Review

The Role of Phytonutrients in Skin Health

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Abstract: Photodamage is known to occur in skin with exposure to sunlight, specifically ultraviolet (UV) radiation. Such damage includes inflammation, oxidative stress, breakdown of the extracellular matrix, and development of cancer in the skin. Sun exposure is considered to be one of the most important risk factors for both nonmelanoma and melanoma skin cancers. Many phytonutrients have shown promise as photoprotectants in clinical, animal and cell culture studies. In part, the actions of these phytonutrients are thought to be through their actions as antioxidants. In regard to skin health, phytonutrients of interest include vitamin E, certain flavonoids, and the carotenoids, β-carotene, lycopene and lutein.

Keywords: photodamage; vitamin E; flavonoids; carotenoids

1. Introduction

Skin is the largest organ in the human body. It is exposed to environmental insult of which UV light is thought to be the most harmful [1]. UV exposure can cause oxidative stress, inflammation, erythema, breakdown of the extracellular matrix, wrinkling and skin cancer [2-4]. In fact, cumulative sun exposure is one of the most important risk factors for both nonmelanoma and melanoma skin cancers [5,6].

Over 1 million skin cancers in the US are attributed to UV exposure per year, making it a significant public health concern [7]. The Healthy People 2010 goals include reducing sun exposure as part of the skin cancer reduction goals [8]. Skin cancer can be categorized as nonmelanoma and
Nonmelanoma skin cancer has an estimated incidence of over 600,000 cases per year in the US. Approximately 500,000 are BCCs and 100,000–150,000 are SCCs [9]. Melanoma has an estimated incidence of between 11 and 14 per 100,000 cases per year in the US among white populations [10].

There are many ways to protect oneself from UV exposure including avoidance of the sun, wearing protective clothing (hats, long-sleeved shirts, treated fabrics, etc.), and topical sunscreen. However, the potential for dietary UV protection is a relatively recent concept. Compared to topical sunscreens which require reapplication and have localized effects, dietary methods are thought to provide continual whole body protection. This promise has led to an explosion in nutrient containing products which are marketed for skin health improvement. Common ingredients of these products, both cosmetic and consumable, are antioxidant nutrients. While a variety of nutrients have been implicated in skin health [11,12], the purpose this review is to evaluate the evidence to date on the role of certain phytonutrients in skin health. Given the implication of oxidative stress being causative in photodamage, antioxidant nutrients are of interest in the prevention of such damage, the nutrients of interest in the study of photoprotection in skin include α-tocopherol, and certain flavonoids and β-carotene, lycopene, and lutein.

2. UV Damage

UV light is grouped as UVA (315–400 nm), UVB (280–315 nm) and UVC (220–280 nm), from lowest to highest energy. The Earth’s atmosphere filters out UVC light and of the remaining UV light to which we are exposed, 95% is UVA. The primary focus of UV research has been UVB light, but this is shifting toward UVA light. UVA is of higher wavelength which enables it to penetrate deeper into the skin [13]. UVB is mostly absorbed by the epidermal layer, but UVA affects the dermal layer as well. UVA has demonstrated carcinogenicity in animal models [14], emphasizing the need to understand and prevent the processes by which UVA exerts its biological effects. Clinical changes seen with photoaging, such as roughness, fine wrinkles, spotty hyperpigmentation, vasodilation, and loss of elasticity, are attributed primarily to UVA as well [15].

UVA is thought to damage the skin primarily via generating reactive chemical species (ROS) upon absorption of photons by riboflavin, porphyrins and heme-containing proteins [14]. ROS, reactive nitrogen species (RNS) as well as further enzymatic production of ROS have been implicated. Reactive species can damage major biomolecules including DNA, proteins and lipids in various skin strata leading to cytotoxicity, mutations and alterations in cell signaling pathways. In cultured human fibroblasts from foreskin biopsies, UVA induced activity in p38 and c-Jun-N-terminal kinase (JNK) of the mitogen-activated protein kinase (MAPK) family, but not extracellular signal-regulated kinases (ERK)[16]. This activation was attributed to singlet oxygen (¹O₂) because of subsequent inhibition by ¹O₂ quenchers. This activation can lead to downstream activation of transcription factors that activate stress-inducible genes, specifically matrix metalloproteinase-1 (MMP-1). MMPs are responsible for the degradation of the extracellular matrix in skin.
3. α-Tocopherol

Vitamin E refers to a family of eight nutrients of which α-tocopherol is the most abundant and biologically active form in the human body (Figure 1). The α form has 3 methyl groups on its chromanol ring and is recognized by the tocopherol transfer protein (TTP) which is responsible for its intracellular transport. This essential lipophilic nutrient is well known for its role as a chain-breaking antioxidant during lipid peroxidation and it protects polyunsaturated fatty acids in cell membranes from oxidation. Vitamin E also has the ability to affect signal transduction and gene expression [17]. Food sources include vegetables, vegetable oils, cereals and nuts [18]. Almonds are a particularly good source of α-tocopherol, containing approximately 26 mg per 100 g [19].

