Circadian disruption and cancer risk: A new concept of stromal niche (Review)

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Abstract. Circadian rhythms regulate a variety of physiological functions. Epidemiological evidence indicates that disruption of these circadian rhythms might be linked to cancer. In general, imbalances in homeostasis, such as immune and hormonal dysfunctions, are thought to be involved in cancer development. The results of a recent study suggested that circadian disruption may induce stromal changes associated with cancer risk, highlighting the importance of the cancer stem cell niche for protecting cancer cells. Current research provides new concepts and clarification regarding the function of the tumor niche, and the new concept of a stromal niche may help us to understand the additional functions of both cancer-associated fibroblasts and the extracellular matrix. In this review, we summarize our current knowledge regarding the role of circadian rhythms in cancer risk and the relevance of the stromal niche in cancer cell survival and progression.

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1. Epidemiological evidence

The use of electric lights means that people in the modern era are exposed to long photoperiods throughout the year. This disruption to the circadian system can induce a wide variety of stresses. Abnormal circadian rhythms, including through exposure to light at night, have been associated with an increased risk of tumorigenesis and a poorer prognosis in carcinomas (1,2). This suggests that the incidence of cancer may continue to increase, in association with the stresses of modern life. The risks of both breast and prostate cancer are high in industrialized societies (3). Continuous hormonal unbalances in shift workers might be caused by circadian disruption and may contribute to the rising risk of cancer. Endocrine target organs, such as the breast and prostate, are thought to be susceptible to psychosocial stresses, including circadian disruption (4). The immunological surveillance system has also been shown to be affected, and may thus not eliminate cancer stem cells (CSCs) effectively (5).

Circadian genes have been shown to function as oncogenes or tumor suppressors at systemic or cellular levels through their involvement in cell proliferation, cell cycle regulation, apoptosis and DNA damage signaling (6,7), indicating a direct effect of circadian effects on cancer cells. However, the kinds of molecular and systemic mechanisms involved in tumor growth under artificial illumination stress remain unknown, and the importance of artificial illumination in promoting tumor growth also needs to be established. We propose an indirect mechanism supporting the survival of potential CSCs and discuss a new concept of tumor niche formation induced by circadian disruption.

2. Cancer stem cell niche

Clinical tumors comprise a heterogeneous cell population including CSCs (8). The malignant phenotype depends not only on the characteristics of the cancer cell itself, but also on the tumor microenvironment. CSCs have to survive for a long time in the body to generate the highly tumorigenic cells responsible for the clinical manifestations of cancer. During this period, the niche helps to shelter CSCs from various insults such as the immune response, and from genotoxic stresses such as chemotherapy (9,10). This suggests that the niche may also play a protective role for CSCs, and may thus contribute to the risk of cancer.
3. Niche-driven tumor progression

Although CSCs appear relatively frequently, they are unable to survive in the healthy body without supportive tissues. As shown in Fig. 1, CSCs are killed by immune cells, or disappear under conditions of hypoxia and malnutrition. In general, CSCs only survive in the primary niche long enough to cause cancer when the microenvironment is supportive. The primary niche is composed of fibroblasts and the extracellular matrix (ECM) (11). Circadian disruption activates the fibroblasts, which produce the autocrine growth factor, WNT10a. These WNT10a-producing fibroblasts secret ECM, and may provide the beneficial conditions required to form an initial tumor niche or microenvironment for CSCs (Fig. 2). It is possible that this process may contribute to the maturation of CSCs, but not their differentiation. Psychosocial stress might activate resident fibroblasts in the body, which together with the ECM, may provide the niche required for CSCs in the preclinical phase. Mouse NIH3T3 cells overexpressing WNT10a can grow rapidly and form tumors in nude mice. As shown in Fig. 3, WNT10a-producing cells promote their own growth and secrete ECM. Interestingly, it has been reported that increased collagen density promotes mammary tumor initiation and progression (12).

It has been proposed that metastasis requires the existence of a metastatic niche to allow the invading cancer cells to survive, colonize and expand to form macrometastases (13). It has been reported that both cancer cells and cancer-associated fibroblasts (CAF) produce angiogenic factors, such as vascular endothelial growth factor and WNT10a, respectively, while it is well known that cancer cells produce reactive oxygen species (ROS) that induce these angiogenic factors (14). Rapid tumor growth with angiogenesis induces extracellular acidosis, which in turn leads to activation of metalloproteinases that destroy the structure of the tumor niche (15). Hypoxic glycolysis is activated and produces acid metabolites, and the subsequent decrease in intracellular pH has been shown to reduce DNA repair activity, resulting in the accumulation of spontaneous mutations following the malignant progression of cancer cells. Thus, solid tumors finally disrupt the niche through degradation of the ECM, and acquire the ability to invade tissues and to metastasize during the clinical phase. The tumor microenvironment is thus crucial for solid tumor development (16), and dysregulation of the pH and disruption of the tumor niche is closely involved in the cancer hallmarks of invasion and metastasis. These observations also suggest that circadian disruption in cancer patients might be related to poor prognosis.

