EDITORIAL

# Cancer and rhythm

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In a recent editorial comment, Denise Duboule [1] emphasized that "animal development is, in fact, nothing but time." In this issue of *Cancer Causes and Control*, several papers will substantiate that not only developmental, but also neoplastic processes may be linked to what Duboule [1] and Halberg [2] referred to as 'chronomics': timing and rhythm. Moreover, there is reason to believe that timing and rhythm may have been underappreciated in current therapeutic settings [3]. The papers in this issue of

**Dedication** This special issue of CCC is dedicated to Gunther Hildebrandt (Marburg, Germany) and Franz Halberg (Minnesota, USA), two great pioneers of Chronobiology.

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*Cancer Causes and Control* are from leading researchers in the field of cancer and body rhythms. Each one addresses a specific aspect of the topic, and their work is cited below in this overview editorial.

Evidence from observational studies is growing [4] that the disturbance of body rhythms, in particular, circadian disruption e.g., shift work [5–8] or chronic jet-lag [9–12] contribute significantly to the development of breast cancer. Figure 1 compares the relative impact of rhythm disturbances to other exposures of significance in breast tumor development [13]. Reproductive risk factors such as parity and age at first birth, age at menarche, and age at menopause each confer a change in risk of roughly 20-30%. Only family history in a first degree confers a relative risk comparable in magnitude to that of female flight attendants. Unlike for other cancers for which primary risk factors have been identified (e.g., smoking and lung cancer risk), to date, no single environmental risk factor has been identified that can account for a major proportion of breast cancers, the incidence of which is still rising. Thus, there is a need for continued, vigorous search of breast cancer risk factors. Disruption of circadian rhythms, which can be caused by a wide variety of factors, may play an important role, not only for breast-but also other cancers, but results are still premature.

Disruption of clock gene function may increase cancer risk: In mice, clock genes stabilize the genome and help maintain important repair mechanisms such as the apoptosis of damaged cells [14]. Per-2 gene deprived mice lack a circadian rhythm [15] and have been shown to develop cancer rapidly and spontaneously. The difference between wild type and rhythm-deprived mice was found to be most striking after exposure to ionizing radiation. In suprachiasmatic nuclei (SCN) ablated mice, disruption of circadian rhythms was associated with accelerated growth of



Life Style Factors

**Fig. 1** Risk and beneficial factors influencing relative breast cancer risks have been extracted from the literature (see Electronic Supplementary Material). Possible chronobiological factors are marked in red, nutritional factors in green, all others in blue. The red line at 1.00 indicates the breast cancer risk of the control cohort documented in the

studies. Higher values e.g. in shift workers and flight attendants correspond to an increased risk. Some factors are known to decrease the risk of breast cancer, e.g. breastfeeding, the number of pregnancies and phytoestrogens contained in vegetables and miso soup

malignant tumors, suggesting that the host circadian clock may play an important role in endogenous control of tumor progression [16, 17].

A similar amplification of cancer promotion through disturbed body rhythms could also be inferred from Fig. 1. A recent meta-analysis of breast cancer risk in flight attendants and shift working women has estimated that the overall relative risk is about 1.5 [18]. There is variability among these studies in what constitutes shift work and of whom the comparison group is comprised, so more must be learned about the impact of occupational circadian rhythm disturbance and risk. In particular, there may be possible interactions with other exposures such as cosmic radiation in the flight attendants.

In animal studies, chronic jet lag suppressed the rhythm in clock gene expression in normal liver and in tumors [19, 20]. Prolonged exposure to constant light promoted growth of transplantable tumors in rats [21, 22] and mice [23], whereas melatonin and rest-activity rhythms were suppressed.

Cancer growth on the other hand may also disturb body rhythms: for example, rhythms of cancer cells degenerate from those of their host organism [24–26]. Chen et al. [27]

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showed deregulation of clock genes in tumor cells compared to normal cells in breast cancers.

