Parathyroid hormone resets the cartilage circadian clock of the organ-cultured murine femur

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Background and purpose — The circadian clock governs endogenous day-night variations. In bone, the metabolism and growth show diurnal rhythms. The circadian clock is based on a transcription-translation feedback loop composed of clock genes including Period2 (Per2), which encodes the protein period circadian protein homolog 2. Because plasma parathyroid hormone (PTH) levels show diurnal variation, we hypothesized that PTH could carry the time information to bone and cartilage. In this study, we analyzed the effect of PTH on the circadian clock of the femur.

Patients and methods — Per2::Luciferase (Per2::Luc) knock-in mice were used and their femurs were organ-cultured. The bioluminescence was measured using photomultiplier tube-based real-time bioluminescence monitoring equipment or real-time bioluminescence microscopic imaging devices. PTH or its vehicle was administered and the phase shifts were calculated. Immunohistochemistry was performed to detect PTH type 1 receptor (PTH1R) expression.

Results — Real-time bioluminescence monitoring revealed that PTH reset the circadian rhythm of the Per2::Luc activity in the femurs in an administration time-dependent and dose-dependent manner. The bioluminescence was measured using photomultiplier tube-based real-time bioluminescence monitoring equipment or real-time bioluminescence microscopic imaging devices. PTH or its vehicle was administered and the phase shifts were calculated. Immunohistochemistry was performed to detect PTH type 1 receptor (PTH1R) expression.

Interpretation — In clinical practice, teriparatide (PTH (1-34)) treatment is widely used for osteoporosis. We found that PTH administration regulated the femoral circadian clock oscillation, particularly in the cartilage. Regulation of the local circadian clock by PTH may lead to a more effective treatment for not only osteoporosis but also endochondral ossification in bone growth and fracture repair.
time information may be carried to bones via a leptin-parasymathetic neuron-bone pathway. The circadian rhythm of bioluminescence in cartilage can be reset with forskolin or synthesized glucocorticoid (dexamethasone) (Gossan et al. 2013, Okubo et al. 2013). Interestingly, plasma PTH levels have been reported to show day-night variations under normal light-dark conditions (Jubiz et al. 1972) and also under “constant routine” conditions, where no external time information is available (el-Hajj Fuleihan et al. 1997). Hinoi et al. (2006) reported that PTH induces Per1 and Per2 expression in organ-cultured fetal metatarsal. We therefore hypothesized that PTH could carry time information to bone and cartilage. In this study, we investigated the effect of PTH on the femoral circadian clock.

Material and methods

Animals and femur sampling

Period2::Luciferase (Per2::Luc) knock-in mice were originally developed by Dr Joseph Takahashi’s group (Yoo et al. 2004). Mice were maintained by intercrossing with their littermates and backcrossing to C57BL/6 mice at our facility. They were kept under 12-h light-dark conditions, and food and water were available ad libitum. For tissue sampling, the mice were deeply anesthetized with isoflurane and killed. We used both male and female mice (4–15 weeks old), as no sex and age differences have been observed in the circadian clock period or phase (Gossan et al. 2013, Okubo et al. 2013). After sterilization with 70% ethanol, femurs were collected and stored in ice-cold PBS. Muscle tissue and tendons were removed.

Femoral organ culture

Femoral organ culture was performed as previously reported (Okubo et al. 2013). In brief, collected femurs were placed in 35-mm cell culture dishes. The culture medium contained phenol red-free D-MEM (Nacalai Tesque, Kyoto, Japan), 1× Glutamax (Life Technologies, Carlsbad, CA), 1 × B-27 supplement (Life Technologies), 10 mM HEPES, 200 μM beetle luciferin (Promega KK, Tokyo, Japan), 100 units/mL penicillin and 100 μg/mL streptomycin (Nacalai Tesque).

Real-time bioluminescence monitoring and PTH stimulation

Femurs were synchronized with forskolin (10 μM) for 1 h and then released into fresh culture medium as previously described (Gibbs et al. 2009, Okubo et al. 2013) The tightly sealed culture dishes were placed under the photomultiplier tube-based real-time bioluminescence monitoring equipment, kept at 35°C, and monitored every 20 min. Human PTH (1-34) (Peptide Institute, Inc., Ibaraki, Japan) was dissolved in distilled water. PTH (1-34) or its vehicle was added and the femurs were set under the bioluminescence counting device again. To study the effect of PTH on the femoral circadian clock, we administered PTH after the second peak (41 h after synchronization) or the trough of Per2::Luc activity (53 h after synchronization).

Microscopic observation

Microscopic bioluminescence observation was performed using a high-sensitivity CCD camera-based microscopic image analyzer and bioluminescence macrosopy (Olympus, Tokyo, Japan). After synchronization, femurs were placed under the CCD camera-based device. After 40 h of observation, 10^{-7} M PTH (1-34) or its vehicle was added and the femurs were placed under the device again.

Immunohistochemistry

Immunohistochemistry was performed by New Histo, Science Laboratory Co., Ltd. (Tokyo, Japan). Briefly, the femurs were fixed with 4% paraformaldehyde for 8 h, which was followed by treatment with 70% ethanol for 3–4 days. The femurs were demineralized with 10% EDTA for 2–4 weeks, and then embedded in paraffin and sliced into 3- or 4-μm sections. After deparaffinization, the samples were incubated in 3% H2O2 for 5 min, washed with 0.01 M PBS, and incubated for 50 min at room temperature in rabbit polyclonal anti-parathyroid hormone receptor 1 antibody (Acris Antibodies GmbH, Herford, Germany) diluted 1:150 in 0.01 M PBS. In the negative controls, anti-rabbit IgG was used instead of the primary antibody. For the secondary antibody, the samples were incubated in Histostar™ mouse (Rb) (Immunostaining Reagent, Anti-Rabbit) (Medical & Biological Laboratories Co., Ltd., Nagoya, Japan) for 5 min and then washed with 0.01 M PBS, followed by visualization of any antibody binding with DAB-peroxidase. Finally, the samples were counterstained with hematoxylin.

