Visualizing and Quantifying Intracellular Behavior and Abundance of the Core Circadian Clock Protein PERIOD2

**Graphical Abstract**

**Highlights**

- Reporter mouse is used for real-time fluorescent imaging of mammalian clock protein PER2
- In contrast to *Drosophila*, localization of PER2 is not subject to circadian gating
- Circadian abundance, mobility, and intracellular dynamics of native PER2 are quantified
- Casein kinase 1 controls nucleocytoplasmic mobility of PER2 alongside circadian period

**In Brief**

Smyllie et al. use a fluorescent reporter mouse to image the circadian dynamics of PER2, a key component of the circadian clock in the suprachiasmatic nucleus and fibroblasts. They reveal marked divergence of the mechanisms of mouse and fly clock cells, and they provide quantitative data to support reappraisal of current models of the mammalian clock.

**Authors**

Nicola J. Smyllie, Violetta Pilorz, James Boyd, ..., Michael R.H. White, Michael H. Hastings, Andrew S.I. Loudon

**Correspondence**

mha@mrc-lmb.cam.ac.uk (M.H.H.), andrew.loudon@manchester.ac.uk (A.S.I.L.)

**Smyllie et al., 2016, Current Biology 26, 1880–1886**

July 25, 2016 © 2016 The Authors. Published by Elsevier Ltd. http://dx.doi.org/10.1016/j.cub.2016.05.018
Visualizing and Quantifying Intracellular Behavior and Abundance of the Core Circadian Clock Protein PERIOD2

Nicola J. Smyllie,1,3 Violetta Pilorz,2,3,4 James Boyd,2 Qing-Jun Meng,2 Ben Saer,2 Johanna E. Chesham,1 Elizabeth S. Maywood,1 Toke P. Krosgager,1 David G. Spiller,2 Raymond Boot-Handford,2 Michael R.H. White,2 Michael H. Hastings,1,* and Andrew S.I. Loudon1,2,*

1Neurobiology Division, Medical Research Council (MRC) Laboratory of Molecular Biology (LMB), Francis Crick Avenue, Cambridge CB2 0QH, UK
2Faculty of Life Sciences, University of Manchester, Oxford Road, Manchester M13 9PT, UK
3Co-first author
4Present address: Sleep and Circadian Neuroscience Institute, Nuffield Department of Clinical Neurosciences, University of Oxford, South Parks Road, Oxford, OX1 3RE, UK
*Correspondence: mha@mrc-lmb.cam.ac.uk (M.H.H.), andrew.loudon@manchester.ac.uk (A.S.I.L.)

http://dx.doi.org/10.1016/j.cub.2016.05.018

Summary

Transcriptional-translational feedback loops (TTFLs) are a conserved molecular motif of circadian clocks. The principal clock in mammals is the suprachiasmatic nucleus (SCN) of the hypothalamus. In SCN neurons, auto-regulatory feedback on core clock genes Period (Per) and Cryptochrome (Cry) following nuclear entry of their protein products is the basis of circadian oscillation [1, 2]. In Drosophila clock neurons, the movement of dPer into the nucleus is subject to a circadian gate that generates a delay in the TTFL, and this delay is thought to be critical for oscillation [3, 4]. Analysis of the Drosophila clock has strongly influenced models of the mammalian clock, and such models typically infer complex spatiotemporal, intracellular behaviors of mammalian clock proteins. There are, however, no direct measures of the intracellular behavior of endogenous circadian proteins to support this: dynamic analyses have been limited and often have no circadian dimension [5–7]. We therefore generated a knockin mouse expressing a fluorescent fusion of native PER2 protein (PER2::VENUS) for live imaging. PER2::VENUS recapitulates the circadian functions of wild-type PER2 and, importantly, the behavior of PER2::VENUS runs counter to the Drosophila model: it does not exhibit circadian gating of nuclear entry. Using fluorescent imaging of PER2::VENUS, we acquired the first measures of mobility, molecular concentration, and localization of an endogenous circadian protein in individual mammalian cells, and we showed how the mobility and nuclear translocation of PER2 are regulated by casein kinase. These results provide new qualitative and quantitative insights into the cellular mechanism of the mammalian circadian clock.

