Photoperiod alters phase difference between activity onset in vivo and mPer2::luc peak in vitro

Carl T. Mickman,1 Jeremy S. Stubblefield,1 Mary E. Harrington,2 and Dwight E. Nelson1

1Department of Biology, University of St. Thomas, St. Paul, Minnesota; and 2Department of Psychology, Smith College, Northampton, Massachusetts

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Mickman C, Stubblefield J, Harrington M, Nelson DE. Photoperiod alters phase difference between activity onset in vivo and mPer2::luc peak in vitro. Am J Physiol Regul Integr Comp Physiol 295: R1688–R1694, 2008. First published September 3, 2008; doi:10.1152/ajpregu.90510.2008.—Photoperiod is a significant modulator of behavior and physiology for many organisms. In rodents changes in photoperiod are associated with changes in circadian period and photic resetting of circadian pacemakers. Utilizing rhythms in vivo behavior and in vitro mPer2::luc expression, we investigated whether different entrainment photoperiods [light:dark (L:D) 16:8 and L:D 8:16] alter the period or phase relationships between these rhythms and the entraining light cycle in Per2::luc C57BL/6J mice. We also tested whether mPer2::luc rhythms differ in anterior and posterior suprachiasmatic nucleus (SCN) slices. Our results demonstrate that photoperiod significantly changes the timing of the mPer2::luc peak relative to the time of light offset and the activity onset in vivo. In both L:D 8:16 and L:D 16:8 the mPer2::luc peak maintained a more stable phase relationship to activity offset, while altering the phase relationship to activity onset. After the initial cycle in culture, the period, phase, and peaks per cycle were not significantly different for anterior vs. posterior SCN slices taken from animals within one photoperiod. After short-photoperiod treatment, anterior SCN slices showed increased-amplitude Per2::luc waveforms and posterior SCN slices showed shorter-duration peak width. Finally, the SCN tissue in vitro did not demonstrate differences in period attributable to photoperiod pretreatment, indicating that period aftereffects observed in behavioral rhythms after long- and short-day photoperiods are not sustained in Per2::luc rhythms in vitro. The change in phase relationship to activity onset suggests that Per2::luc rhythms in the SCN may track activity offset rather than activity onset. The reduced amplitude rhythms following long-photoperiod treatment may represent a loss of coupling of component oscillators.

SEASONAL CHANGES IN DAY LENGTH or photoperiod are associated with significant plasticity in behavior and physiology. In mammals, photoperiod changes induce changes in reproductive state, coat color, and thermoregulation, as well as changes in behavior and activity levels (3). Behavioral activity rhythms in mice and hamsters entrained to long- or short-day photoperiods reveal long-lasting changes in the timing of activity onset and activity duration, as well as the period, and photic responsiveness of circadian activity rhythms (15). Coincidence of a light-sensitive circadian clock phase with light exposure during long photoperiods (but not during short photoperiods) may serve as a mechanism for photoperiod detection (5).

In mammals the suprachiasmatic nucleus (SCN) of the hypothalamic functions as a light-responsive circadian pacemaker, maintaining its own endogenous rhythm and synchronizing rhythms in other tissues (20). A complex interaction of up to 15 genes, known as clock genes, and their products drives cellular circadian oscillations (7). Of the known clock genes, the period1 and period2 genes are particularly important for circadian responses to environmental light cues. Light increases transcription of mPer1 and mPer2 in the SCN, initiating a cascade that can reset and entrain endogenous circadian rhythms (18).

Exposure to short-day vs. long-day photoperiods changes the timing and expression patterns of mPer1 and mPer2 in vivo. Per2 mRNA remained elevated above baseline in Syrian hamsters for longer durations when housed under long days than when under short days, even after hamsters became refractory to the effects of short days (2, 24). Similar effects are observed for per1 expression in rats (22), Syrian (10) and Siberian hamsters (9, 15), mice (19), and sheep (8). A long-lasting change in duration and amplitude of clock gene expression may provide a “memory” of photoperiod in the circadian clock system. Such a long-lasting change should be able to be preserved in vitro if an isolated SCN is capable of maintaining this altered gene expression pattern, similar to what has been seen in studies of spontaneous firing rates (e.g., Ref. 12).

Most SCN cells show peak Per1 expression linked to the time of activity offset under changing photoperiods, although, interestingly, a group of cells in the anterior SCN seems to show a peak linked to activity onset (6). These variations in phase of SCN neurons may lead to changes in the overall circadian waveform that allow the SCN to preserve a code of day length (6, 13, 16, 17, 25).

