A Glimpse on Melanoma - Risk Factors and Treatment

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Abstract
Melanoma is the most dreadful form of skin cancer that begins in melanocytes. It is the leading cause of death from skin disease. It can occur in any part of the body that contains melanocytes. Several internal and external risk factors of the body may contribute to the development of many cancer melanomas such as unexpected exposure to UV radiation, weakened immune system, family history etc. Melanoma diagnosed at early stages is typically cured by surgical excision, but advanced stages are often deadly. Cancer stem cells (CSC) are recently proposed to be the cancer initiating cells responsible for tumorigenesis and contribute to cancer resistance. In the present review I have discussed about melanoma incidence, detection with various modes of treatment.

Keywords: Melanoma; Melanocytes; Skin cancer; Signs and Symptoms; Risk factors; Detection of melanoma; Therapies for treatment

Abbreviations: MM: Malignant melanoma; QOL: Quality of life; UV: Ultraviolet; ART: Adaptive radiotherapy; TRT: Targeted internal radionuclide therapy TSC: tumor stem cells; CSC: Cancer stem cells; BCG: Bacillus Calmette-Guerin; MSCs: mesenchymal stem cells; ABC: ATP binding cassette

Introduction
Cancer is the condition where cells are marked by their unregulated proliferation [1]. It is the major cause of death in the world after cardiovascular diseases [2]. It generally develops when normal cells in a particular part of the body begin to grow out of control. Tumor consists of numerous cancer cells, the size and shape of a tumor may not be an accurate measure as it is an abnormal growth [3]. The increase in the tumor size also increases the risk of complications and chance of reoccurrence [4].

There are different types of cancers; all types of cancer cells continue to grow abnormally. Some types of cancer cells often travel to other parts of the body through blood circulation or lymph vessels (metastasis), where they begin to grow.

Human skin regulates heat and water loss from the body [5], while preventing the access of deadly chemicals or microorganisms. Skin of healthy individual comprises of three main layers: subcutaneous tissue, dermis and epidermis [6]. Skin is also affected by psoriasis [7], Kaposi’s sarcoma [8] linked with inflammation, and angiogenesis and immunosuppression.

When an injury occurs, the skin repairs itself through the proliferation and growth of dermal and epidermal cells [9]. But in deep skin lesions, destruction of the dermal and epidermal layers may take place thus the regeneration process is slowed down and complications arise. The first step for production of reconstructed skin involves the proliferation of dermal (fibroblasts) and epidermal cells (keratinocytes and melanocytes). Melanomas are usually an irregular pigmented lesion of varying size, but appearance may be atypical [10]. In case of metabolism of cancer cells and also for cell proliferation mitochondria plays a vital role [11].

In the treatment of large skin injuries, a higher number of applicable fibroblasts and keratinocytes are required [12]. In case of human keratinocytes media formulations, factors such as insulin, transferrin and lipid emulsion (containing cholesterol, α-tocopherol acetate, fatty acids, Tween 80, and Pluronic F-68) should be assessed [13].

Melanoma
Melanoma is the alarming and most aggressive type of skin cancer which begins in melanocytes. Melanocytes are the cells that make the dark pigment called melanin which is responsible for the colour of skin. Melanoma may not be confined only to skin but also spreads to other pigmented tissues of the body, such as eye, bowel, and the intestines [14]. If it is confined to the skin, it can often be cured by surgery. However if it has spread, melanoma is usually incurable because it does not respond to most treatments. Recently clinicians have been trying a combination of chemotherapy and immunotherapy in the hope of improving the outcome [15].

The progression of Cancer depends upon physiological changes [16]. Metastatic melanoma may cause nonspecific paraneoplastic symptoms, including loss of appetite, nausea, vomiting and fatigue [17]. Metastasis of early melanoma is possible, but relatively rare: less than a fifth of melanomas diagnosed early become metastatic. Brain metastases are particularly common in patients with metastatic melanoma [18].

Though the incidence of Melanoma is still increasing, the mortality rate has remained unchanged. Alteration of Lymph node is the only most important prognostic factor for stage I/II melanoma patients. Right now, the standard of care with regard to the staging of these patients is the surgical sentinel node procedure. Ultrasound is not customary for the diagnostic work-up of primary melanomas [19]. In light of advanced and promising adjuvant therapies, the need for ultrasound staging might increase at speed. The advanced stages of melanoma are inevitably resistant to common therapeutic agents.