Results

Generation and Validation of PERIOD2::VENUS Mouse

We used homologous recombination to knock in a fluorescent tag at the Per2 locus, an equivalent strategy to that used for the PER2::LUC mouse, which is known to exhibit wild-type (WT) PER2 behavior [8]. Venus was fused to exons 19–23 of mPer2 (Figure S1A). The presence of PER2::VENUS protein expression was confirmed by fluorescence microscopy in the brain and in lung fibroblasts (Figures 1A and 1B). As well as strong fluorescence in the suprachiasmatic nucleus (SCN), limited expression was observed in the pituitary cortex, thalamus, and hippocampus (Figure S1B). Importantly, the spatial distribution of PER2::VENUS co-localized completely with PER2 immunoreactivity (-ir) in Per2WT/Venus SCN (Figures S1C–S1E).

To test for normal circadian function in Per2VENUS/Venus animals, we first assessed wheel-running behavior. They entrained effectively to a 12-hr light/12-hr dark schedule (12:12 LD), and they exhibited consolidated circadian activity patterns of wheel-running when placed in constant conditions (Figures 1C, S1F, and S1G). There were no significant differences between WT and Per2VENUS mice in the distribution, structure, or robustness (measured by non-parametric circadian rhythm analysis) of circadian behavior. After crossing with Per1-luc reporter mice, Per2VENUS/Venus SCN organotypic slices expressed robust, high-amplitude circadian bioluminescence rhythms (Figures 1D and S1H). The circadian periods of behavioral and SCN rhythms were not significantly different between WT and Per2VENUS mice (Figures 1E and 1F). Thus, PER2::VENUS did not compromise molecular pacemaking in the SCN or effective circadian control over behavior. To confirm that Per2VENUS did not encode a loss-of-function mutation, Per2VENUS mice were crossed to Per1−/− mice. In the absence of PER1, WT PER2 is a necessary and sufficient component of the circadian pacemaker [9]. Per2VENUS/Per1−/− mice exhibited robust and sustained wheel-running and SCN bioluminescence rhythms (Figures S1I and S1J), with comparable periods to Per2WT/WT, Per1−/− mice (Figure S1K). Thus, Per2VENUS encodes a functional allele of...
PER2\textsuperscript{Venus} mice were then crossed with CK1\textsubscript{ε}Tau mutants to test whether PER2::VENUS can interact with CK1\textsubscript{ε}, a key modulator of PER2 stability and circadian period \cite{10}. In WT animals, the CK1\textsubscript{ε}Tau/Tau mutation shortened period from \(~24\) hr (Figures S1I, S1J, and S1L) \cite{10}. \textit{Per2Venus/Venus} mice showed comparable acceleration of SCN and behavioral rhythms. Thus, \textit{Per2Venus} encodes an endogenous fusion protein that functions competently within the mammalian clock.

---

**Figure 1.** PER2::VENUS Fusion Protein Is a Competent Circadian Clock Protein Suitable for Real-Time Imaging

(A) PER2::VENUS fluorescence across the mouse brain, at the peak time of SCN expression (ZT12). Inset shows a close up of the SCN. Scale bar, 1 mm.

(B) Bright-field and fluorescence confocal images show \textit{Per2Venus} lung fibroblasts. Scale bar, 20 \textmu m.

(C) Representative, double-plotted wheel-running actograms for \textit{Per2WT} (left) and \textit{Per2Venus} (right) animals. Mice were entrained on a 12:12 LD cycle, followed by a schedule of constant conditions (dim red light, represented by shaded gray).

(D) Representative, de-trended \textit{Per1}-luc bioluminescence rhythms of SCN slices from \textit{Per2WT} (left) and \textit{Per2Venus} (right) mice are shown.

(E) Mean ± SEM circadian periods for wheel-running are shown (n\textsubscript{WT} = 6; n\textsubscript{WT/V} = 8; n\textsubscript{V/V} = 7).

(F) Mean ± SEM circadian periods for SCN slices (n\textsubscript{WT} = 6; n\textsubscript{WT/V} = 7; n\textsubscript{V/V} = 9). One-way ANOVA revealed no significant effect for either measure.

(G) Snapshots from confocal real-time imaging show PER2::VENUS fluorescence in representative \textit{Per2WT} (top panel), \textit{Per1 null} (middle panel), and CK1\textsubscript{ε}Tau (lower panel) in SCN slices.