![Figure 1. Chemical structure of α-tocopherol (AT).](image)

α-Tocopherol concentrations in skin can be increased with oral or topical delivery [18] and are decreased with UV exposure [20]. Distribution among the layers of human skin has been determined (Table 1). Within human upper arm stratum corneum, a gradient of highest to lowest α-tocopherol concentration was found from lower to upper layers [21]. α-Tocopherol’s activity as an antioxidant, combined with its presence in human skin, provide biologic plausibility for its role as a photoprotectant in skin.

| Skin Layer            | Concentration          | Reference         |
|-----------------------|------------------------|-------------------|
| Epidermis             | 31 ± 3.8 nmol/g tissue | Shindo et al. [20]|
| Dermis                | 16.2 ± 1.1 nmol/g tissue | Shindo et al. [20]|
| Stratum corneum       | 33 ± 4 nmol/g tissue   | Thiele et al. [19]|
| Sebum                 | 76.5 ± 1.5 nmol/g sebum| Thiele et al. [21]|

3.1. Epidemiologic Studies

In a review of epidemiologic literature, McNaughton et al. found evidence of a weak protective effect of vitamin E in nonmelanoma carcinoma (BCC and SCC)[22]. Of the 12 studies examined, two identified an inverse relationship between intake and BCC risk while one demonstrated a positive relationship. Six found no relationship between plasma or intake of vitamin E and BCC risk. Four studies found no association between vitamin E and SCC. More recently, two studies in an Australian population reported no association between serum α-tocopherol and nonmelanoma carcinoma but a positive association between dietary α-tocopherol, including supplements, and BCC [23,24]. However, in these two studies a single baseline serum measurement or dietary assessment was compared to
cancer incidence during an eight year period. These unexpected findings may reflect the possible inability of a single measure of vitamin E to accurately reflect vitamin E status and a change in eating habit cannot be ruled out over the 8 year period. Furthermore, these studies analyzed the association between cancer, which is a long-term outcome, and short-term markers of vitamin E intake or status such as dietary assessment and serum concentration. Cancer may be closer related to skin concentrations which were not compared in any of these studies.

There are no epidemiological studies assessing the relationship between topically applied α-tocopherol and skin cancer.

3.2. Clinical Trials

Clinical trials suggest that dietary vitamin E may act as a photoprotectant with the aid of other antioxidants and topical vitamin E may provide protection as well, although reports have not been entirely consistent.

One study showed subjects treated with 400 IU/d α-tocopherol for eight weeks had reduced skin malondialdehyde (MDA) after UV exposure but this intervention provided no protection from erythema, the reddening of the skin commonly referred to as sunburn [14]. Wernignaus et al. [25] showed no photoprotective effect after 6 months of daily dietary supplementation of α-tocopherol acetate (400 IU) in 12 subjects of skin types II-IV. Both clinical measure of minimal erythema dose (MED), the minimum dose of UV necessary to induce erythema in a subject, and histologic measure of sunburn cells were compared. No difference was seen in skin α-tocopherol concentrations between treatment and placebo groups. Plasma α-tocopherol concentrations increased by 65% in the treatment group and 18% in the placebo group, so there was no question of bioavailability. Although a very modest increase in MED was observed in the treatment group after 6 months, the sample size of this study may have been too small to observe significant changes. It is possible that α-tocopherol will provide the most benefit in subjects with a limited range in skin types of increased risk for photodamage (very light to light). Additionally, repetitive UV doses lower than necessary to produce erythema are enough to cause local immunosuppression and skin carcinogenesis [15], suggesting that other markers of damage may be important such as DNA damage.

Other dietary studies have shown vitamin E as a photoprotectant when combined with other antioxidants. Fuchs et al. [26] reported no protection when vitamin C (3 g) and α-tocopherol (2 g or 3000 IU) were supplemented individually, but in combination these nutrients provided protection as assessed by increased MED after 50 days of daily supplementation. Eberlein-Konig et al. [27] also showed increased MED in subjects supplemented daily with a combination of vitamin C (2 g) and α-tocopherol (1000 IU) after only 8 days. A similar study showed increased MED in subjects supplemented daily with α-tocopherol (1200 IU) for 7 days, but when combined with vitamin C (2 g) a greater increase in MED was observed [28]. Placzek et al. [29] demonstrated a photoprotective effect after 3 months of daily dietary supplementation with vitamin C (1 g) and α-tocopherol (500 IU). Participants showed increased resistance to UVB-induced sunburn, as assessed by MED, and protection from DNA damage. This study provides evidence of long-term photoprotection of lower doses than had been previously tested.
Clinical trials suggest that vitamin E applied topically also provides photoprotection. Chung et al. demonstrated topical application of 5% vitamin E (24 hours prior to UV exposure) inhibited UV-induced MMP-12 mRNA by 47% [30]. Other groups have reached opposing conclusions on the protection that vitamin E can provide from erythema. One group demonstrated reduced erythema when vitamin E was applied prior to UV exposure but not after [31]. Another group found vitamin E application after UV exposure protective, but not pretreatment [32].

