Potent Role of Antioxidant Molecules in Prevention and Management of Skin Cancer

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Abstract

Evidence reported that cancers are spreading every nook and corner of the world at an alarming rate. Skin diseases like chronic skin inflammations, psoriasis and skin cancers have also been burning topic in today. Protections of a biological system are often hampered while skin gets damaged. Factors like UV, radiation, viruses, chronic diseases, genetic predispositions, food habits and environmental exposures might lead to skin cancers. In addition to these, urbanization and globalization may also contaminate the environment that may eventually modify several biological and genetic functions. USA, Europe and Australia are in the most dangerous zone to be exposed. We basically performed detailed search of PubMed, Google Scholar and Science Direct for literature search and collecting related information. On the other hand, experiments suggested that antioxidant components such as phenolic acid derivatives, flavonoids and flavonol found to be preventive against cancer cell proliferations. Moreover, antioxidants have been also evaluated as a protective agent against chronic inflammatory diseases as well. These molecules may participate as an additional therapy which could exert synergistic effects while applying with other chemotherapeutic agents. Our literature findings and hypothetical figure may establish a good correlation between skin cancer and antioxidant therapy. Therefore this study will be focused on skin cancer biology and some possible management strategies using antioxidant phyto-nutrients.

Keywords: Skin; Radiation; Cancer; Chemotherapy; Free radicals; Antioxidants

Introduction

Skin is the most outer part of the body which also known as the largest organ of the body. It protects internal organs, heals automatically and lasts a lifetime [1]. However, mammalian skin is composed of generally epidermis and dermis. Skin helps in several communications like absorption, excretion, heat regulation and water resistance. In addition, these layers prevent and protects from injuries, pathogens, heat and other harmful exposures [2,3].

According to National Center for Health Statistics, around 1,685,210 new cancer population and 595,690 cancer deaths are projected to occur in United States during 2016 only [4]. Currently, cancers have been a real burden and the cost of this life threatening situations has become unattainable for the lower and middle class people; as a result, affected subjects lose their property as well as savings. Similarly, it is also a major challenge to the health care providers to ensure the effective treatment for everyone though researchers are working very hard to discover competent remedies to minimize this concern [5].

However, chronic exposures, human papipoma virus, UV-radiation, and dietary intake have been identified as the major culprit of skin cancer. Several reports found that using moisturizing cream may increase the risk of developing skin cancers through UV [6,7]. Melanoma, squamous-cell skin cancer and basal-cell skin cancer are observed mostly in today's medical investigations. Basal cells constantly divide to form daughter cells to replace the squamous cells. This type of cancer is observed through skin cells in the lowest layer of the epidermis which is the most seen skin carcinoma [8]. Squamous-cell skin cancers are the second most prominent owing to UV exposures; hence, clinical appearance is highly variable and often has the tendency to spread towards other organs [9]. On the other hand, excess sun burn in the childhood and adulthood may cause Melanoma [10] which further linked with melanoma once there is any mutations in the germ line p16 are occurred [11]. Several sign and symptoms have been reported on these such as burning, itching, moderate to severe pain, ulcer, edema, redness and often bleeding [12].

Free radical often reacts with cell membrane, endoplasmic reticulum, necessary enzymes and other genetic materials result in cellular necrosis. In addition, necrotized cells often secret various potent pro-inflammatory markers like Tumor necrosis factors,
Where UV A and UVB found to have mostly responsible for skin carcinogenesis, and thereby identifying non-toxic strong antioxidants to diminish skin cancer is an important area of investigation. Antioxidants strongly prevent both photo-carcinogenesis and skin tumor progression in animal model, in part, by reducing free radicals and reactive oxygen species. These molecules have also been proven highly potent by inhibiting endogenous TNF-α in animal skin, a prime mediator in skin tumor initiator. Furthermore, evidences often suggested that antioxidant inhibit mitogenetic and cell survival signaling and initiate apoptosis for cell suppression. Finally, phenolic acid derivatives and flavonoids effectively modulate cell-cycle regulators and check points toward reserve of proliferation, and often arrest growth in G0–G1 phases of a cell cycle [17-19]. Therefore, this study will describe the probable molecular reasons for developing skin cancer along with that a series of therapeutic approaches based on antioxidant molecules will be exposed to prevent skin carcinoma.