(H) Snapshots from confocal real-time imaging show PER2::VENUS in fibroblasts. Scale bar, 20 \textmu m.

(I) Mean fluorescence measures from recordings in (G) are shown.

(J) Mean fluorescence measures from recordings in (H) are shown.

See also Figure S1, Table S1, and Movie S1.
Intracellular Circadian Dynamics of Endogenous PERIOD2

We next analyzed the rhythmicity of the PER2::VENUS protein.

A clear circadian oscillation of PER2::VENUS abundance was detected by western blot in temperature-entrained lung fibroblasts (Figures S1M and S1N). PER2::VENUS was also highly and rhythmically expressed in the SCN (Figures S1O and S1P).

At the peak of PER2 expression (zeitgeber time 12 [ZT12]), in the SCN, PER2::VENUS was detected in effectively all arginine vasopressin (AVP)-immunoreactive (ir) and vasoactive intestinal peptide (VIP)-ir neurons but in <10% of gastrin-releasing peptide (GRP)-ir neurons (Figure S2; Table S1). At the trough of the cycle (CT0), only a few AVP-ir cells expressed PER2::VENUS (Table S1).

We next tested the utility of PER2::VENUS as a real-time circadian reporter, using confocal microscopy. Both SCN slices and lung fibroblasts exhibited stable, high-amplitude circadian oscillations of fluorescence throughout 70 hr of recording (Figures 1G and 1H; Movie S1). Circadian expression of PER2::VENUS was well defined in the SCN with a Per1−/− background and accurately reported period shortening by the CK1ε mutation. Thus, PER2::VENUS is a high-fidelity real-time reporter of the behavior of an endogenous clock protein in SCN neurons and fibroblasts.

We next determined the macrodynamics of PER2::VENUS.

Using cycloheximide to inhibit translation in SCN slices, we revealed that PER2::VENUS has a half-life of ~2 hr, comparable to that of PER2::LUC (Figures S3C–S3E) [10]. Consistent with proteasomal degradation of Per2WT [11], application of the proteasomal inhibitor MG132 at CT12 increased PER2::VENUS levels above those of vehicle-treated lung fibroblasts (Figure S3F) and SCN slices (Figures S3G and S3H). The nuclear export inhibitor leptomycin B, applied at CT12, significantly increased the half-life of PER2::VENUS, suggesting that nuclear export facilitates degradation (i.e., PER2 is subject to degradation in the cytoplasm; Figures S3I and S3J). Moreover, synchronization between cells in SCN slices was reduced following nuclear export blockade, suggesting that timely nuclear export and degradation are necessary for transcriptional-translational feedback loop (TTFL) timing.

Gated nuclear entry of Per protein is considered a pivotal feature of circadian timekeeping. We examined the subcellular localization of PER2::VENUS in the SCN at different points across the LD cycle. Although the overall abundance of PER2::VENUS changed across the cycle, its subcellular localization did not (Figure 2A). Mander’s M1 coefficient indicated almost complete co-localization of Venus with nuclear DAPI signal at all phases (Figure 2B): at no point was PER2::VENUS excluded from the nucleus, with exclusively cytoplasmic localization. To discount the possibility of a transient gate for cytoplasmic retention and nuclear entry, cellular localization was monitored in living SCN slices imaged during the rising phase (CT0–CT9) of PER2 expression. Again, when detected,
PER2::VENUS was observed in the nucleus at all time points (Figure 2C). This was also the case in fibroblasts, where (weak) cytoplasmic fluorescence oscillated in phase with strong nuclear fluorescence (Figures S3K–S3M). Thus, in marked contrast to the temporal gating that delays nuclear fluorescence (Figures S3K–S3M). Thus, in marked contrast to the temporal gating that delays nuclear fluorescence (Figures S3K–S3M).