3.3. Animal Studies

UVB irradiated mice fed α-tocopheryl acetate had less cancer incidence in a dose-dependent manner [33]. Dietary α-tocopherol esters are readily cleaved and free α-tocopherol is absorbed into the circulatory system. It should be noted that the doses used in this study were considered toxic given that more treated mice died compared to control mice. A more recent experiment tested an oral combination of vitamin C, vitamin E, pycnogenol (a pine bark extract rich in polyphenols and procyanidins), and evening primrose oil in mice [34]. The nutrient cocktail treatment significantly inhibited UVB-induced wrinkling and many associated events, including expressions of MMPs, MAPK and activation of activator protein (AP)-1 transcriptional factor. While these results support a role for nutrient effects in photoprotection, specific effects of vitamin E could not be determined.

Studies in mice treated topically with α-tocopherol provide strong evidence of photoprotection [18]. Mice treated topically with 25 mg α-tocopherol three times per week for three weeks before and during twelve weeks of UVB irradiation had significantly less incidence of skin cancer compared to untreated mice [35]. Four additional murine studies demonstrated photoprotection, assessed as skin tumor incidence, from topical α-tocopherol treatment [36-39]. A follow-up study using this same α-tocopherol treatment reduced induction of DNA damage by UVB [40]. Later studies by this group found acetate and succinate esterified forms of α-tocopherol applied in the same manner did not provide the same chemoprevention [41]. This difference may be due to the limited capacity to cleave acetate or succinate when applied topically. This limitation may compromise the antioxidant activity given that ester forms do not have a free hydroxyl group on the aromatic ring necessary for scavenging free radicals.

An additional 15 animal studies have consistently found topical pretreatment with vitamin E photoprotective using variety of outcome measures including erythema, lipid peroxidation, wrinkling and sunburn cell formation [42-56].

3.4. In Vitro

A human skin model has previously shown absorption of topical α-tocopherol -glucoside, α-tocopherol and α-tocopherol acetate into epidermal and dermal compartments [57]. α-Tocopherol was not included in this study. Pig skin biopsies have been cultured showing pretreatment with topical α-tocopherol reduced UVB-induced cytotoxicity, apoptosis and lipid peroxidation [20]. In cultured mouse skin, media pretreatment of 9 µM α-tocopherol-6-O-phosphate, a water-soluble form of α-tocopherol, reduced UVB-induced apoptosis and lipid peroxidation [58]. Media pretreatment of α-tocopherol in human melanocytes prevented UVA- and UVB-induced glutathione loss and diminished apoptosis [59].
4. Flavonoids

Flavonoids are a family of over 5000 compounds found in plants. Flavonoid structure is based on a flavane nucleus, which contains two benzene rings separated by an oxygen containing pyrane ring. Individual flavonoids compounds have been reported to be radical scavengers, UVA absorbent, cytoprotective, anti-inflammatory anti-apoptotic, and to inhibit DNA damage and to affect cellular signaling pathways [60-62]. Collectively, flavonoids act in vitro as antioxidants and induce quinone reductase activity, a marker of chemopreventive activity [63]. Addition of almonds, a rich source of flavonoids, to human diet has reduced biomarkers of oxidative stress [64]. Predominant dietary flavonoids include isorhamnetin, catechin, kaempferol, epicatechin, quercetin and epigallocatechin-3-gallate (EGCG)(Figure 2)[65].

**Figure 2.** Chemical structure of predominant polyphenols.

(+) Catechin

(−) Epicatechin

Epigallocatechin-3-gallate

Quercetin

Kaempferol

Isorhamnetin

Absorption of flavonoids to human skin has not been tested. However, radioactivity was detected in the skin of mice fed $^3$H-labeled EGCG, a closely related flavanol found in green tea [66]. From a single topical dose of EGCG, maximal concentrations of 1,366 and 411 ng/mL were measured in mouse epidermis and dermis, respectively [67].
4.1. Epidemiologic Studies

Epidemiologic evidence suggests an association between flavonoid-rich foods and skin cancer incidence. A case-control study in an Italian population found a negative correlation between cutaneous melanoma and daily tea drinking, high consumption of vegetables, particularly carrots, cruciferous and leafy vegetables, and fruits, especially citrus [68]. In the “Food Habits in Later Life” study, tea consumption was negatively correlated with actinic skin damage in Anglo-Celtic Australians [69].