Skin Cancer

Cancer, one of the most complicated system in a biological subject which deals with several factors such as alteration of DNA bases, mutation in genetic materials, mutation or blocking of tumor suppressive gene p53 [20,21]. As a result, normal cell of the body lost their control and start replicating their progenitor cells for an enormous numbers which finally act as immortal cells [22]. Every cell has some predetermined functions and they remain in one place for their own survival and biological importance but in cancer, these cells have the ability to travel or spread toward other organs and take control over there. These cells then become malignant tumor and may be threat for a subject [23]. In today’s world, men are more prone to prostate cancer, and women are mostly vulnerable towards breast and ovarian cancers but it is skin cancer which affects both the genders [24]. Merkel cell carcinoma, Kaposi sarcoma, Cutaneous (skin) lymphoma, Skin adrenal tumors and different types of sarcomas are rarely observed on skin [25,26]. The most common reason for skin cancer is either sunburn or excess UV exposure, however, family history also plays a major role [27]. Several reports have been suggested where excess UV can cause skin carcinoma in the laboratory animals [28,29]. So far UVA, UVB and UVC have been investigated where UVA and UVB found to have mostly responsible for skin carcinoma, on the contrary, UVC can’t reach to the earth due to protection of Ozone layer. Extreme Sun light/UV may penetrate to the skin nucleus and alter/damage genetic code which further leads to prostate cancer, and women are mostly vulnerable towards breast and environmental toxins like smoking, high oxygen atmosphere, which deals with several factors such as alteration of DNA bases, mediator in skin tumor initiator. Furthermore, evidences often suggested that antioxidant inhibit mitogenetic and cell survival signaling and initiate apoptosis for cell suppression. Finally, phenolic acid derivatives and flavonoids effectively modulate cell-cycle regulators and check points toward reserve of proliferation, and often arrest growth in G0–G1 phases of a cell cycle [17-19]. Therefore, this study will describe the probable molecular reasons for developing skin cancer along with that a series of therapeutic approaches based on antioxidant molecules will be exposed to prevent skin carcinoma.

Limitations of Chemotherapies

The prevalence of cancer in USA and other western countries is very high and often found tumor who lead normal life [36]. At the initial stage of the tumor progression the treatment on any cancer is easier and to some extend very much curable, however once it reaches towards the metastasis it become more complex and it most cases the treatment seems to be more resistant and the chances of cure become very narrow. Currently, three types of treatment protocol are now being used in the clinic to treat cancer. Radiation which kills individual cells, full or partial removal of tumor cells by surgery and chemotherapies which work against particular cells [37]. In contrast, the treatment of cancer is becoming very difficult owing to following different patterns by the tumor cell. Tumor cells are very distinct and always vary within individuals [38,39]. In addition, it is really difficult to deal with cancer subjects because cancer is a constellation of at least 200 other diseases and it has been often reported that cancer cells are not homogenize [24]. It is always suggested that cancer therapies should optimize most tumor cells and it must be monitored for minimal effect to the normal cells [40]. Unfortunately, all cancer therapies produce severe side effects, adverse effects and are very costly. In majority if the cases, single drug therapy is not sufficient to treat carcinoma. Multiple therapies with several cycles are necessary to optimize the cancer cells. Alopecia, bone marrow depression, severe vomiting, fatigue, rash and heart failure are associated with chemotherapies [41,42]. Most of the anti-tumor drugs are not well studied and many of these are not approved by FDA. Anti-cancer molecules are often contraindicated during pregnancy and several studies suggested that these drugs may interact with foetus and may cause death [43,44]. Besides, several deaths have been reported while given chemotherapies on many clinical evaluations [45,46]. Hepatic and kidney failure have been documented when platinum based chemotherapy was provided [47,48]. Cardiac arrest or arrhythmia have been documented when monoclonal antibodies were administrated to the cancer subjects and sometimes the subjects must be admitted in ICU during prior treatment which increase overall cost [49-51].