**Quantitative Analysis of PERIOD2 Intracellular Mobility and Abundance**

Fluorescence correlation spectroscopy (FCS) was used to measure mobility and molecular abundance of PER2::VENUS in skin fibroblasts (Figure 3A). Circadian changes in PER2::VENUS concentration were observed in fibroblast nuclei (Figure 3B); but, importantly, calibration of fluorescence intensity to FCS-calculated concentration per nuclear volume enabled absolute quantification of the molecular abundance of PER2::VENUS across the circadian cycle (Figures 3C and S4A–S4D). This revealed a 10-fold amplitude cycle, with peak expression of ~15,000 molecules per nucleus (>90% of cellular total; Figure 3D). Interestingly, when FCS-derived auto-correlations were fit to a two-component diffusion model, a bimodal distribution of diffusion coefficients was determined for nuclear PER2. This indicates that it exists in at least two molecular/dynamic states, possibly as bound and unbound fractions (Figure 3E). The more mobile fraction had a diffusion coefficient of ~8 μm²/s⁻¹. Furthermore, at CT12 the number of nuclear PER2::VENUS molecules, their diffusion rates, and the proportion displaying the slowly diffusing component were not significantly different in CK1εfibroblasts compared to WT (Figures S3E–S3G). This suggests that PER2 mobility within the nucleus is not affected in the mutant. Thus, we have presented the first quantitative measures of intracellular dynamics of a mammalian circadian clock protein.

Fluorescence recovery after photobleaching (FRAP) was used to further examine the intracellular behavior of PER2::VENUS. Data obtained by photobleaching of fibroblast nuclei agreed with the FCS-based calculations that intranuclear PER2 mobility was unaffected by the CK1εmutation (Figure S4H). Furthermore, after bleaching nuclear fluorescence, full recovery of nuclear fluorescence did not occur within the experimental time frame, confirming that the bulk (>90%) of PER2::VENUS was nuclear (Figure S4I). FRAP also was measured in SCN slices (Figure 4), yielding diffusion coefficients within cytoplasm or nucleus of ~0.2 μm²/s⁻¹. This is comparable to the slow diffusing component revealed by FCS in fibroblasts (Figure 3E). The diffusion coefficient was calculated for both cytoplasm and nucleus, at CT12 and CT2. Intranuclear mobility was comparable for both cytoplasm and nucleus at both time points (Figure 4D). PER2 mobility was not reduced in the nucleus, which may have been expected for a transcriptional regulator. Translocation of PER2 into the nucleus is critical for the circadian TTFL \[6, 7\]. Thus, CT12 is the critical point for the circadian TTFL [6, 7]. Thus, T₁/₂ of FRAP was measured after photobleaching either the whole nucleus or cytoplasm (Figures 4B and 4C; Movie S2) to quantify between-compartment mobility. Importantly, this showed that there is no compartmental or temporal restriction over the mobility of PER2::VENUS (Figure 4E), in agreement with the confocal time-lapse imaging data (i.e., movement between compartments was comparable in both directions).

CK1ε activity is an important regulator of the TTFL, where CK1-mediated phosphorylation of PER proteins licenses them for ubiquitination and proteasomal degradation. To test whether CK1 modulates PER2 mobility, we treated SCN slices with CK1 inhibitor PF670462, which slowed Per1::Luc oscillations to \[28.6 ± 0.30 \text{ hr (n = 10, p < 0.01 versus pre-treatment)}\].
Intracompartment mobility was unaffected by treatment with PF670462 (Figures S4J and S4K); but, surprisingly, both at CT2 and CT12 it significantly decreased the T₁⁄₂ of FRAP (Figure 4E; i.e., the rates of PER2::VENUS bi-directional translocation were accelerated). Furthermore, the magnitude of the increases in T₁⁄₂ were consistent across time points (CT2/CT12) and direction of movement (Figure S4K). Thus, decreasing CK1 activity increased PER2::VENUS translocation rates in an unbiased manner. In conclusion, low CK1ε/δ activity, which significantly lengthened the period of the SCN TTFL, was associated with accelerated nucleo-cytoplasmic shuttling of PER2::VENUS.

DISCUSSION

The PER2::VENUS mouse enables direct qualitative and quantitative observations of the complex spatiotemporal behavior of the PER2 protein, in a physiologically relevant setting. Our results suggest a pivotal role for the dynamic balance of nuclear entry and export for the determination of circadian period.