4. Cancer-associated fibroblasts and tumor microenvironment

Fibroblasts are an abundant cell type in connective tissues. They produce ECM components and various cytokines, and contribute to the formation of a structural network through tissue remodeling (17). The tumor environment includes structural and cellular components. The cellular components are the so-called stromal cells, including both resident and circulating cells such as macrophages, inflammatory cells, endothelial cells, myofibroblasts and CAFs (18). Increasing evidence indicates that CAFs are a main player in the hallmarks of cancer such as angiogenesis, invasion, metastasis and inflammation, which are critical factors for the development of solid tumors (Fig. 4). However, the origin of CAFs remains unknown (19). Mesenchymal stem cells contribute to the formation of tumor-associated stroma containing cellular components such as myofibroblasts and fibro-
Furthermore, the detailed roles of CAFs are still unclear, and not all the cytokines produced by CAFs have yet been identified. It is possible that the properties of CAFs important during the early stage of tumor development differ from those involved in the late stage. CAFs may be activated by factors in the microenvironment, such as hypoxia (24), and by physiological conditions such as circadian rhythms. We demonstrated that disruption of circadian rhythms can promote tumor growth through WNT10A-dependent angio/stromagenesis, associated with increased levels of oxidative stress (25). Both endothelial cells and stromal cells may be activated by WNT10A signals from non-tumor cells such as CAFs. WNT signaling has been
classified into ‘canonical’ and ‘non-canonical’ pathways. In addition, β-catenin expression has been observed in endothelial cells in newly-formed tumor vessels, suggesting that WNT/β-catenin signaling plays a role in tumor angiogenesis. WNT signaling is also known to play an important role in cancer and stem cell biology (26), indicating that WNT10A might affect not only the tumor microenvironment, but also the CSCs themselves.

5. Tumor-driven niche development

The generation of CSCs and cancer progression involve a long-term and complicated series of processes. Accumulating evidence suggests that psychosocial stress can influence cancer cell growth via many processes. Cancer cells induce inflammation (27). Both tumor-associated macrophages and CAFs are critical for cancer progression (28). In addition, both cancer cells and activated stromal cells produce high levels of ROS and cytokines. ROS induce not only DNA damage following malignant transformation, but also CAF activation. These tumor-host interactions alter the local tumor environment and contribute to tumor growth. Activated CAFs produce ECM around cancer cells, and CAF-dependent ECM production may support the formation of a new niche allowing the development of local micrometastases around the primary tumor. Finally, the development of an angiogenic niche around the main tumor supports invasion and metastasis. Fig. 5 shows a typical case that supports this idea. Tumor-associated connective tissue is often observed in front of the main gastric tumor. Both azan staining and immunohistochemical studies showed that this region was rich in collagen and microvessels (Fig. 5B and C), and the vascular smooth muscle cells in the microvessels and the fibroblastic cells observed in the collagen deposits were positive for WNT10A (Fig. 5D). WNT10A may thus contribute to the formation of the angio/stromagenic niche. We also suggest that the tumor-associated connective tissue functions as an angio/stromagenic niche around the primary tumor. Bateman (29) also proposed that cancer cells modify the stroma of remote organs and create a premetastatic niche.

WNT10A mutations are associated with autosomal recessive ectodermal dysplasia (30). In addition, the expression of WNT signaling antagonists has been shown to be downregulated in fibroblasts in keloids, which are an aggressive type of wound-healing tissue (31). These previous reports indicate that WNT signaling is involved in both tissue repair and wound healing. An earlier hypothesis suggests that cancer results from uncontrolled wound-healing (32). This is supported by the observation that WNT10A expression was markedly increased in fibroblastic cells in the hyperplastic stroma of keloid tissue, suggesting its function as an angio/stromagenesis gene in tumor progression.

There are some limitations associated with experiments using mice. We cannot exclude the possibility that other physiological and/or hormonal factors, such as melatonin, may affect the growth of implanted cancer cells in mouse models. Melatonin suppression through exposure to artificial light at night leads to carcinogenesis of target endocrine organs (33). However, serum melatonin cannot be measured in mice because the pineal gland is too small. Thus although the subcutaneous injection of rapidly growing human cancer cells into nude mice provides a setting in which tumor growth can be assessed in a relatively short time span, an orthotopic model that more accurately reproduces the interactions between tumor cells and their microenvironment is required to confirm these results.