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Received July 12, 2011; Accepted September 24, 2011; Published September 26, 2011

Citation: Soumya D (2011) A Glimpse on Melanoma - Risk Factors and Treatment. J Cancer Sci Ther S17. doi:10.4172/1948-5956.S17-004

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In particular, epigenetic approach [20] might open interesting perspectives in combination with conventional ones. In my view, further investigations are needed to be conscious of the toxicity of these treatments in order to promote the survival of melanoma patients.

The therapeutic options remain confined for advanced MM, and those directed to the neoplastic cells have not brought major survival advantage so far. Immunotherapy is another focused choice [21]. By incorporating active and passive target techniques we can improve the efficiency of imaging techniques, allowing for earlier diagnosis and treatment of lethal conditions such as malignant growth [22]. Malignant tumors such as breast cancer, gastric cancer may be metastatic cause and basal cell carcinomas, melanoma, and angiosarcoma can be localized cause [23].

There are various types of Melanoma, which include the following.

- Superficial spreading melanoma: This type of melanoma is more commonly observed in the locations such as the legs of women and the backs of men. These are often barely raised and have a variety of colors.
- Nodular melanoma: This type of melanoma is known to progress faster and more likely to spread. Approximately 20% of melanomas take their origin as deeper, blue-black to purplish lumps.
- Lentigo maligna: This type of melanoma tends to occur on locations like the face, which are exposed to the sun regularly. They appear like a large, irregularly shaped or colored freckle and develop gradually. It takes many years to evolve into a fatal melanoma.
- Acral-lentiginous melanoma: This is the more common type of melanoma in nonwhites. Often they appear as irregular enlarging black flat spot (macule) on the palm and sole, rarely on a mucosal surface, such as the vulva or vagina.

Other occasional forms of melanoma may occur under the nails, on the palms and soles, in the eye, or at times even in the internal parts of the body.

**Risk Factors of Melanoma**

Many human organs can take several years to turn cancerous and have many causative factors. In industrialized nations cancer is more common, but there has been a growth in cancer rates in developing countries also, as these nations adopt the diet and lifestyle habits of industrialized countries [24].

Several internal and external factors of the body may contribute to the advancement of many types of cancer. Also hormonal influences may be responsible for the carcinogenic risk [25]. With reference to the role of immune system in cancer, psychosocial influences on immune function provide a mechanism of association between psychosocial factors (like interpersonal aggression) and prediction of cancer [26]. The actual cause of melanoma is not known but the risk factors that increase development of the disease are known. Risk factors for melanoma include:

- Unprotected exposure to UV radiation from natural sunlight or artificial sources, such as tanning beds/lamps
- Serious blistering sunburns
- Having fair skin and also having naturally red hair
- Family or personal history of melanoma
- Bearing many suspicious moles
- Weakened immune system

Other risk factors for melanoma include age, gender. Gender has been shown to be an essential and independent forecaster of clinical outcome and survival in cutaneous melanoma, with premenopausal females (but not women older than 60 years) experiencing an improved prognosis [27].

Xeroderma pigmentosum, Giant congenital melanocytic nevi, Dysplastic nerveous syndrome, atypical nevi, many acquired, melanocytic nevi and immunosuppression are other risk factors. The absolute treatment is with surgical excision. Adjuvant therapies such as chemotherapy, immunotherapy, and radiation therapy can be used in advanced stages [28]. The occurrence of melanoma is increasing worldwide, with a growing fraction of patients with advanced disease for which diagnosis remain standing [29]. Therapeutic options are limited despite advances in immunotherapy and targeted therapy.

**Signs and symptoms:** Early signs of melanoma are changes to the shape or color of existing moles. At later stages, the mole may itch, ulcerate or bleed. Early signs of melanoma include:

- Irregularity of mole
- Blurred or ragged edges
- Alteration in the pigmentation of mole.
- Increase in the size and shape of mole
- Itching or tenderness of mole

Later symptoms include: pain in mole, swelling of lymph node, weight loss, chronic cough, gray skin, seizures etc.