Here we tested whether in vivo entrainment to long- or short-day photoperiod alters the circadian rhythm of Per2::Luc bioluminescence (27) in the mouse SCN in vitro. First, we determined whether photoperiod pretreatment altered the phase relationship between the light:dark (L:D) cycle, the activity rhythm in vivo, and the phase of Per2::Luc rhythms in vitro. We predicted that the phase of Per2::Luc rhythm in vitro would be influenced by photoperiod and that the Per2::Luc rhythms within many regions of the SCN in vitro would track the time of activity offset, as suggested by prior studies (6, 13). Second, we tested the hypothesis that short and long photoperiods would change Per2::Luc rhythm amplitude and duration in vitro in a manner similar to that measured for mPer2 expression in vivo. Finally, we tested whether period aftereffects...
observed in behavioral rhythms in vivo after long- and short-day photoperiods (15) would also be observed in Per2::Luc rhythms in vitro.

MATERIALS AND METHODS

Research was performed in accordance with the “Guiding Principles in the Care and Use of Animals” (American Physiological Society), and protocols were reviewed and approved by the Institutional Animal Care and Use Committees of the University of St. Thomas and Smith College, an Association for Assessment and Accreditation of Laboratory Animal Care-certified facility.

mPer2::Luc bioluminescence. Adult male and female C57BL/6J mPer2 luc (mPeriod2 luc [reverse]) mice were bred from in-house colonies (founders courtesy of Dr. E. Herzog, Washington University, St. Louis, MO; originally derived from the colony of Dr. J. Takahashi, Northwestern University, Evanston, IL). For the initial experiment mice were group housed (4 per cage) for 3–4 wk under L:D 16:8 (16-h light:8-h darkness, lights on at 0400; n = 13) or 4–5 wk under L:D 8:16 (lights on at 0800; n = 12). In a subsequent trial we tested mice entrained to L:D 18:6 for 4 wk with lights on at 0300 (n = 9).

For all experiments food and water were available ad libitum. Mice were 9–14 wk of age at the time of slice preparation and were homozygous for the mPer2 luc gene.

Mice were killed in the light (at times ranging from 2 to 10 h before lights off) by halothane overdose. The brains were rapidly removed and placed in ice-cold HBSS (Invitrogen, Carlsbad, CA). Coronal sections (300 μm) were made from the anterior and posterior SCN with a vibratome. The majority of the SCN was generally contained within these two sections; therefore, medial SCN is represented in both sections. We compared anterior and posterior SCN sections from the same animal as well as across animals. Hypothalamic sections were trimmed to the SCN and optic chiasm and placed on culture plate inserts (model PICMORG50; Millipore, Billerica, MA) in 35-mm petri dishes (BD FacIon, Franklin Lakes, NJ) with 1.2-ml sterile-filtered (Nalgene, Rochester, NY) culture media (composed of 1×DMEM (cat. no. D5030; Sigma-Aldrich, St. Louis, MO) supplemented with 1×B-27 (Invitrogen), 4 mM l-glutamine (Invitrogen), 25 mM glucose (Sigma), 4.2 mM Na3PO4 (Sigma), 10 mM HEPES (Sigma), 25 U/ml penicillin G (Invitrogen), 34 μM streptomycin sulfate (Invitrogen), and 100 μM beetle luciferin (Promega, Madison, WI). Cultures were maintained at 35.8°C, and bioluminescence was recorded for 1-min durations at 10-min intervals over 7 days with a photomultiplier tube and turntable (Lumicycle, Actimetrics, Evanston, IL).

The Lumicycle analysis package was used to analyze the bioluminescence rhythms. Cycle 0 data (from time of slice preparation through midnight of that evening) were excluded from the phase and period analyses, and only slice records containing three mPer2::Luc peaks were used. Records meeting these criteria showed significant rhythms as assessed by χ² periodogram. Lumicycle was used to fit a low-order polynomial to each data set to determine the baseline, which was then subtracted from the raw data. Analyses were then performed on the baseline-subtracted rhythms. mPer2::Luc peaks were determined by finding the time of maximum amplitude between successive troughs following running-mean smoothing. Using the times of mPer2 peak, linear regression was used to determine period and phase angle difference between the mPer2 rhythms and the entraining L:D cycle on day 0 for individual slices. To test for an effect of euthanasia time on circadian phase, slices within each photoperiod were assigned to either an early or late time of euthanasia relative to lights off. Within each photoperiod, mice were killed within 6 h of one another. Slices were designated as early if they were prepared within the first 3 h of this range and late if prepared after this duration. Statistical comparisons between times of mPer2::Luc peak under different photoperiods were determined with Watson-Williams’ F-test and Watson’s U²-test for circular data analyses (Oriana, version 2.02c; Kovach Computing). Differences in circadian period were determined with ANOVA and the Tukey-Kramer method (SAS), whereas differences in waveform were determined with ANOVA followed by Bonferroni tests (SPSS). Statistical significance was determined at P < 0.05.

