pointed out that the source and purity of DHA were not specified. Concerns have arisen about the possible mutagenic properties of DHA, and there are conflicting reports in the literature. Pham et al³⁹ demonstrated in 1980 that DHA is mutagenic in the *Salmonella* mutagenicity assay. More recently, Petersen et al²³ discovered DHA to induce DNA damage, cell-cycle block, and apoptosis in cultured keratinocytes. However, in another study, Petersen et al²⁰ reported DHA to delay UV-radiation-induced photocarcinogenesis in hairless mice. Other investigators have shown DHA to have a lack of mutagenicity or even to be antimutagenic. Akin and Marlowe¹⁷ observed no increase in cancer in mice after application of topical DHA for 80 weeks. Furthermore,

Chan et al⁴⁰ reported that the products of the Maillard browning reaction are antimutagenic. Taken as a whole, the medical literature seems to support the safety of topical application of DHA. Safety concerns still remain with regard to the use of DHA during pregnancy and the possible theoretical risks involved in inhaling DHA or other components of the tanning solutions in spray-on booths.

CONCLUSION

Civilization has a long history of preoccupation with skin color. The golden-bronze look is seemingly attractive and one that many aspire to achieve. Therefore, the public will continue to engage in UV-radiation-induced tanning despite known health risks. To decrease the adverse effects associated with this practice, physicians must continue to advocate a safe alternative. Sunless tanning with DHA-based formulations should be recommended to patients desiring a tanned appearance. Patients should also be informed that use of sunless tanning products is not sufficient sun protection. Formulations with DHA result in a sun protection factor (SPF) of only 2 and offer no protection against ultraviolet-B light.^{8,34} In addition, the SPF gained from the sunless tanning product is effective only for a short period of time, not for the duration of the artificial tan. It is recommended that patients use a traditional sunscreen with an SPF of at least 15 in conjunction with the sunless tanner.¹⁵

With more cosmetically appealing products now available, sunless tanning is becoming a safe alternative to the "traditional" tan. The long-term risks, if any, of consistent use of DHA in humans have not been fully elucidated. However, the compound appears to be safe. It is prudent for dermatologists to continue to recommend this alternative for skin tanning and to inform patients of the need for added SPF through the application of traditional sunscreen. Until society deems pale, nontanned skin attractive, we must continue to pursue alternatives to decrease UV-radiation-induced photocarcinogenesis.