4.2. Clinical Trials

The acute and long-term photoprotective effects of consumption of a flavanol-rich cocoa beverage was studied in double-blind, placebo controlled clinical trials. Epicatechin and its stereoisomer catechin are the major flavanols found in this beverage, contributing 61 and 20 mg/serving, respectively to a total of 329 mg cocoa flavanols. Two hours after a single dose of the flavanol-rich cocoa, dermal blood flow and oxygen saturation of hemoglobin significantly increased by 1.7- and 1.8-fold, respectively [70]. Long-term, daily ingestion of the same flavanol-rich cocoa beverage decreased UV-induced erythema by 15% after only six weeks [71]. After twelve weeks, additional protection was evident as erythema was decreased by 25%. Blood flow of cutaneous and subcutaneous tissues, skin density and skin hydration were also increased while skin roughness and scaling were significantly decreased. None of these skin improvements was observed in the control group.

Green tea contains the flavanols catechin and epicatechin. An eight week clinical trial evaluating the clinical and histological effects of combined topical and oral green tea supplementation (300 mg twice daily) found improvements in elastic tissue content [72]. However no significant differences in clinical grading of appearance were found. All subjects were asked to apply sunscreen daily which may have attenuated differences between groups.

Immunosuppression was inhibited in participants after a single topical application of either green or white tea treatment (2.5 mg/cm²) as evaluated by hypersensitivity to dinitrochlorobenzene [73]. The same topical treatments prevented DNA damage. Another study found a single topical dose of green tea treatment (1–4 mg/2.5 cm²) protective of DNA damage, in a dose-dependent manner [74]. Long-term topical application of green tea treatment significantly reduced UV-induced p53 expression and apoptosis in keratinocytes, but not erythema [75].

Topical application of an extract of Culcitium reflexum H.B.K. leaves which contain isorhamnetin, quercetin and kaempferol significantly decreased erythema when applied after UVB irradiation [76].

4.3. Animal Studies

Investigations of foods rich in flavanoid nutrients demonstrate their efficacy as photoprotectants against UV carcinogenesis in animal models. Grape seed proanthocyanidins are polymers of flavanol monomers such as catechin and epicatechin [74]. In SKH-1 hairless mice, dietary supplementation with these compounds resulted in a dose-dependent reduction in photocarcinogenesis. Green tea as well as black tea consumption also resulted in significantly fewer UV-induced tumors in mice [77]. The administration of green tea polyphenols in drinking water or the topical application of EGCG, a
major tea flavanol, also induced partial regression or inhibition of tumor growth of established skin papillomas in mice [78]. Gensler et al. also investigated EGCG in BALB/cAnNHsd mice. They found induction of skin tumors by UV irradiation was significantly reduced by topical but not by oral treatment of EGCG [79]. Other research has focused on acute photoprotection of EGCG. In Wistar rats, topical EGCG treatment for 30 minutes prior to UVA irradiation significantly decreased sunburn cell occurrence and dermo-epidermal activation compared to untreated rats [80]. The same protection was not observed with topical EGCG application 30 minutes after UVA exposure. UVA-induced oxidative stress was significantly attenuated in Sprague-Dawley rats by intraperitoneal injections of quercetin prior to UV exposure [81]. Markers of oxidative stress included increased plasma malondialdehyde (a product of lipid peroxidation), decreased erythrocyte glutathione peroxidase and reductase activities, and decreased catalase and superoxide dismutase activities. Topical quercetin treatment was found protective of UVB-induced oxidative stress in mouse skin as well [82].

Positive photoprotective results from the above animal studies lead researchers to investigate the mechanism of protection. Dietary supplementation of green tea polyphenols in mice, during 8 weeks of UVB irradiation, inhibited UV-induced expression of MMPs, including MMP-2, -3, -7, and -9 [83]. All of these MMPs have been shown to be involved in the degradation of types-I and -III collagen fragments generated by collagenases and type IV collagen of the basement membrane.

4.4. In Vitro Studies

In human dermal fibroblasts, both quercetin and kaempferol inhibited MMP-1 (collagenase) activity and expression induced by UVA [34]. EGCG inhibited MMP-2 and -9 activity (gelatinases)[84]. Adult human skin fibroblasts and normal human epidermal keratinocytes, cultured separately, had significantly less DNA damage from UVA irradiation when treated with EGCG [85].

Ex vivo cultured human skin experiments demonstrated pretreatment with topical green and white tea (2 mg/cm²) provided protection from UV induced immunosuppression, measured by depletion of Langerhans cells [73]. Another ex vivo human skin experiment found topical cocoa polyphenol extract (containing catechin and epicatechin) improved skin structure after five days of application [86]. Glycosaminoglycans increased as well as collagen I, II and IV expression, indicating improved elasticity and skin tonus, respectively. In UVA exposed artificial skin models, composed of differentiated epidermis on a dermal substitute, pretreatment with topical EGCG decreased expression of MMP-1 and -3 and activity of MMP-2 and -9 [87]. Tissue inhibitor of metalloproteinase-1 (TIMP-1) expression increased simultaneously.