An Overview of Antioxidants

Oxidative stress mostly found to be free radical-mediated which further correlates with the response of cellular metabolism and catabolism [52]. In addition, free radicals are mainly generated from mitochondria during ATP production and cellular response in a biological subject. Sometimes drug molecules produce free radicals when given in higher dosage [53]. There are numerous evidences where free radical-mediated oxidative stress linked with many diseases like cardiovascular dysfunctions [54], hepatic damages [55], renal injury [56], diabetes [57], neuro-degeneration [58], aging [59] and other life threatening events [60]. Free radicals are often blamed for acute and chronic inflammation [61] which further drawn with cancers [62,63]. Several approaches against cancer cells have been undertaken targeting reactive oxygen species and often found good results while taken with other synthetic drugs [64]. On the other hand, antioxidants are the molecules which fight against oxidative stress. These are also known as ‘free radical scavengers’. Basically, polyphenols and flavonoids are considered as anti-oxidants which exert several protective mechanisms and found to be effective among in vivo, in vitro and human subjects [65]. Foods and supplements based
antioxidants have been proved beneficial against free radical mediated damages, at the same time they protect nucleus and necessary enzymes [57,66]. There are multiple studies have been undertaken to investigate anti-cancer activities of antioxidant molecules and some breakthrough outcomes were successfully documented [67,68].

**Role of Antioxidant on Skin Cancers**

Skin, the biggest organ of the body is protected by several layers. Multiple protective mechanisms have been proposed so far through animal and human studies. Chronic UV exposure-induced skin diseases are mainly caused by the excessive oxidative stress that further leads to inflammation and finally results in DNA and necessary protein damages [35]. DNA protection using antioxidants therapy has been a good choice for clinical trials. Phenolic acid and flavonoids are being reported extremely well against chemoprevention [69]. It is reported that in the TPA-induced skin inflammation model citrus peel extract prevented expression of inducible nitric oxide synthase, cyclooxygenase-2, ornithine decarboxylase, and vascular endothelial growth factor in mouse skin. The study also noticed inhibitory effects of citrus on 7,12-dimethylbenz[a]anthracene (DMBA)/TPA-induced skin tumour formation and reduced tumor incidence, tumor weight and tumor multiplicity of papillomas at 20 weeks treatment [70]. The blocking properties of naringin were observed when in BALB/c mice skin irradiated with UVB. The expressions of interleukin-1β (IL-1β), interleukin-6 (IL-6), interleukin-8 (IL-8) and cyclooxygenase-2 (COX-2) were significantly lowered in Ha CaT cells by primarily blocking MAPK pathways [71]. Src family kinases (SFKs), basically known as oncogenes, are recently drawing attention for their association in melanoma skin cancers; on the other hand, naringin may be an effective molecule for the treatment of melanoma as a natural inhibitor of c-Src [72]. Retinoids such as 13-cis-retinoic acid and retinyl palmitate exhibit a good preventive effect in chemically induced papillomas and carcinomas of the skin in female Swiss mice after three weeks of induction of cancer. The study showed inhibitory effects of retinoids on skin carcinoma [73]. Topical application of vitamin E in a dose-dependent manner has been shown to reduce the incidence of UV-induced skin cancer in mice by exhibiting antioxidant mechanism. Study shows that dimer and trimer products may contribute in avoidance of UV-induced photo-damage in animal [74]. Vitamin C was also found as photo-protectant while applied to mice and pig skin before exposure to ultraviolet radiation (Figure 1) [75,76].

Figure 1: Role of anti-oxidants on skin cancer cell.