The PER2::VENUS allele was validated as clock competent, a property shared with PER2::LUC [8, 12]. Luciferase reporter systems, however, are not suitable for measuring fast events, such as nucleo-cytoplasmic shuttling of proteins. They produce a dim signal necessitating a long integration time for detection, and their inherent requirement for luciferin substrate can generate luciferase-chemistry-dependent artifacts. Fluorescent proteins do not suffer from these problems, but there are, of course, some potential limitations, including altered stability of the endogenous protein. The half-life of PER2::VENUS was similar to PER2::LUC, suggesting that the Venus tag did not alter the stability of the PER2 protein. PER2::LUC had a slightly shorter half-life, but this is in line with published literature reporting that luciferase-tagged reporters have a shorter half-life than their endogenous counterparts [13]. The PER2::VENUS half-life may be a more accurate estimate of PER2 stability, as it is a direct measure of the protein, rather than using enzymatic luciferase activity, which is an indirect measure. Prolonged fluorescence imaging can cause both phototoxicity and photobleaching, but for PER2::VENUS we found that it was possible to image SCN slices and fibroblast cultures over at least 70 hr without loss of fluorescence or circadian competence. Although there was limited detection of cytoplasmic PER2::VENUS in fibroblasts, photobleaching of the entire cytoplasm only reduced the nuclear fluorescence by <2% at CT12; thus, it is unlikely to contribute to the overall behavior of PER2 in this cell type. Overall, PER2::VENUS is a useful and faithful reporter for monitoring PER2 dynamics over shorter and longer timescales.

In contrast to the Drosophila ortholog, dPer, which accumulates in the cytoplasm prior to nuclear entry [3, 4], PER2 is not subject to a circadian gate in SCN neurons and in fibroblasts. This contrasting behavior may be explained by the proteins having different hetero-dimerization partners (dTIm and CRY, respectively). Our data do not preclude nuclear gating of other clock factors, but there is, so far, no evidence for this. For example, snapshot immunostaining of PER1 in SCN highlights nuclear expression and no cytoplasmic restriction [14]. Progression of the TTFL in the mammalian clock, therefore, is not dependent on gated nuclear translocation; rather, it is achieved by graded nuclear accumulation of the negative regulators.
PER2::VENUS enabled us to quantify the number of PER2 molecules per cell and determine the dynamic changes in protein mobility. At CT12, PER2::VENUS was present at a concentration of ~15–20 nM in fibroblast nuclei, equating to >10,000 molecules. Interestingly, stochastic simulations of the mammalian circadian clock predicted that stability within the virtual TTFL required >4,000 molecules [15]. Thus, our real observations are of the same order of magnitude and can inform future development of in silico models. The Per2VENUS mouse also facilitates analysis of how the intracellular behavior of PER2 directs the properties of the clock. Both FRAP and FCS revealed diffusion coefficients principally in the range of 0.1 to 1.0 μm²s⁻¹. This is broadly compatible with data from other dynamic transcription factor proteins [16]. FCS also identified a more mobile fraction in fibroblasts, with a coefficient of ~10 μm²s⁻¹. The obtained fast and slow diffusion constants (~10 and ~1 μm²s⁻¹), as well as free and bound fractions, are comparable to estimates made by FCS for the in vivo binding of glucocorticoid receptor [17].

Numerous studies link CK1 to the regulation of PER2 function and localization [18–20]. Both CK1εCas and hPer2S662G are mutants that have a short circadian period, with altered phosphorylation of PER2. The former exhibits rapid clearance of PER2 from the nucleus [10, 21] and the latter decreased nuclear retention [22–24]. These are two aspects of the same phenomenon. Our data demonstrate that CK1 contributes to the translocation of PER2 through the nuclear pores. Thus, CK1 is a critical regulator of PER2 mobility and circadian period, although the contribution of various types of mutation to short periods is not fully resolved [25, 26].

In conclusion, PER2::VENUS operates as a functional circadian protein, and it is sufficient to sustain timekeeping in the absence of WT Per proteins. We believe that the fundamental observations we have presented will support a significant reappraisal of the mammalian clock mechanism and provide valuable observational data on an endogenous clock protein that will inform the development of enhanced quantitative models of cellular circadian timekeeping.

SUPPLEMENTAL INFORMATION

Supplemental Information includes four figures, one table, Supplemental Experimental Procedures, and two movies and can be found with this article online at http://dx.doi.org/10.1016/j.cub.2016.05.018.

AUTHOR CONTRIBUTIONS

N.J.S. and V.P. contributed equally to this work. N.J.S. designed, performed, and analyzed all SCN-related experiments. V.P. contributed to the generation and initial behavioral validation of the mouse, with input from E.S.M., Q.-J.M., and R.B.-H. J.B. conducted the FCS experiments with input from M.R.H.W. and D.G.S. B.S. managed breeding and generated primary skin fibroblasts for FCS experiments. J.E.C. supervised breeding and conducted wheel-running experiments. T.P.K. performed western blotting experiments with input from Q.-J.M., J.B., D.G.S., and E.S.M.