5. Carotenoids

Carotenoids are a family of compounds of over 600 fat-soluble plant pigments [88]. Fruits and vegetables are major sources of carotenoids [89]. Carotenoids are considered to be efficient scavenger of singlet molecular oxygen [90,91]. In fact, β-carotene (a major dietary carotenoid), at physiological concentrations, has been demonstrated to interact with UVA effects in keratinocytes by mechanisms that included singlet molecular oxygen quenching [92]. Other dietary major dietary carotenoids include lycopene, and lutein [89].
Carotenoids concentrations in human skin have been reported to range from 0.2 to 0.6 nmol/g wet weight [93]. Carotenoid skin concentrations have been shown to vary by location on the body (forehead > palm > dorsal > inside arm = back of hand)[94-97]. Further, skin concentrations can be increased with supplementation [71] and are lower in persons with oxidative stress (e.g., smokers)[94]. Similarly, plasma and skin carotenoids decrease with UV exposure [98,99]. Based on the presence of carotenoids in human skin and the known biologic activities of carotenoids, including quenching singlet oxygen species, carotenoids have been hypothesized to protect the skin from UV damage.

5.1. Epidemiological Studies and Fruit & Vegetable Intakes

Epidemiological studies have provided some evidence that fruit and vegetable intake may improve skin health and decrease the risk of skin cancer. However, in most cases, the specific effect of an individual carotenoid was not evaluated. Nonetheless, they provide evidence to support a role for carotenoids, and other phytonutrients, in skin health.

In a cross-sectional study investigating the association between skin wrinkling and dietary intake, a significant negative association was found with eggs and green leafy vegetables [69]. Eggs are a highly bioavailable source or lutein [100] and green leafy vegetables are good sources of both β-carotene and lutein [89]. This relationship between carotenoids intake and skin wrinkling may be due to an ability to prevent extracellular matrix breakdown [101]. An association of green leafy vegetables with decreased risk of skin cancer has also been reported [102]. In this prospective study, a higher consumption of fruits and vegetables was related to decreased risk of SCC risk by 54%. This protective effect was mostly explained by the association with green leafy vegetables.

In an observational study of an Italian population, high intake (at least 3 times per week) of dark green leafy vegetables was associated with decreased risk of cutaneous melanoma after adjustment for sex, age, education, hair color, skin phototypes, number of nevi, presence of freckles and sunburns in childhood [68]. In another observational study which looked at BCC and SCC, a decreased risk of SCC in persons with a history of skin cancer and high dietary intake of lutein and zeaxanthin was reported [23]. Other studies show a mix of results including no association between serum lutein/zeaxanthin and BCC or SCC [24] and a positive association between serum lutein/zeaxanthin and risk of SCC in participants with a history of BCC [103]. Dietary lutein (combined with zeaxanthin) showed a slightly positive relationship with SCC and BCC incidence in women [104,105]. However, vitamins A, C, and E, folate and total carotenoids were also found to have a positive relationship. The mixed results of these studies could be due to limitations in dietary assessment or individual variability in the ability to accumulate certain phytonutrients in skin tissue.

5.2. β-Carotene

β-Carotene is in the carotene class of carotenoids (Figure 3). It is a strongly-colored red-orange pigment. As a carotene with beta-rings at both ends, it is the most common form of carotene. It is a precursor of vitamin A. Rich sources of β-carotene include yellow and orange fruits, such as mangoes and papayas, orange root vegetables such as carrots and yams and in green leafy vegetables such as spinach, kale, sweet potato leaves, and sweet gourd leaves [89]. β-Carotene inhibits free radical and
singlet oxygen-induced lipid peroxidation in liposomes and the biologic system. It is a photoprotective agent and is thought to quench photochemical reactions in the epidermis involving singlet oxygen and oxygen radicals generated by UV exposure [106].

**Figure 3.** Chemical structure of β-carotene.

5.2.1. Epidemiologic Studies

The epidemiologic studies that have evaluated a relationship between β-carotene and skin cancer risk have provided little support for a protective role of this carotenoid. No studies have found a relationship between plasma β-carotene and risk of BCC [107-114]. The few studies have examined SCC also found no relationships [110,112,114,115]. However, in a prospective study among 1001 randomly selected adults, dietary intake of β-carotene in the second tertile was associated with increased risk of BCC, with no trend [116].