**Preventive Measures**

The earlier the melanoma is diagnosed and treated, the better chance the patient has of surviving. But when left untreated, it spreads to other organs in the body. The high rate of metastasis is the main reason melanoma is considered the deadliest skin cancer. As “Prevention Is Better than Cure” measures should be followed to prevent occurrence of melanoma. Not all melanomas can be prevented, but can reduce the risk.

- The best way to reduce the risk of melanoma is to limit exposure to strong sunlight and other sources of UV light. And also tanning beds and sunlamps
- Protecting skin with clothing
- Usage of sunscreen

Vitamin D has many health benefits and it may aid to lower the risk for some cancers. The major source of vitamin D synthesis in humans is through the skin exposed to solar UV radiation [30]. If possible, it is better to obtain vitamin D from diet rather than from the sun. Abnormal or suspicious moles have to be checked and removed. And routine follow-up is to be carried out [31]. If the moles are altering into suspicious skin lesions, medical attention should be accomplished in a right away [32]. Also skin, scalp, feet, nails, and genital area should surely be examined as melanoma can appear on parts of the body that
people do not consider checking. Genetic counseling and testing should be as a measure for reducing the risk of melanoma as gene mutation increases the risk of melanoma.

Detection of melanoma: Unlike infectious diseases, cancers arise from our own cells towards which the immune system has been tolerated. Therefore, effective immune responses against cancer would be challenging than it is against infectious diseases [33]. Generally, Melanoma [34] diagnosed at early stages is cured by surgical excision alone [35], but in advanced stages it is often fatal [36]. It is crucial to detect melanoma at early stages as surgical excision is the only means of life saving. However, analyzing benign pigmented lesions and early melanomas can be challenging and hence there is a need for reliable markers that would boost in the diagnosis [37]. Still it is unexplored whether screening for early detection of melanoma enhances survival [38].

Treatment

Current advances in the diagnosis and treatment of cancer have led to an increase in the survival and hence, there is a greater emphasis on quality over quantity of survival [39]. Effective treatment for advanced melanoma is still lacking. And hence it is better to limit exposure to ultraviolet radiation which remains the most effective way to reduce the risk of melanoma [40]. In-vivo testing devices include the devices or sensors that can be used to detect cancer [41].

Therapies for Treatment: Traditional approaches for the encounter of cancer include surgery, chemotherapy, and radiation therapy [42]. A new idea in cancer therapy is to enhance the selectivity of cytotoxic or cell-killing molecules by directly delivering them to cancer cells via the targeting of antigens that are unique to or highly expressed on the surface of cancer cells [43]. Fibroblast-derived cytokines have an impact on the immune system in tumor tissue [44].

The treatment becomes more complex when melanoma grows deeper into the skin or spreads. Surgeries may be needed to treat. Other treatments for melanoma include:

- **Lymphadenectomy:** This is the surgery which involves the removal of lymph nodes.
- **Immunotherapy:** This therapy aids the patient’s immune system fight the cancer e.g.: interferon injections.
- **Chemotherapy:** Therapy that kills the cancer cells using chemicals.
- **Radiation therapy:** Radiations are used to kill the cancerous cells in this therapy.

Cancer treatment involving chemotherapy has toxic side effects; hence drug dosage given to patient should be limited. As a result, all of the tumor tissue may not be exposed to a lethal dose of the drug. The use of nanocarriers like liposomes and micelles can enhance the pharmacological properties of conventional chemotherapeutics [45]. Chemotherapy is associated with higher toxicity and lower quality of life (QOL) [46]. The secondary metabolites from microbes play a crucial role in generating chemotherapeutics [47]. It is known that different microorganisms are the source for various anticancer molecules [48]. A combination of treatment with radiotherapy and chemotherapy, or surgery can improve the prognosis of patients [49].

Many malignant tumours are hypoxic in tumor regions. This is clinically important as tumor hypoxia may increase resistance to radiation therapy and also to some forms of chemotherapy [50]. HIF-1α is involved in inflammatory pathways and angiogenesis and provides a therapeutic target. It also acts as a clinical biochemical marker for diagnosis of cancer [51].