It has also been argued that rhythm disturbance is not a carcinogenic effect in itself [28]: Light at night induces a decrease of plasma melatonin levels [4, 29] via specialized retinal photoreceptors that were only recently discovered [30–32]. Since both blind people [33–36] as well as workers in photographic laboratories [37] were reported to have lower breast cancer incidence rates than the general population, melatonin has been implicated. However, among cancer patients, not only changes in melatonin levels, but also changes in their circadian rhythms were found, in some cases independent of their melatonin levels [38, 39]. Untreated breast and prostate cancer patients, on the other hand, were found to show tumor-size dependent depressions of the circadian amplitude of circulating melatonin, which was accompanied by parallel neuroendocrine disturbances involving particularly prolactin and TSH [40]. Also colorectal cancer patients [39, 41] as well as patients with lung and stomach cancer [42] had lower levels of melatonin in plasma resp. 6-sulfatoxymelatonin in urine than healthy control subjects, suggesting a possible link between low melatonin levels and the enhanced

development of cancer in humans. Overall, a strong interaction between rhythm disturbances and melatonin levels appear most likely: a recent study demonstrates the interplay between melatonin levels, rhythm disturbances and tumor growth: blood from human donors collected during a normal night was able to suppress proliferative activity in rats xenografted with human breast cancer [43]. Blood collected during the day or after a white light exposure during the night did not. In sum, only subjects that had not been exposed to light at night produced enough melatonin to suppress proliferative cell growth in this animal model. So, melatonin may also be beneficial in cancer treatment if administered at chronobiologically determined optimal times of the day [44]: according to this study, evening administration stabilized quality of life and prevented cancer-related cachexia. Significantly less patients showed a progressive disease in the melatonin group. Further evidence [45] supports the notion that melatonin is beneficial in human cancer treatment regardless of the timing of its administration. Other studies, finally, suggest melatonin as a possible remedy against jet-lag: Manfredini et al. [46] and Cardinali et al. [47] administered melatonin to athletes after transmeridian flights. Cardinali's athletes displayed fewer jet lag symptoms and coped better than those of Manfredini. The former group received an additional 'Zeitgeber' training, which appears to increase melatonin's efficacy. With growing data, we therefore argue that circulating melatonin levels may be only one aspect of the cancer-protective effects of stable biological rhythms. While rhythm disturbances are likely to aggravate tumor growth, the use of well chosen "Zeitgebers" could be of additional value to prevent tumors or to support tumor therapy.

"Zeitgebers", as defined by Aschoff in the 1960s [48], are cues, which help organisms to establish and stabilize their biologic rhythms. Such cues may be parameters like morning light, which indicates the beginning of the day. Temperature changes and meals, as well as the administration of drugs at a certain phase of the circadian cycle are other examples of such "Zeitgebers".

"Zeitgebers" have also been used as therapeutic elements is chrono-chemotherapy [49–51]. Chrono-chemotherapy implies drug administration to cancer patients at what is considered 'the right time of the day', following the 'Kairos' principle.<sup>1</sup> However, although such timing has been shown to successfully reduce the adverse effects of toxic agents [52], it has so far only rarely resulted in an increase of survival rates when compared to conventional, constant-rate infusion [53]. Thus, and because chemotherapy in itself reduces the amplitude of body rhythms [54– 56], it might not work outside the framework of an accompanying "rhythm therapy", which is yet to be established in oncology.

By using several "Zeitgebers" in the right order, it is possible to develop novel rhythm therapies. What are the basic elements of such a rhythm therapy? Klages [57] distinguishes between 'tact' and 'rhythm': 'tact' refers to identical things happening during identical periods, whereas 'rhythm' refers to similar things happening during similar periods. Today, it has been established that biological rhythms are not rigid but flexible; thus, a rhythm therapy should also take flexibility into account. For example, the circadian phase of a patient could be determined prior to intervention and be considered when designing a patient's treatment regimen. Determining whether the patient is an "owl" or a "lark" (evening/ morning type [2, 50, 58] could thus lead to a better adaptation of the therapy.

To date, "Zeitgeber" training has been applied in a variety of diseases. For example, light at the right time of the day seems to substantially improve the condition of patients suffering from seasonal affective disorder [59]; modification of the sleep pattern intended to reset the circadian clock has been successfully used to improve psychiatric depression [60]; a rhythm therapy has been for patients following developed stroke (http:// www.viewzone.com/rhythm therapy.html). We too, have recently shown that both the number of accidents as well as quality of sleep in stressed construction workers were significantly improved by applying a rhythm therapy [61]. This involved not only the introduction of additional "Zeitgebers" (e.g. working breaks in a pattern adopted from the human basic rest and activity cycle), but also the enhancement of body rhythms by resonance phenomena and by coordinating different body rhythms through art therapy [62, 63]. We further observed that excitation of the biological system with ultradian short period rhythms increases the amplitude of its long period rhythms, thereby enlarging the circadian cycle of heart rate and heart rate variability [61], both of which are connected to sleep quality.