5.2.2. Clinical Trials

The photoprotectiveness of β-carotene was first demonstrated with a study of healthy volunteers receiving 180 mg β-carotene/day for 10 weeks or a placebo [117]. Compared to the placebo group, threshold MED was significantly higher in the supplemented group, although there was no significant difference between the two groups in the degree of erythema. Similarly, in a placebo-controlled study involving 30 mg/day β-carotene for 10 weeks, there was a significantly decrease in the intensity of erythema induced by sunlight [118]. Lee _et al._ examined the effects of increasing does of β-carotene (30–90 mg/day for 24 weeks) on UV-induced erythema [119]. In a study evaluating the immune response to supplementation with either 30 mg/day β-carotene or a placebo was studied in 24 men (19–30 years)[120]. Subjects ingested the supplement with a low carotenoid diet for 70 days. From day 28, the subjects were exposed to UV light 12 times over 16 days. No reductions in plasma carotenoid concentrations were observed in either group. Delayed-type hypersensitivity (DTH) skin responses decreased in the placebo group but were unchanged in the supplemented group. The investigators concluded that β-carotene protected against photosuppression of the immune system and that BC supplementation may be beneficial in those with compromised immune function. 

Herraiz _et al._ conducted a similar study involving 32 elderly men (55–79 years) receiving either 30 mg/day β-carotene or a placebo with a low carotenoid diet for 47 days [121]. Repeated exposure to UV light resulted in a decrease in DTH skin responses in the placebo group but not in the supplemented group and plasma β-carotene concentrations were unchanged. Higher plasma β-carotene concentrations were related to higher resistance to immunosuppression after exposure to UV light.

The photoprotectiveness of β-carotene(24 mg/day) was compared to that of a carotenoid mixture of β-carotene, lycopene and lutein (8 mg/day each) in a 12 week intervention trial [122]. Erythema intensity after irradiation with a solar light simulator was determined at baseline and at 6 and 12 weeks
of supplementation. The intensity of erythema 24 hours after irradiation was significantly decreased to a similar level in both groups.

A recent meta-analysis of controlled clinical trials reported an overall protective effect of β-carotene supplements against sunburn [123]. Although the studies to date suggest that high dose supplementation with β-carotene may be of benefit to skin health, there is a question of safety with these high intakes. Two intervention trials involving individuals at high risk for cancer, there was a significantly increased incidence of lung cancer in those that received the β-carotene supplement [124,125].

5.2.3. Animal Studies

Minami et al. evaluated the role of lipid peroxidation products in MMP-9 activation and the effect of dietary β-carotene in MMP-9 activation. In this study, hairless mice were subjected to periodic UVA-irradiation for 8 weeks [126]. The amount of peroxidized cholesterol (primary product of lipid peroxidation in biomembranes) in the skin significantly increased by the UV exposure. The activity and protein level of MMP-9 were elevated, as well as the wrinkling and sagging formation. Adding β-carotene to the diet of the mice during the period of irradiation suppressed the activity and expression of MMP-9 along with the wrinkling and sagging formation. The amount of peroxidized cholesterol was also significantly lowered. These investigators concluded that dietary β-carotene prevents the expression of MMP-9, in part by inhibiting photodynamic action involved in the formation of peroxidized cholesterol, the result of which was less wrinkling and sagging.

5.2.4. In Vitro Studies

β-Carotene, along with the carotenoids astaxanthin, and canthaxanthin, were evaluated for systemic photoprotection in human dermal fibroblasts [127]. Fibroblasts were exposed to moderate doses of UVA which resulted in apoptosis and increased oxidative damage and expression of oxidative stress-response enzymes. When the carotenoids were delivered to fibroblasts 24 hours before exposure to UVA, astaxanthin exhibited a pronounced photoprotective effect and counteracted all of the measured UVA-induced alterations to a significant extent. β-Carotene only partially prevented the UVA-induced decline of catalase and superoxide dismutase activities, but it increased membrane damage and stimulated heme-oxygenase-1 expression. Furthermore, β-carotene dose-dependently induced caspase-3 activity following UVA exposure. However, canthaxathin had no effect on oxidative damage, except for heme-oxygenase-1 expression, which was increased. The data suggest that among the carotenoids tested, astaxanthin is the most protective towards photo-oxidative changes in cell culture.

5.3. Lycopene

Lycopene (Figure 4) is a carotene found in primarily in tomatoes but also in other red fruits and vegetables, such as red carrots, watermelons and papayas [89]. Unlike β-carotene, lycopene has no vitamin A activity [128]. Lycopene is considered to be the most the most efficient biological carotenoid singlet oxygen quencher [129]. A suggestion that lycopene may be protective in skin health
comes from the work of Darvin et al. in which it was reported that there was a significant correlation between skin roughness and lycopene concentrations in the skin [130]. Topical application of lycopene contained in monoglyceride or triglyceride microemulsions was found to result in lycopene penetration in the stratum corneum (6- and 3.6-fold, respectively) and in viable layers of porcine ear skin (from undetected to 172.6 and 103.1 ng/cm², respectively) compared to a control solution. Furthermore, the antioxidant activity of skin treated with lycopene was found to be 10-fold higher than untreated skin [131].