ACKNOWLEDGMENTS

All animals were cared for in accordance with the UK Animals (Scientific Procedures) Act of 1986 with local ethical approval. We thank the biomedical facilities' staffs at the MRC-LMB Ares facility and the University of Manchester, We also thank Nick Barry (MRC-LMB) for expert assistance with confocal microscopy. This work was funded by Biotechnology and Biological Sciences Research Council (BBSRC) grants (BB/E022553/1 to A.S.I.L., BB/K003097/1 to A.S.I.L. and M.R.H.W., and BB/E023223/1 to M.H.H.) and the MRC (MC_U105170643 to M.H.H. and MR/K015865/1 A.S.I.L. and M.R.H.W.).

Received: March 18, 2016
Revised: May 3, 2016
Accepted: May 5, 2016
Published: June 30, 2016

REFERENCES

1. Shearman, L.P., Sriram, S., Weaver, D.R., Maywood, E.S., Chaves, I., Zheng, B., Kume, K., Lee, C.C., van der Horst, G.T.J., Hastings, M.H., and Reppert, S.M. (2000). Interacting molecular loops in the mammalian circadian clock. Science 288, 1013–1019.
2. Koike, N., Yoo, S.H., Huang, H.C., Kumar, V., Lee, C., Kim, T.K., and Takahashi, J.S. (2012). Transcriptional architecture and chromatin landscape of the core circadian clock in mammals. Science 338, 349–354.
3. Meyer, P., Saez, L., and Young, M.W. (2006). PER-TIM interactions in living Drosophila cells: an interval timer for the circadian clock. Science 317, 222–229.
4. Shafer, O.T., Rosbash, M., and Truman, J.W. (2002). Sequential nuclear accumulation of the clock proteins period and timeless in the pacemaker neurons of Drosophila melanogaster. J. Neurosci. 22, 5946–5954.
5. Tamamini, F., Yagita, K., Okamura, H., and van der Horst, G.T.J. (2005). Nucleocytoplasmic shuttling of clock proteins. Methods Enzymol. 393, 418–435.
6. Yagita, K., Tamamini, F., Yasuda, M., Hoeijmakers, J.H.J., van der Horst, G.T.J., and Okamura, H. (2002). Nucleocytoplasmic shuttling and mCRY-dependent inhibition of ubiquitylation of the mPER2 clock protein. EMBO J. 21, 1301–1314.
7. Yagita, K., Yamaguchi, S., Tamamini, F., van Der Horst, G.T.J., Hoeijmakers, J.H.J., Yasui, A., Lorous, J.J., Dunlap, J.C., and Okamura, H. (2000). Dimerization and nuclear entry of mPER proteins in mammalian cells. Genes Dev. 14, 1353–1363.
8. Yoo, S.H., Yamazaki, S., Lowrey, P.L., Shimomura, K., Ko, C.H., Buhr, E.D., Siepka, S.M., Hong, H.K., Oh, W.J., Yoo, O.J., et al. (2004). PERIOD2:LUCIFERASE real-time reporting of circadian dynamics reveals persistent circadian oscillations in mouse peripheral tissues. Proc. Natl. Acad. Sci. USA 101, 5339–5346.
9. Maywood, E.S., Chesham, J.E., Smyllie, N.J., and Hastings, M.H. (2014). The Tau mutation of casen kinase Tε sets the period of the mammalian pacemaker via regulation of Period1 or Period2 clock proteins. J. Biol. Rhythms 29, 110–118.
10. Meng, Q.J., Logunova, L., Maywood, E.S., Gallego, M., Lebiecki, J., Brown, T.M., Slodke, M., Semikhodskii, A.S., Glossop, N.R.J., Piggins, H.D., et al. (2008). Setting clock speed in mammals: the CK1 epsilon tau mutation in mice accelerates circadian pacemakers by selectively destabilizing PERIOD proteins. Neuron 58, 78–88.
11. Eide, E.J., Woolf, M.F., Kang, H., Woolf, P., Hurst, W., Camacho, F., Velihaber, E.L., Giovanni, A., and Virshup, D.M. (2005). Control of mammalian circadian rhythm by CKI epsilon-regulated proteasome-mediated PER2 degradation. Mol. Cell. Biol. 25, 2795–2807.