In sum, the elements of an optimal rhythm therapy should incorporate:

- Use of several "Zeitgebers" to complement each other instead of a simple Zeitgeber at a time;
- the use of ultradian short period "Zeitgebers" (to enhance long period circadian rhythms by excitation, coupling and resonance);
- measures to coordinate various body rhythms;
- flexible and individualized rhythms instead of metronomic periods
- Kairos instead of Chronos (right timing instead of arbitrarily set timing)

<sup>&</sup>lt;sup>1</sup> Kairos ( $\kappa \alpha i \rho o \zeta$ ) is the Greek word for the right time in contrast to Chronos ( $\chi \rho o v o \zeta$ ), the term for time in general.

We suggest that maintenance of a strong circadian rhythm in everyday life [64] and the use of rhythm therapy when disease occurs [65], may not only help to substantially reduce breast cancer incidence and mortality in industrialized countries, but, more generally, will also support the introduction of new avenues in the prevention and treatment of cancer [3].

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## References

- 1. Duboule D (2003) Time for chronomics? Science 301:277
- Halberg F, Cornelissen G, Wang Z, et al. (2003) Chronomics: circadian and circaseptan timing of radiotherapy, drugs, calories, perhaps nutriceuticals and beyond. 101
- Moser M, Frühwirth M, Penter R, Winker R (2006) Why life oscillates? – from a topographical towards a functional chronobiology. Cancer Causes Control 17:591–599
- 4. Stevens RG, Rea MS (2001) Light in the built environment: potential role of circadian disruption in endocrine disruption and breast cancer. Cancer Causes Control 12:279–287
- Schernhammer ES, Laden F, Speizer FE, et al. (2001) Rotating night shifts and risk of breast cancer in women participating in the nurses' health study. J Natl Cancer Inst 93:1563–1568
- Tynes T, Hannevik M, Andersen A, Vistnes AI, Haldorsen T (1996) Incidence of breast cancer in Norwegian female radio and telegraph operators. Cancer Causes Control 7:197–204
- Davis S, Mirick DK, Stevens RG (2001) Night shift work, light at night, and risk of breast cancer. J Natl Cancer Inst 93:1557–1562
- 8. Hansen J (2001) Increased breast cancer risk among women who work predominantly at night. Epidemiology 12:74–77
- Rafnsson V, Tulinius H, Jonasson JG, Hrafnkelsson J (2001) Risk of breast cancer in female flight attendants: a population-based study (Iceland). Cancer Causes Control 12:95–101
- Pukkala E, Auvinen A, Wahlberg G (1995) Incidence of cancer among Finnish airline cabin attendants, 1967–92. BMJ 311:649– 652
- Haldorsen T, Reitan JB, Tveten U (2000) Cancer incidence among Norwegian airline pilots. Scand J Work Environ Health 26:106–111
- Blettner M, Zeeb H, Langner I, Hammer GP, Schafft T (2002) Mortality from cancer and other causes among airline cabin attendants in Germany, 1960–1997. Am J Epidemiol 156:556–565
- Key TJ, Verkasalo PK, Banks E (2001) Epidemiology of breast cancer. Lancet Oncol 2:133–140
- Fu L, Lee CC (2003) The circadian clock: pacemaker and tumour suppressor. Nat Rev Cancer 3:350–361
- Fu L, Pelicano H, Liu J, Huang P, Lee C (2002) The circadian gene Period2 plays an important role in tumor suppression and DNA damage response in vivo. Cell 111:41–50