**Figure 4.** Chemical structure of lycopene.

5.3.1. Epidemiologic Studies

Serum or plasma concentrations of lycopene were found to not be related with either BCC or SCC [110,113,114]. These results are contrary to observations in clinical trials (see below).

5.3.2. Clinical Trials

Intervention trials have been conducted to evaluate the effect of lycopene supplementation on erythema prevention [132,133]. The lycopene intervention was for 10–12 weeks and volunteers consumed one of five sources of lycopene: tomato paste (16 mg/day lycopene); carrot juice (10 mg/day lycopene, 5.1 mg/day β-carotene; lycopene supplement (9.8 mg/day lycopene, 0.4 mg/day β-carotene); tomato extract (8.2 mg/day lycopene, 0.4 mg β-carotene); or synthetic lycopene (10.2 mg/day lycopene). In all studies serum lycopene significantly increased at study end. Skin carotenoids levels also increased with all interventions. However, only total carotenoids could be quantitated, as reflection spectroscopy was the method of assessment. With the exception of synthetic lycopene, all lycopene treatments were found to be effective in being photoprotective, as measured by erythema formation. Of note, is that the synthetic lycopene was the only intervention to contain no other carotenoids. Additionally, the tomato-based interventions contain other constituents including the carotenoids phytofluene and phytoene. It is possible that these other carotenoid contribute to photoprotectiveness since they absorb light in the UV range.

Darvin et al. evaluated the relationship between skin lycopene concentrations and skin roughness [130]. Lycopene was analyzed quantitatively on the forehead skin of 20 volunteers aged between 40 and 50 years. There was, no significant correlation between age and skin roughness but, a significant correlation was obtained between the skin roughness and the lycopene concentration (R = 0.843).

These findings suggest that higher levels of lycopene in the skin effectively lead to lower levels of skin roughness, however, since other antioxidants were not evaluated, it if difficult to conclude that this relationship was specific to lycopene.
5.3.3. *In Vitro* Studies

Lycopene has been shown to inhibit proliferation of several types of cancer cells through arrest of tumor cell-cycle progression, IGF-1 (insulin-like growth factor 1) signaling transduction, and induction of apoptosis [134,135]. Wu *et al.* [136] found that lycopene inhibited PDGF-BB (platelet-derived growth factor-BB)-induced signaling and cell migration in human cultured skin fibroblasts through direct binding to PDGF-BB. Trapping of PDGF by lycopene also compromised melanoma-induced fibroblast migration and attenuated signaling transduction in fibroblasts simulated by melanoma-derived conditioned medium. These findings suggest that lycopene may interfere with tumor–stroma interactions. These investigators concluded that the trapping activity of lycopene on PDGF suggests that it may act as an inhibitor on stromal cells, tumor cells and their interactions, which may contribute to its anti-tumor activity.

5.4. Lutein

Lutein is a member of the xanthophyll class of carotenoids (Figure 5). Carotenoids are a family of more than 600 lipid-soluble compounds that are plant pigments. All carotenoids have a backbone composed of conjugated double bonds which is essential for their antioxidant activity. Xanthophylls, the most polar of carotenoids, are unique to this family because of their functional hydroxyl groups.

Lutein is commonly found in fruits and vegetables. The richest source of lutein is leafy green vegetables but also can be found in significant amounts in broccoli, corn and peas [137,138]. Although levels are not high, lutein is also found in egg yolk, which is considered to be a highly bioavailable source of lutein [139]. There is no established Dietary Recommended Intake (DRI) for lutein, but 6 mg is associated with a reduced risk of cataracts and age-related macular degeneration [140-142]. Most Americans only get 1 to 2 mg/day [143-146].

Lutein has a well-established role in the eye health, acting as an antioxidant and absorbing blue light. Lutein also has an established presence in epidermal and dermal compartments of the skin and contributes to skin color [97].

![Figure 5. Chemical structure of lutein.](image)

5.4.1. Epidemiologic Studies

Epidemiological studies have provided some evidence that the level of lutein intake may improve skin health and decrease the risk of skin cancer.

In a cross-sectional study investigating the association between skin wrinkling and dietary intake, a significant negative association was found with eggs and green leafy vegetables [69]. This relationship between lutein intake and skin wrinkling may be due to lutein’s ability to prevent extracellular matrix breakdown [101]. An association of green leafy vegetables with decreased risk of skin cancer has also
been reported [102]. In this prospective study, a higher consumption of fruits and vegetables was related to decreased risk of SCC risk by 54%, but this protective effect was mostly explained by the association with green leafy vegetables.