REFERENCES

1. Draeels ZD. Self-tanning lotions: are they a healthy way to achieve a tan? *Am J Clin Dermatol*. 2002;3:317-318.
2. Draeels ZD. Cosmetics to imitate a summer tan. *Skin Therapy Lett*. 2000;6:3-4.
3. Albert MR, Ostheimer KG. The evolution of current medical and popular attitudes toward ultraviolet light exposure: part 3. *J Am Acad Dermatol*. 2003;49:1096-1106.
4. Fu JM, Dusza SW, Halpern AC. Sunless tanning. *J Am Acad Dermatol*. 2004;50:706-713.
5. Feldman SR, Liguori A, Kucenic M, et al. Ultraviolet exposure is a reinforcing stimulus in frequent indoor tanners. *J Am Acad Dermatol*. 2004;51:45-51.
6. Zeller S, Lazovich D, Forster J, et al. Do adolescent indoor tanners exhibit dependency? *J Am Acad Dermatol*. 2006;54:589-596.
7. Sheehan DJ, Leshner JL. The effect of sunless tanning on behavior in the sun: a pilot study. *South Med J*. 2005;98:1192-1195.
8. Levy SB. Dihydroxyacetone-containing sunless or self-tanning lotions. *J Am Acad Dermatol*. 1992;27:989-993.
9. Hornung RL, Magee KH, Lee WJ, et al. Tanning facility use: are we exceeding Food and Drug Administration limits? *J Am Acad Dermatol*. 2003;49:655-661.
10. Rigel EG, Leibold M, Rigel AC, et al. Daily UVB exposure levels in high-school students measured with digital dosimeters. *J Am Acad Dermatol*. 2003;49:1112-1114.
11. Thieden E, Philipsen PA, Heydenreich J, et al. UV radiation exposure related to age, sex, occupation, and sun behavior based on time-stamped personal dosimeter readings. *Arch Dermatol*. 2004;140:197-203.
12. Diffey BL. A quantitative estimate of melanoma mortality from ultraviolet A sunbed use in the U.K. *Br J Dermatol*. 2003;149:578-581.
13. Karagas MR, Stannard VA, Mott LA, et al. Use of tanning devices and risk of basal cell and squamous cell skin cancers. *J Natl Cancer Inst*. 2002;94:224-226.
14. Demko CA, Borawski EA, Debanne SM, et al. Use of indoor tanning facilities by white adolescents in the United States. *Arch Pediatr Adolesc Med*. 2003;157:854-860.
15. Dajani Z, Swetter SM, Demierre MF, et al. Sun protection factor content and warning statements for sunless tanning products: an examination of retail outlets and the Internet. *J Am Acad Dermatol*. 2005;53:919-920.
16. Goldman L, Barkoff J, Blaney D, et al. Investigative studies with the skin coloring agents dihydroxyacetone and glyoxal: preliminary report. *J Invest Dermatol*. 1960;35:161-164.
17. Akin FJ, Marlow E. Non-carcinogenicity of dihydroxyacetone by skin painting. *J Environ Pathol Toxicol Oncol*. 1984;5:349-351.
18. Wittgenstein E, Berry HK. Staining of the skin with dihydroxyacetone. *Science*. 1960;132:894-895.
19. Maibach HI, Klignan AM. Dihydroxyacetone: a suntan-simulating agent. *Arch Dermatol*. 1960;82:505-507.
20. Petersen AB, Na R, Wulf HC. Sunless skin tanning with dihydroxyacetone delays broad-spectrum ultraviolet photocarcinogenesis in hairless mice. *Mutat Res*. 2003;542:129-138.
21. Enders D, Voith M, Lenzen A. The dihydroxyacetone unit—a versatile C(3) building block in organic synthesis. *Angew Chem Int Ed Engl*. 2005;44:1304-1325.
22. Johnson JA, Fusaro RM. Therapeutic potential of dihydroxyacetone. *J Am Acad Dermatol*. 1993;27:989-993.
23. Petersen AB, Wulf HC, Gniadecki R, et al. Dihydroxyacetone, the active tanning ingredient in sunless tanning lotions, induces DNA damage, cell-cycle block and apoptosis in cultured HaCaT keratinocytes. *Mutat Res*. 2004;560:173-186.
24. Taylor CR, Kwangskuth C, Wimberly J, et al. Turbo-PUVA: dihydroxyacetone-enhanced photochemotherapy for psoriasis. *Arch Dermatol*. 1999;135:540-544.
25. Puccetti G, Leblanc RM. A sunscreen-tanning compromise: 3D visualization of the actions of titanium dioxide particles and dihydroxyacetone on human epiderm. *Photochem Photobiol*. 2000;71:426-430.
26. Fesq H, Brockow K, Strom K, et al. Dihydroxyacetone in a new formulation—a powerful therapeutic option in vitiligo. *Dermatology*. 2001;203:241-243.
27. Mahler HI, Kulik JA, Harrell J, et al. Effects of UV photographs, photoaging information, and use of sunless tanning lotion on sun protection behaviors. *Arch Dermatol*. 2005;141:373-380.
28. Kaur M, Liguori A, Lang W, et al. Induction of withdrawal-like symptoms in a small randomized, controlled trial of opioid blockade in frequent tanners. *J Am Acad Dermatol*. 2006;54:709-711.
29. Suga Y, Ikejima A, Matsuba S, et al. Medical pearl: DHA application for camouflaging segmental vitiligo and piebald lesions. *J Am Acad Dermatol*. 2002;47:436-438.
30. Fusaro RM, Johnson JA. Topical photoprotection for hereditary polymorphic light eruption of American Indians. *J Am Acad Dermatol*. 1991;24:744-746.

A REVIEW OF SUNLESS TANNING

31. Asawanonda P, Oberlender S, Taylor C. The use of dihydroxyacetone for photoprotection in variegated porphyria. *Int J Dermatol*. 1999;38:916-925.
32. Rice EG. Dihydroxyacetone naphthoquinone protection against photosensitivity. *Dermatologica*. 1976;153:38-43.
33. Fusaro RM, Johnson JA. Protection against long ultraviolet and/or visible light with topical dihydroxyacetone. implications for the mechanism of action of the sunscreen combination, dihydroxyacetone/naphthoquinone. *Dermatologica*. 1975;150:346-351.
34. Kullavanijaya P, Lim HW. Photoprotection. *J Am Acad Dermatol*. 2005;52:937-958.
35. Kaye ET, Levin JA, Blank IH, et al. Efficiency of opaque photoprotective agents in the visible light range. *Arch Dermatol*. 1991;127:351-355.
36. Johnson JA, Fusaro RM. Protection against long ultraviolet radiation: topical browning agents and a new outlook. *Dermatologica*. 1987;175:53-57.
37. Obeid OA, Bittar ST, Hwalla N, et al. Effect of diet supplementation with glutamine, dihydroxyacetone, and leucine on food intake, weight gain, and postprandial glycogen metabolism of rats. *Nutrition*. 2005;21:224-229.
38. Morren M, Dooms-Goossens A, Heidebuchel M, et al. Contact allergy to dihydroxyacetone. *Contact Dermatitis*. 1991;25:326-327.
39. Pham HN, DeMarini DM, Brockmann HE. Mutagenicity of skin tanning lotions. *J Environ Pathol Toxicol*. 1979;3:227-231.
40. Chan RI, Stich HF, Rosin MP, et al. Antimutagenic activity of browning reaction products. *Cancer Lett*. 1982;15:27-33. ■