- Filipski E, King VM, Li X, et al. (2002) Host circadian clock as a control point in tumor progression. J Natl Cancer Inst 94:690–697
- Filipski E, King VM, Li X, et al. (2003) Disruption of circadian coordination accelerates malignant growth in mice. Pathol Biol (Paris) 51:216–219
- Megdal SP, Kroenke CH, Laden F, Pukkala E, Schernhammer ES (2005) Night work and breast cancer risk: a systematic review and meta-analysis. Eur J Cancer 41:2023–2032
- Filipski E, Delaunay F, King VM, et al. (2004) Effects of chronic jet lag on tumor progression in mice. Cancer Res 64:7879–7885
- van den Heiligenberg S, Depres-Brummer P, Barbason H, Claustrat B, Reynes M, Levi F (1999) The tumor promoting effect of constant light exposure on diethylnitrosamine-induced hepatocarcinogenesis in rats. Life Sci 64:2523–2534
- Dauchy RT, Blask DE, Sauer LA, Brainard GC, Krause JA (1999) Dim light during darkness stimulates tumor progression by enhancing tumor fatty acid uptake and metabolism. Cancer Lett 144:131–136
- 22. Blask DE, Dauchy RT, Sauer LA, Krause JA, Brainard GC (2003) Growth and fatty acid metabolism of human breast cancer (MCF-7) xenografts in nude rats: impact of constant light-induced nocturnal melatonin suppression. Breast Cancer Res Treat 79:313–320
- 23. Bartsch C, Bartsch H (2006) The anti-tumor activity of pineal melatonin and cancer enhancing life styles in industrialized societies. Cancer Causes Control (this volume)
- Keith LG, Oleszczuk JJ, Laguens M (2001) Circadian rhythm chaos: a new breast cancer marker. Int J Fertil Womens Med 46:238–247
- 25. Mormont MC, Langouet AM, Claustrat B, et al. (2002) Marker rhythms of circadian system function: a study of patients with metastatic colorectal cancer and good performance status. Chronobiol Int 19:141–155
- 26. Simpson HW, Pauson AW, McArdle C, Griffiths K, Turkes A (1990) Loss of the progesterone-associated metabolic rhythm in cancer-associated breast tissue. Prog Clin Biol Res 341A:47–53
- Chen ST, Choo KB, Hou MF, Yeh KT, Kuo SJ, Chang JG (2005) Deregulated expression of the PER1, PER2 and PER3 genes in breast cancers. Carcinogenesis 26:1241–1246
- Gauger MA, Sancar A (2005) Cryptochrome, circadian cycle, cell cycle checkpoints, and cancer. Cancer Res 65:6828–6834
- Czeisler CA, Shanahan TL, Klerman EB, et al. (1995) Suppression of melatonin secretion in some blind patients by exposure to bright light. N Engl J Med 332:6–11
- Brainard GC, Hanifin JP, Greeson JM, et al. (2001) Action spectrum for melatonin regulation in humans: evidence for a novel circadian photoreceptor. J Neurosci 21:6405–6412
- Berson DM, Dunn FA, Takao M (2002) Phototransduction by retinal ganglion cells that set the circadian clock. Science 295:1070–1073
- Brainard GC, Hanifin JP, Rollag MD, et al. (2001) Human melatonin regulation is not mediated by the three cone photopic visual system. J Clin Endocrinol Metab 86:433–436
- Hahn RA (1991) Profound bilateral blindness and the incidence of breast cancer. Epidemiology 2:208–210
- Feychting M, Osterlund B, Ahlbom A (1998) Reduced cancer incidence among the blind. Epidemiology 9:490–494
- Verkasalo PK, Pukkala E, Stevens RG, Ojamo M, Rudanko SL (1999) Inverse association between breast cancer incidence and degree of visual impairment in Finland. Br J Cancer 80:1459–1460
- Kliukiene J, Tynes T, Andersen A (2001) Risk of breast cancer among Norwegian women with visual impairment. Br J Cancer 84:397–399
- Olsen JH, Jensen OM (1987) Occupation and risk of cancer in Denmark. An analysis of 93,810 cancer cases, 1970–1979. Scand J Work Environ Health 13(Suppl 1):1–91