In an observational study of an Italian population, high intake (at least 3 times per week) of lutein-rich dark green leafy vegetables was associated with decreased risk of cutaneous melanoma after adjustment for sex, age, education, hair color, skin phototypes, number of nevi, presence of freckles and sunburns in childhood [68]. In another observational study which looked at BCC and SCC, a decreased risk of SCC in persons with a history of skin cancer and high dietary intake of lutein and zeaxanthin was reported [23]. Other studies show a mix of results including no association between dietary lutein-zeaxanthin and BCC or SCC [24] and a positive association between serum lutein-zeaxanthin and risk of SCC in participants with a history of BCC [103]. Dietary lutein (combined with zeaxanthin) showed a slightly positive relationship with SCC and BCC incidence in women [104,105]. However, vitamins A, C, and E, folate and total carotenoids were also found to have a positive relationship. The mixed results of these studies could be due limitations in assessment of dietary lutein or individual variability in the ability to accumulate lutein in skin tissue, as has been noted for the macula [147-149].

5.4.2. Clinical Trials

Results from clinical trials looking at photoprotection support a role for lutein in skin health. One of the earliest of these studies showed oral supplementation with 24 mg carotenoids/day (including 8 mg/day each β-carotene, lutein and lycopene) improved UV-induced erythema in humans after only 6 weeks of supplementation in a 12 week trial [71]. Others have shown an increase in skin hydration and superficial skin lipids with twice daily oral supplementation (45 mg vitamin C, 5 mg tocopherol, 3 mg lutein and 2.5 mg α-lipoic acid). Also, skin lipid peroxides (a measure of oxidation) were decreased [150]. While these studies suggest a role for antioxidants in skin health, it is difficult to determine the individual contribution of lutein.

Palombo et al. [150] provide strong evidence that increased lutein intake may improve skin health. This double-blind placebo controlled trial was designed to study the efficacy of oral supplementation (lutein 10 mg/day, zeaxanthin 0.6 mg/day) or topical application (lutein 100 ppm/day, zeaxanthin 4 ppm/day) on five physiology parameters (surface lipids, hydration, photoprotective activity, skin elasticity, skin lipid peroxidation). This was a 12 week intervention. Oral or topical administration of these carotenoids significantly improved these measures, with oral administration providing better protection when measured by changes in lipid peroxidation and photoprotective activity in the skin following UV irradiation. However, the combined oral and topical administration provided the greatest protection.

5.4.3. Animal Studies

A study in hairless mice has shown protection from photocarcinogenesis (both number and size of tumors) with lutein supplementation [151]. In this study, mice received either a lutein/zeaxanthin-supplemented diet or a standard nonsupplemented diet. Dorsal skin of female Skh-1 hairless mice was exposed to UVB radiation in doses required for photoaging and for
photocarcinogenesis. Clinical evaluations were performed weekly, and the animals were sacrificed 24 h after the last UVB exposure. For photoaging experiments, skin fold thickness, suprapapillary plate thickness, mast cell counts and dermal desmosine content were evaluated. For photocarcinogenesis, samples of tumors larger than 2 mm were analyzed for histological characterization, hyperproliferation index, tumor multiplicity, total tumor volume and tumor-free survival time. Results of the photoaging experiment found that skin fold thickness and number of infiltrating mast cells following UVB irradiation were significantly less in lutein/zeaxanthin-treated mice when compared to irradiated animals on the standard diet. In the photocarcinogenesis experiment, tumor-free survival time increased, tumor multiplicity decreased and total tumor volume decreased in lutein/zeaxanthin-treated mice compared to control irradiated animals fed the standard diet. The results from this study suggest that dietary lutein/zeaxanthin supplementation protects the skin against UVB-induced photoaging and photocarcinogenesis.

5.4.4. In Vitro

Lutein protected rat kidney fibroblasts from UVA-induced oxidative stress, including decreases in the antioxidant enzymes, catalase and superoxide dismutase [152]. Furthermore, in human lens epithelial cells, pretreatment with lutein decreased UVB-induced lipid peroxidation and activation of p38 and JNK [153]. Lutein significantly inhibited MMP-1 expression in dermal fibroblasts and melanoma cells [101]. Results from these in vitro studies lend mechanistic support to a lutein’s role in skin health.

6. Conclusions and Summary

Skin cancer is a significant public health concern. The skin is vulnerable to UV damage, which can lead to cancer. Possible mechanism by which this occurs include oxidative damage and inflammation. A diet rich in phytonutrients, including α-tocopherol, flavonoids, β-carotene, lycopene and lutein, may produce continual whole body protection from such damage. Epidemiological studies support a role for diets high in these food components and decreased risk of photoaging and cancer. However, the data are not entirely consistent. Stronger evidence comes from intervention studies which find supplementation with α-tocopherol, flavonoids, β-carotene, lycopene, lutein and various nutrient combinations can protect from shorter-term markers of UV damage including lipid peroxidation, erythema, MMP expression, DNA damage, and apoptosis. Similar results are observed from topical treatments of these nutrients. Further investigation of optimal doses and mechanism of protection are needed to better target and prevent photodamage with dietary and/or topical treatments.

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