12. Tahara, Y., Kuroda, H., Saito, K., Nakajima, Y., Kubo, Y., Otsuka, M., Fuse, Y., Ohura, Y., et al. (2012). In vivo monitoring of peripheral circadian clocks in the mouse. Curr. Biol. 22, 1029–1034.
13. Millar, A.J., Short, S.R., Chua, N.H., and Kay, S.A. (1992). A novel circadian phenotype based on firefly luciferase expression in transgenic plants. Plant Cell 4, 1075–1087.
14. Field, M.D., Maywood, E.S., O’Brien, J.A., Weaver, D.R., Reppert, S.M., and Hastings, M.H. (2000). Analysis of clock proteins in mouse SCN demonstrates phylogenetic divergence of the circadian clockwork and resetting mechanisms. Neuron 25, 437–447.
15. Forger, D.B., and Peskin, C.S. (2005). Stochastic simulation of the mammalian circadian clock. Proc. Natl. Acad. Sci. USA 102, 321–324.
16. Bagnall, J., Boddington, C., Boyd, J., Brignall, R., Rowe, W., Jones, N.A., Schmidt, L., Spiller, D.G., White, M.R., and Paszek, P. (2015). Quantitative dynamic imaging of immune cell signalling using lentiviral gene transfer. Integr. Biol. (Camb.) 7, 713–725.
17. Stasevich, T.J., Mueller, F., Michelman-Ribeiro, A., Rosales, T., Knutson, J.R., and McNally, J.G. (2010). Cross-validating FRAP and FCS to quantify the impact of photobleaching on in vivo binding estimates. Biophys. J. 99, 3093–3101.
18. Akashi, M., Tsuchiya, Y., Yoshino, T., and Nishida, E. (2002). Control of intracellular dynamics of mammalian period proteins by casein kinase I epsilon (CKI epsilon) and CKIdelta in cultured cells. Mol. Cell. Biol. 22, 1693–1703.
19. Takano, A., Isojima, Y., and Nagai, K. (2004). Identification of mPer1 phosphorylation sites responsible for the nuclear entry. J. Biol. Chem. 279, 32578–32585.
20. Vielhaber, E., Eide, E., Rivers, A., Gao, Z.H., and Virshup, D.M. (2000). Nuclear entry of the circadian regulator mPER1 is controlled by mammalian casein kinase I epsilon. Mol. Cell. Biol. 20, 4888–4899.
21. Dey, J., Carr, A.J.F., Cagampang, F.R.A., Semikhodskii, A.S., Loudon, A.S.I., Hastings, M.H., and Maywood, E.S. (2005). The tau mutation in the Syrian hamster differentially reprograms the circadian clock in the SCN and peripheral tissues. J. Biol. Rhythms 20, 99–110.
22. Toh, K.L., Jones, C.R., He, Y., Eide, E.J., Hinz, W.A., Virshup, D.M., Ptácek, L.J., and Fu, Y.H. (2001). An hPer2 phosphorylation site mutation in familial advanced sleep phase syndrome. Science 291, 1040–1043.
23. Vanselow, K., Vanselow, J.T., Westermark, P.O., Reischl, S., Maier, B., Korte, T., Herrmann, A., Herzl, H., Schlosser, A., and Kramer, A. (2006). Differential effects of PER2 phosphorylation: molecular basis for the human familial advanced sleep phase syndrome (FASPS). Genes Dev. 20, 2660–2672.
24. Xu, Y., Toh, K.L., Jones, C.R., Shin, J.Y., Fu, Y.H., and Ptácek, L.J. (2007). Modeling of a human circadian mutation yields insights into clock regulation by PER2. Cell 128, 59–70.
25. Vanselow, K., and Kramer, A. (2007). Role of phosphorylation in the mammalian circadian clock. Cold Spring Harb. Symp. Quant. Biol. 72, 167–176.
26. Zhou, M., Kim, J.K., Eng, G.W.L., Forger, D.B., and Virshup, D.M. (2015). A Period2 phosphoswitch regulates and temperature compensates circadian period. Mol. Cell 60, 77–88.