Data analysis

For detection of phase shift, the data were smoothened by a 24-h moving average and the damping effect was corrected by using a 24-h standard deviation (Izumo et al. 2006). The difference in the periods between pre- and post-stimulation was calculated. The circadian period was defined as the peak-to-peak interval. Statistical analysis was performed with 1-way ANOVA followed by Dunnett’s test.

Ethics

Animal experiments were performed with approval from the Experimental Animals Committee, Kyoto Prefectural University of Medicine (date of issue: April 1, 2012; registration number: M24-240).

Results

When a low concentration of PTH (10^{-11} M) was administered
To elucidate the spatial distribution of bioluminescence, we studied the effect of PTH on the circadian clock in femurs using the CCD camera-based device (Figure 2). We administered $10^{-7}$ M PTH 41 h after synchronization. This is because the largest phase shift was evoked under this condition (Figure 1). Before administration of PTH, strong Per2::Luc signals were observed in cartilaginous tissues such as the epiphysial cartilage of the growth plate and articular cartilage, whereas Per2::Luc signals in the ossified bone were extremely weak, as we have reported previously (Okubo et al. 2013) (Figure 2A). No obvious change was observed from treatment with vehicle, and the next peak came 16 h after administration (Figure 2B–D). With PTH treatment, signal in the growth plate gradually increased and peaked 8 h after PTH administration, and then continued circadian oscillations (Figure 2B–D). Similar PTH-induced phase shifts were observed in the femoral head, the greater and lesser trochanter, and the distal end (data not shown). We also confirmed that PTH1R was expressed in the growth plate cartilage, by immunohistochemistry (Figure 2E).

Discussion

We found that PTH reset the circadian rhythm of the Per2::Luc activity in the femurs in both a time-dependent and a dose-dependent manner. By CCD camera-based imaging analysis, the phase shift of Per2::Luc activity was observed in the growth plate and articular cartilage. These results suggest that PTH induces the phase shift of the circadian clock of cartilage. In the suprachiasmatic nucleus, a light pulse induces Per1 and Per2 mRNA expression and resets the circadian clock in both a time-dependent and an intensity-dependent manner (Shigeyoshi et al. 1997). Field et al. (2000) reported that light induced both Per mRNA expression and Per protein expression in the suprachiasmatic nucleus, although protein peaks came a few hours later. Similarly, PTH administration has been reported to induce Per1 and Per2 mRNA expression in chondrogenic cells or tissues. Hinoi et al. (2006) observed that PTH administered at $10^{-7}$ M induced both Per1 and Per2 promoter activity in chondrogenic ATDC5 cells via the cAMP-PKA-CRE pathway. They also showed that both Per1 mRNA expression and Per2 mRNA expression were induced 1–2 hours after administration of $10^{-7}$ M PTH to organ-cultured fetal metatarsals. Taken together with our data, this indicates that PTH may reset the circadian clock of cartilage through alteration in levels of expression of Per mRNA and protein.

Not only PTH, but also PTHrP can exert physiological effects through the activation of PTH1R (Kopic and Geibel 2013). In the growth plate, where we confirmed PTH1R expression, hormones and paracines including PTHrP tightly control endochondral ossification (Lanske et al. 1996, Kronenberg 2003). Interestingly, the bone length of both PTHrP knock-out mice and clock gene BMal1 knock-out mice is shorter (Schipani et al. 1997, Bunger et al. 2005). PTHrP...
synthesis is mainly regulated by Indian hedgehog (IHH) in the growth plate (Chung et al. 2001). Takarada et al. (2012) reported that the expression of Ihh clearly shows the circadian rhythm. Our findings raise the possibility that PTHrP carries time information to cartilage. Despite the increasing number of studies on the cartilage circadian clock, the local clock function still remains unclear. Importantly, bone length of the chondrocyte-specific Bmal1-deficient mouse is shorter than that of its wild-type equivalent (Takarada et al. 2012). Since the secretion of the systemic hormones that control bone growth presumably has normal diurnal rhythms in such circumstances, the local circadian clock must have a role as a regulating mechanism in bone growth. Our results suggest that the local circadian clock is regulated by PTH. Thus, the regulation of the local circadian clock by PTH may lead to a treatment for diseases of bone growth.

In clinical practice, teriparatide is widely used for treatment of osteoporosis, but not in young patients with bone growth disease. Endochondral ossification occurs not only in the growth plate but also in the fracture site. Teriparatide treatment has been found to accelerate fracture healing, due to an increase in chondrocyte recruitment and in the rate of differentiation, in a mouse fracture model (Kakar et al. 2007). Our results suggest that fracture repair may be a target of chronotherapy using teriparatide, and further studies on the circadian clock of cartilage are therefore needed to determine the most effective timing of teriparatide administration.

In summary, we have demonstrated that PTH administration induced phase shifts of oscillation of the circadian clock in the growth plate and articular cartilage of a bone organ culture system. These results suggest that PTH regulates the circadian clock of cartilage.
KY, NO, YM, and T Kubo conceived and designed the experiments. NO, YM, TKun, TH, and HI performed the experiments. NO, HF, YM, TKun, TH, and YU analyzed the data. NO, HF, YM, MA, TKub, and KY wrote the paper. All the authors approved the final manuscript.

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