6. Drug resistance

The molecular mechanisms responsible for the cellular sensitivity to anticancer agents have been extensively studied in cancer cell lines (16). However, drug resistance is also influenced indirectly by the tumor microenvironment. Cisplatin resistance is affected by several factors that influence intracellular drug accumulation, including the levels of cellular thiols and DNA
Figure 5. Stomach cancer with remodeling of connective tissue as an angio/stromagenic niche. (A) Hematoxylin and eosin stained squamous cell carcinoma at the gastro-esophageal junction. Tumor penetrates the serosa (A-1). Carcinoma arranged in sheet-like nests or irregular cords confined to the serosa. Yellow dotted line indicates the front line of the cancer partitioned from the stroma (A-2). Scar-like stromal reaction, in which peripheral desmoplasia with proliferating spindle-shaped cells and numerous well-developed blood capillaries, is found throughout the front line of carcinoma progression (A-3). (B) Azan staining. In desmoplastic stromal areas, spindle-shaped cells and blood capillaries were counterstained red (B-1 and 2), as were nuclei, fibrin and epithelial hyaline bodies. Collagen fibers and mucus were counterstained blue, together with the basal lamina of the blood capillaries (B-3). (C) CD34 (red) - AE1/AE3 (brown) double staining method. Immunohistochemically, carcinoma cells were positive for AE1/AE3 (brown) and angioendothelial cells were positive for CD34 (red) (C-1 and 2). In stromal areas, spindle-shaped cells and the basal lamina of blood capillaries were stained red (C-3). (D) AE1/AE3 (red) - WNT10 (brown) double staining method. Immunohistochemically, carcinoma cells were positive for AE1/AE3 (red) and the stromal area was positive for WNT10 (brown) (D-1 and 2). In the stromal area, spindle-shaped cells and the basal lamina of blood capillaries were stained brown (D-3).
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Circadian disruption, tumor niche and development of solid tumor

Psychosocial stress → Activated stromal niche with activation of fibroblasts and ECM and cytokine production

WNT10A-dependent angio/stromagenesis

CSCs

Malignant transformation

Continued proliferation

Malignant progression

Evade apoptosis

Activated CAFs and TAMs → VEGF-dependent angiogenesis

Solid tumor development in clinical phase

Invasion

Angio/stromagenic niche as local tumor niche

Remote organ stroma

Premetastatic niche

Metastasis

Metastatic niche

Figure 6. Schematic summary.

repair activity (34). Our own studies showed that glutathione biosynthesis was upregulated by activating transcription factor 4 (ATF4), which is also regulated by the circadian transcription system (35,36). ATF4 expression was induced by oxidative stress through the Nrf2 transcription factor (37). The role of histone acetyltransferase (HAT) gene expression in the development of drug resistance has not been extensively studied, though it has been shown that HAT genes such as CLOCK and TIP60 are overexpressed, and are involved in glutathione biosynthesis and DNA repair (38), indicating that the system protecting against various stresses might be regulated periodically. These results indicate that similar mechanisms to those observed in CAFs may contribute to niche-dependent drug resistance.

The circadian transcription system thus drives the expression of genes that regulate the cell cycle, DNA repair, and thiol production, which are involved in drug sensitivity, indicating that the circadian rhythm may contribute to the efficacy of chemotherapy in cancer patients, with implications for side-effects and patient outcomes, including prognosis (39). New methods are required to understand the status of the circadian rhythm in an individual patient. WNT16B expression has recently been reported to be upregulated in fibroblasts by chemotherapy, and to promote epithelial-mesenchymal transition in neoplastic prostate epithelium through paracrine signaling (40). They also showed that WNT16B promoted the survival of cancer cells after chemotherapy, suggesting that the tumor environment functions as a stromal chemoresistant niche.

7. Conclusions

WNT10a is a key molecule in the development of the tumor niche. WNT10a-dependent activation of the tumor niche not only supports the emerging links between the circadian rhythm, oxidative stress and tumor progression at the molecular level, but also alerts us to the potentially adverse effects of artificial light. Further studies are needed to clarify whether WNT10A-Frizzled binding mediates cell proliferation in both endothelial cells and stromal cells. Examining WNT10A receptors and their associated signal transduction pathways may provide valuable insights into the role of circadian
rhythms in tumor progression. A greater understanding of the complexity of the tumor microenvironment and the role of the tumor niche will lead to further advances in cancer treatment.

Long-term disturbance and disruption of the circadian rhythm contributes to the development of cancer from the preclinical to clinical phases through the evolution of a highly specialized tumor niche (Fig. 6). As expected, WNT10A expression is controlled by a clock gene (41). These results suggest that researchers should consider the relevance of chronobiology based on the results of occupational science related to shift work. Improved understanding of the circadian rhythm will also allow the further development of chronotherapy in cancer patients.

References