An efficient and alternative to the current therapies for disseminated melanoma treatment is Targeted internal radioisotope therapy (TRT) [52]. Adaptive radiotherapy (ART) is a feedback method of treatment that optimizes according to patient-specific information measured during the course of treatment [53]. Cutaneous metastases from carcinoma are relatively uncommon in clinical practices when compared to metastases in various other organs of the body [54]. In the effective method of treatment for the metastatic melanoma cells to various other organs, Yttrium-90 microspheres radioembolization can be used [55]. Several toxicities and complications arise with radioembolization [56]. Epigenetic cancer therapy is becoming one of the most widely studied therapeutic agents for a vast array of diseases, including cancer [57].

The recent targeted biologic therapies will block essential functions of cancer cells and tumour stroma. A growing number of therapies, alone or in combination with other therapies (chemotherapy, hormone therapy, radiotherapy), will be a better adaptation of treatment to patients and disease characteristics [58]. Photo dynamic therapy [59] which was first applied in 1970s is another mode of control and treatment of melanoma cancers. It would be effective in combination with conventional methods. Tissue-engineered skin or Bioengineered skin can be used as an alternative to add various other cell types (keratinocytes, melanocytes, adipocytes, endothelial and immunological cells, etc.) according to needs [60].

In case of defective expression, DNA repair genes may affect the status of tumor DNA repair causing resistance to therapy, and affect the outcome of cancer and survival of patients. DNA repair enzymes and DNA repair inhibitors [61] are emerging as therapeutics in cancer diagnosis and development of cancer biomarkers to fight against cancer.

Anti-cancer drug resistance could be developed as a strategy for research based on the evolution of drug resistance in malignant tissues [62].

**Vaccines:** Vaccines remain a promising but experimental treatment for melanoma [63]. Cancer vaccines, activating the immune system against specific antigens, have demonstrated clinical benefit and are now in development [64]. Vaccines are a novel means in the treatment of melanoma. They are unique in their potential ability to induce long-lasting immune responses that can selectively attack and kill tumor cells. At present a variety of vaccines are being tested in clinical trials. Melanoma vaccines have minimal toxicity and appear to be much safer than traditional therapy with interferon that causes serious toxicity in up to two thirds of patients. Vaccines can induce antibody and/or cellular immune responses. Early trials of BCG vaccine-based immunotherapy for melanoma consistently show a trend toward improved clinical results in patients treated with BCG compared with observation alone [65].

TLR (Toll-like receptors) are exploring to be used in combination with cancer vaccines and hold significant promise for enhancing the efficacy of immunotherapeutics for promoting tumor-specific immunity [66].

Immunoglobulins with undefined specificity [67] are required
for the development and viability of cancer cells from various tissue origins in human. Therefore, target-specific anti-cancer drugs can be developed, if we can identify unique biomarkers that can differentiate cancer cell-expressed immunoglobulins from those originated from normal B cells.

**Palliative care:** This care does not treat the cancer but can relieve symptoms and improve a patient’s quality of life. When melanoma spreads, palliative care can aid in controlling the pain and other symptoms.

**Drugs:** Melanoma is a kind of cancer which represents about 1% of all tumors and is responsible for 75% of skin cancer deaths. Dacarbazine is currently the only approved chemotherapy drug for the standard treatment of metastatic melanoma stage IV, but its therapeutic response is not satisfactory and only 15-20% of patients with favorable prognosis have increased survival rate [68].

- Melanoma can become a model in cancer therapy. At present there are two drugs namely ipilimumab and vemurafenib with different characteristics and effectiveness. Vemurafenib acts immediately, with a median latency of 6-7 months before developing resistance, ipilimumab with a slower (months) but more lasting (years) action [69].

- For the future success of melanoma therapeutics, the use of combination treatments in which different regulatory pathways or the immunological response are targeted seems to be a promising tool [70].

- Clinical trials and preclinical testing are the basis for the standard approach to evaluate novel compounds for cancer treatment after drug synthesis or discovery [71].

- A new technology of novel double lumen catheter for drug delivery in skin is on its trails for its applications in humans [72]. It aids in preventing infections.