- Kos-Kudla B, Ostrowska Z, Kozlowski A, et al. (2002) Circadian rhythm of melatonin in patients with colorectal carcinoma. Neuroendocrinol Lett 23:239–242
- Vician M, Zeman M, Herichova I, Jurani M, Blazicek P, Matis P (1999) Melatonin content in plasma and large intestine of patients with colorectal carcinoma before and after surgery. J Pineal Res 27:164–169
- 40. Bartsch C, Bartsch H, Fluchter SH, Mecke D, Lippert TH (1994) Diminished pineal function coincides with disturbed circadian endocrine rhythmicity in untreated primary cancer patients. Consequence of premature aging or of tumor growth? Ann N Y Acad Sci 719:502–525
- Khoory R, Stemme D (1988) Plasma melatonin levels in patients suffering from colorectal carcinoma. J Pineal Res 5:251–258
- 42. Kvetnaia T, Kvetnoy I, Bartsch H, Bartsch C, Mecke D (2001) Melatonin in patients with cancer of extra-reproductive location. In: Bartsch C, Bartsch H, Blask D, Cardinali D, Hrushesky W, Mecke D (eds) The pineal gland and cancer: neuroimmunoendocrine mechanisms in malignancy, Springer Verlag, Berlin, pp 177–196
- 43. Blask DE, Brainard GC, Dauchy RT, et al. (2005) Melatonindepleted blood from premenopausal women exposed to light at night stimulates growth of human breast cancer xenografts in nude rats. Cancer Res 65:11174–11184
- 44. Lissoni P, Chilelli M, Villa S, Cerizza L, Tancini G (2003) Five years survival in metastatic non-small cell lung cancer patients treated with chemotherapy alone or chemotherapy and melatonin: a randomized trial. J Pineal Res 35:12–15
- Vijayalaxmi, Thomas CR, Jr., Reiter RJ, Herman TS (2002) Melatonin: from basic research to cancer treatment clinics. J Clin Oncol 20:2575–2601
- 46. Manfredini R, Manfredini F, Conconi F (2000) Standard melatonin intake and circadian rhythms of elite athletes after a transmeridian flight. J Int Med Res 28:182–186
- Cardinali DP, Brusco LI, Lloret SP, Furio AM (2002) Melatonin in sleep disorders and jet-lag. Neuroendocrinol Lett 23(Suppl 1):9–13
- Aschoff J (1965) Circadian rhythms in man. Science 148:1427– 1432
- 49. Levi F, Zidani R, Misset JL (1997) Randomised multicentre trial of chronotherapy with oxaliplatin, fluorouracil, and folinic acid in metastatic colorectal cancer. International Organization for Cancer Chronotherapy. Lancet 350:681–686
- Mormont MC, Levi F (1997) Circadian-system alterations during cancer processes: a review. Int J Cancer 70:241–247

- Hrushesky (2003) Circadian cancer chronotherapy: The Necessity for Complementary Preclinical Investigation. webside 3
- Levi F (2002) From circadian rhythms to cancer chronotherapeutics. Chronobiol Int 19:1–19
- 53. Levi FA, Zidani R, Vannetzel JM, et al. (1994) Chronomodulated versus fixed-infusion-rate delivery of ambulatory chemotherapy with oxaliplatin, fluorouracil, and folinic acid (leucovorin) in patients with colorectal cancer metastases: a randomized multiinstitutional trial. J Natl Cancer Inst 86:1608–1617
- Ekholm EM, Salminen EK, Huikuri HV, et al. (2000) Impairment of heart rate variability during paclitaxel therapy. Cancer 88:2149–2153
- 55. Ohdo S, Koyanagi S, Suyama H, Higuchi S, Aramaki H (2001) Changing the dosing schedule minimizes the disruptive effects of interferon on clock function. Nat Med 7:356–360
- Li XM, Vincenti M, Levi F (2002) Pharmacological effects of vinorelbine on body temperature and locomotor activity circadian rhythms in mice. Chronobiol Int 19:43–55
- 57. Klages L (1923) On the essence of Rhythm (in German)
- Hildebrandt G, Moser M, Lehofer M (1998) Chronobiology und chronomedicine—biologic rhythms medical consequences (in German). Hippokrates, Stuttgart
- Terman JS, Terman M (1999) Photopic and scotopic light detection in patients with seasonal affective disorder and control subjects. Biol Psychiatry 46:1642–1648
- 60. Riemann D, Konig A, Hohagen F, et al. (1999) How to preserve the antidepressive effect of sleep deprivation: A comparison of sleep phase advance and sleep phase delay. Eur Arch Psychiat Clin Neurosci 249:231–237
- Moser M, Frühwirth M, Lackner H, et al. (2000) Baufit—stress on the building site—made visible by the heartbeat. AUVA-Report (Vienna) 38:55–70
- Bettermann H, von Bonin D, Fruhwirth M, Cysarz D, Moser M (2002) Effects of speech therapy with poetry on heart rate rhythmicity and cardiorespiratory coordination. Int J Cardiol 84:77–88
- Cysarz D, von Bonin D, Lackner H, Heusser P, Moser M, Bettermann H (2004) Oscillations of heart rate and respiration synchronize during poetry recitation. Am J Physiol Heart Circ Physiol 287:H579–H587
- Stevens RG (2005) Circadian disruption and breast cancer: from melatonin to clock genes. Epidemiology 16:254–258
- Hrushesky WJ (2001) Tumor chronobiology. J Control Release 74:27–30