A potent antioxidant molecule, Coenzyme Q 10, generally exerts several mechanisms in cancer, inflammation and aging. When Coenzyme Q10 applied on human dermal and epidermal cells, it protects against cell death induced by reactive oxygen species in keratinocytes [77]. A population-based case-control study was performed to investigate the correlation between citrus peel use and black tea intake on squamous cell carcinoma of the skin tissue. Preventive effects of citrus peel and black tea were statistically noticed on human subjects [78]. A study was undertaken with a large number of healthy women whom solar stimulated UV were given on the skin and β-carotene (120 mg) was provided. The study found to be protective by inducing skin lycopene and reducing oxidative markers [79].

The hypothetical pathway explains that there are various extracellular effects can activate the signaling of the growth factor. One of the major reasons of skin cancer is the UV-B from UV radiation. UV-B can influence the mutation of K-RAS which is a upstream signaling kinase of MAPK family. Even though K-RAS signaling is essential for normal cell proliferation and apoptosis but when mutated it is a key player of activating certain signaling pathway that leads to the binding of different transcription factor with DNA that will cause the activation of different cytokines which will participate in the
progression of cancer cell. It has been most commonly known that chronic inflammation leads to the progression of cancer. In skin cancer many studies suggested that activation of TNFR1 by its binding with TNF-α facilitate the synthesis of different inflammatory cytokines. All these factors together participate in the progression of tumor cell. White people are more vulnerable towards the skin cancer than the black and mixed race people. One of the possible reason is the production of melanin is really low in the skin of white people compared to black. However, certain polyphenols and flavons can prevent the mutation from taking place. The underlining reason is yet unknown, due to the lack of study in this particular area it is yet to be discover, but one hypothesis is the hyper activation of the growth factor can be prevented by polyphenols and flavons. They also prevent the phosphorylation of certain kinases, which leads to the apoptosis of those cells. All these activity made them a potential molecule against skin cancer (Table 1).

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Outcomes of the study</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model: Solar simulated irradiation of skin</td>
<td>Caspase 3, 7 density were lowered</td>
<td>[80]</td>
</tr>
<tr>
<td>Disease induced by: Radiation</td>
<td></td>
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<tr>
<td>Treatment: Ferulic acid</td>
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<tr>
<td>Dose: 0.5% Ferulic acid</td>
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<tr>
<td>Model: Human keratinocytes</td>
<td>Decreased DNA synthesis, cell growth inhibition, DNA synthesis and Blockade of the cell cycle at the S/G2-M phase and cell death by necrosis.</td>
<td>[81]</td>
</tr>
<tr>
<td>Disease induced by: Se methyl selenocysteine</td>
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<td></td>
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<tr>
<td>Treatment: CusO₄</td>
<td></td>
<td></td>
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<tr>
<td>Dose: N/A</td>
<td></td>
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<tr>
<td>Model: CD-1 mice</td>
<td>Decreased protein expression of ODC, cox-2, nitric oxide synthase, and blocked NF-κB and phosphatidyl inositol 3-kinase (PI3K)/Akt signaling in tumor promotion</td>
<td>[82]</td>
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<tr>
<td>Disease induced by: Genetic</td>
<td></td>
<td></td>
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<tr>
<td>Treatment: Lupeol</td>
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<tr>
<td>Dose: 1–2 mg/mouse</td>
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<tr>
<td>Model: Heterozygous DBA/2 MnSOD KO mice</td>
<td>suppressed mitosis without interfering with apoptosis, decreased protein carbonyls and reduced the activity of activator protein-1, and prevented the level of proliferating cellular nuclear antigen</td>
<td>[83]</td>
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<tr>
<td>Disease induced by: 12-O-tetradecanoylphorbol-13-acetate</td>
<td></td>
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<tr>
<td>Treatment: SOD mimetic</td>
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<tr>
<td>(MnTE-2-PyP5+)</td>
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<td>Dose: 5 ng/mouse</td>
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<tr>
<td>Model: Human</td>
<td>Changed in lipid peroxidation and photo-protective activity in the skin.</td>
<td>[84]</td>
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<tr>
<td>Disease induced by: UV light irradiation</td>
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<tr>
<td>Treatment: Lutein and zeaxanthin</td>
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<tr>
<td>Dose: Lutein 5 mg/zeaxanthin 0.3 mg 2/day</td>
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<tr>
<td>Model: Mouse</td>
<td>inhibited UVB-induced oxidative stress, significant prevention of UVB-induced depletion of antioxidant enzymes, and Inhibition of a single UVB irradiation-induced phosphorylation of ERK1/2 and MAPK family.</td>
<td>[7]</td>
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<tr>
<td>Disease induced by: Solar ultraviolet (UV) radiation</td>
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<td>Treatment: (-)-Epigallocatechin-3-gallate (EGCG)</td>
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<td>Dose: 1 mg/cm² skin area</td>
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<tr>
<td>Model: Skh:2 Hairless Mice</td>
<td>showed no signs of toxicity and had significantly less acute and chronic skin damage induced by UV irradiation, and reduced inflammation and pigmentation.</td>
<td>[29]</td>
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<tr>
<td>Disease induced by: Ultraviolet Irradiation</td>
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<tr>
<td>Treatment: Vitamin E</td>
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<td>Dose: 5% oral and topical RRR-α-tocopheryl acetate</td>
<td>reduction of the UV-induced erythemas, and Parallel reduction of the lipoperoxide level was observed.</td>
<td>[85]</td>
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<tr>
<td>Model: Human</td>
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<tr>
<td>Disease induced by: Acute UV radiation</td>
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<tr>
<td>Treatment: Antioxidant complex – vitamins (lycopene, β-carotene, α-tocopherol), selenium</td>
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<tr>
<td>Dose: 3 mg of tomato lycopene, 3 mg of natural α- and b-carotene 5 mg of natural α-tocopherol and 37.5 mg of organic selenium</td>
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</tbody>
</table>
### Table 1: Role of antioxidant molecules on various cancer induced skin cells.