Natural sources i.e. honey can be used against cancer and invasive infections. Many anticancer compounds obtained from *Ziziphus sp.* such as betulnic acid are selective inhibitor drugs for the growth of human melanoma [73] cell lines by causing apoptosis.

Cancer stem cells (CSC) are the cancer initiating cells responsible for tumorigenesis and contribute to cancer resistance. Tumor stem cells may display ATP binding cassette (ABC) transporters which have been reported as TSC markers in melanoma & osteosarcoma. For instance ABCB5 has been reported as a marker for a subset of CD133+ melanoma stem cells [74]. Cancer stem cells can arise from mesenchymal stem cells (MSCs) recruited to the tumor microenvironment [75]. MSCs [76] take their origin from fetus mesoderm layer and in the adult reside in different tissues like bone marrow stem cells (BMSCs), dermal stem cells etc. Human MSCs in the absence of ascorbic acid produce minimal amounts of collagen, which in turn aids in the inhibition of proliferation [77]. Tissue-specific stem cells could be the source of the original tumor. The existence of cancer stem cells is mostly accepted by the scientific community [78]. The identification of cancer stem cell with specific maintenance marker and its role in the differentiation would provide critical information for advancing towards the long-term goal of developing novel therapeutic strategies and will reduce the incidence of tumor recurrence for cancer patients [79]. Haematopoietic stem cells were widely used in transplantation experiments, especially in the treating leukemia and other cancers [80].

Melanoma cell line was cultivated in soft agar to check-out the effect of sugarcane cystatin in the anchorage-independent growth. Sugarcane cystatin CaneCPI-4 inhibits melanoma development *in vivo* by angiogenesis disruption and prevention of melanoma invasion, migration and anchorage-independent growth. CaneCPI-4 efficiently inhibited melanoma cell invasion *in vitro* using a Transwell invasion assay [81]. The current dogma in the field of cancer is that cell lines contain at least two subpopulations of cells: cancer stem cells and non-cancer stem cells [82]. Response rate, relapse free survival, disease stabilization rate overall survival, progression free survival in different phases are considered to be the principal primary endpoints used in advanced melanoma [83].

Proteomics has been employed recently to identify new disease related biomarkers for cancer diagnosis and development of targeted treatment [84]. Research studies are going for the employment of novel biomarkers in the diagnosis of cancers [85].

The ultimate goal of cancer therapy is to develop therapy with higher specificity for target tissues or cells. Certainly in the case of conventional chemotherapy agents such as doxorubicin, cisplatin etc., systemic administration results in cytotoxic effects not only into tumor cells, but also in healthy tissues.

**Emergence of Nanomedicine**

Nanomedicine has arrived into the cancer diagnostics and therapeutics [86]. Nanotechnologies cite a great opportunity to develop new products against cancer [87]. Nanotechnology definitely promises to serve as drug delivery carriers for the more challenging traditional drugs used for the treatment and management of chronic diseases such as cancer [88]. Nanoparticles or nanoporoporous particles functionalized with organic groups can be used as biomarkers, tracer, and drug delivery systems with much functionality for the treatment of cancer [89].

Using targeted nanoparticles to deliver chemotherapeutic agents in cancer therapy offers many advantages to improve drug or gene delivery thus to overcome many problems associated with conventional chemotherapy [90]. Liposomal nanoparticle therapeutics containing cytotoxic factors may provide the base for potentially more effective and less toxic anti-cancer treatment strategies due to their improved pharmacokinetics, reduced systemic toxicity, and increased intratumoral or intracellular delivery [91]. From the past few years there has been a growing attentiveness by various scientists to employ dendrimers for targeted delivery in cancer therapeutics [92].

**Conclusion**

In the present article I have discussed risk factors and various therapies for the treatment of melanoma. The use of combination treatments in which different regulatory pathways or the immunological response are targeted seems to be a promising tool for the future success of melanoma therapeutics. The Ultimate goal of cancer therapy is to develop therapy with higher specificity for target tissues or cells. Vaccines are a novel approach in the treatment of melanoma. Besides, there has been a growing attentiveness by various scientists to employ dendrimers for targeted delivery in cancer therapeutics [92].
Acknowledgement

I would like to thank my friends for their support in writing the review article.

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