| Model: Mice | Disease induced by: UV-induced skin cancer | Treatment: Vitamin E and epigallocatechin gallate | Dose: Olive oil on the dorsal skin with a moist cotton swab | Delayed the onset of and to reduce the incidence rate of skin cancer development. | [28] |
| Model: SK-mel28 and A375 human melanoma cells | Disease induced by: Genetically induced | Treatment: Resveratrol | Dose: 30, 60, or 100 mM all-trans-resveratrol | Induced phosphorylation of ERK1/2 in A375 | [88] |
| Model: Healthy women | Disease induced by: Solar stimulated light | Treatment: β-carotene | Dose: 120 mg | Skin lycopene found to be increased and reduced oxidative damages. | [79] |
| Model: Human | Disease induced by: UV light | Treatment: Carotenoids and carotenoids plus vitamin E | Dose: 25 mg total carotenoids/d and 335 mg (500 IU) RRR-α-tocopherol/d | Dorsal black skin was significantly diminished | [87] |
| Model: Cell culture | Disease induced by: UV or ozone | Treatment: Vitamin E | Dose: N/A | Combated oxidative stress induced by UV | [88] |
| Model: Hairless mice | Disease induced by: UV exposure | Treatment: β-carotene | Dose: 10% carotenoids | Reduced skin tumor progression | [89] |

### Conclusion and Future Directions

Evidences suggested that oxidative stress plays a pivotal role in mutation of the different genes, and the expression of many cytokines and chemokines leads to the progression of tumor. It also prevents mitochondrial biogenesis as well, resulting cellular reproduction gets hampered; on the other hand, antioxidants found to be very useful for biogenesis. Although antioxidant molecules have been highly appreciated in the treatment of skin diseases like skin cancers, negative findings are reported too. Both animal and small clinical trials are showing good activities when antioxidant molecules are applied on various skin cancers. Additional larger and wide randomized clinical trials are needed to provide clear scientific reports about the potential benefits of taking antioxidant supplements during skin cancer treatment.

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### Conflict of Interest

The authors declare no conflict of interest.

### References


inflammation, fibrosis, iron overload, hepatic dysfunction and renal injury in acetaminophen induced rats. J Drug Metab Toxicol 7: 2-7.