Behavioral responses to long- and short-day photoperiods. In a separate experiment, male C57BL/6J mice were obtained from Jackson Laboratories (Bar Harbor, ME) at 3–4 wk of age and housed under L:D 16:8 or L:D 8:16 (n = 123 and 121 per group, respectively) with food and water available ad libitum. After 3 wk mice were transferred to individual cages containing running wheels and micro-watches to monitor locomotor activity. Running wheel activity rhythms were recorded and analyzed with a Chronobiology Kit (Stanford Software, Santa Cruz, CA). After preentrainment mice were released into constant darkness (D:D) for 1 wk to allow expression of free-running activity rhythms. These behavioral data were analyzed to determine the circadian period and activity duration (onset and offset) for the rhythm of locomotor activity over the first 7 circadian cycles after L:D 16:8 or L:D 8:16. Activity onsets and offsets were analyzed with eye-fit regression. Phase of activity onset and offset in D:D were extrapolated to the final cycle of L:D to determine the phase relationship between the behavioral rhythms and the entraining light cycle. Statistical differences between activity phases were determined with circular statistics (Oriana, version 2.02c; Kovach Computing) using Watson-Williams’ F-test and Watson’s U²-test. Statistical differences in circadian period were determined with ANOVA and the Tukey-Kramer method (SAS). Significance was determined at P < 0.05.

RESULTS

Comparisons of entrainment phase for rhythms of bioluminescence and behavioral activity. After entrainment to L:D 16:8 or L:D 8:16, rhythms of mPer2::Luc bioluminescence were obvious for both anterior and posterior SCN slices. The mPer2::Luc rhythms from anterior slices were more variable in bioluminescence baseline (Fig. 1) and phase (Figs. 2 and 3) than posterior slices. Rhythms from three randomly chosen animals from each housing condition are shown in Fig. 1.

Significant differences in phase were measured after entrainment to different photoperiods (Figs. 2 and 3). For anterior slices the peak for mPer2::Luc rhythms was +5.88 ± 2.72 h (circular mean ± circular SD, mean relative to lights off; n = 9) after L:D 16:8 and −0.67 ± 1.63 h (n = 10) after L:D 8:16. For posterior slices the peak was +6.27 ± 1.08 h (n = 10) after L:D 16:8 and +0.13 ± 0.98 h (n = 7) after 8:16. The effect of photoperiod on mPer2::Luc phase was significant for both anterior and posterior SCN slices [Watson-Williams F-test: F(1, 15) = 120.0, P = 1.5 × 10⁻⁸; F(1, 17) = 35.4, P = 1.6 × 10⁻⁵, respectively], but there was no significant phase difference between slices obtained from anterior or posterior SCN within the two photoperiods [Watson-Williams F-test: L:D 8:16 slices: F(1, 15) = 1.299, P = 0.271; L:D 16:8 slices: F(1, 17) = 0.164, P = 0.60].

Entrainment to long- or short-day photoperiods also altered running wheel activity rhythms (Fig. 3). Under L:D 16:8 activity onset was significantly earlier (+0.87 ± 0.87 h) than for mice entrained with L:D 8:16 [−0.13 ± 0.78 h; Watson-Williams F-test: F(1, 242) = 86.1, P < 0.001]. Photoperiod also significantly changed activity offset. After L:D 16:8, offset was significantly earlier (−10.50 ± 2.35 h) compared with activity offset in L:D 8:16 [−15.35 ± 2.18 h; Watson-Williams F-test: F(1, 242) = 270.7, P < 0.001].

Thus photoperiod altered the phase difference between activity onset in vivo and mPer2::Luc peak in vitro. This can be
seen in Fig. 3. Under long photoperiods, the in vitro Per2::Luc rhythm peaked earlier relative to activity onset than under short photoperiod. The phase of peak Per2::Luc to activity offset was more conserved as photoperiod was altered, while phase relative to activity onset, dark onset, and light onset were all altered after different photoperiods.

We verified that the time of slice preparation did not influence the phase of subsequent mPer2::luc rhythms in vitro. There were no significant differences between the phase of the bioluminescence peak in vitro in early versus late slice preparation times for L:D 8:16 slices (−0.08 ± 1.13 vs. +0.88 ± 1.55 h, respectively, relative to lights off; Watson-Williams F-test: F(1, 16) = 2.083, P = 0.168) or for L:D 16:8 slices (6.35 ± 2.30 vs. 5.27 ± 1.40 h, respectively; Watson-Williams F-test: F(1, 15) = 0.381, P = 0.546). Within 2–10 h before lights off, differing slice preparation times had no significant effect in mPer2::luc phase in vitro.

Comparison of waveform parameters for bioluminescence after L:D 16:8 and L:D 8:16. To determine the influence of photoperiod on mPer2::luc rhythm waveform, we measured peak width (using the times of baseline crossing) and amplitude (mag-
plitude of peak over mean level) for the day 1 in vitro data. Peak width showed a significant difference across groups \( F(3, 33) = 4.276, P = 0.012 \). Bonferroni post hoc tests indicated that this was due to a smaller peak width in the posterior slices from the short photoperiod compared with the anterior slices in the long photoperiod (mean \( \pm SE: 12.19 \pm 0.25 \) vs. \( 14.54 \pm 0.69 \) h). The groups also showed differences in amplitude of the waveform \( F(3, 33) = 9.59, P < 0.001 \), with post hoc tests revealing that the anterior slices from the short photoperiod showed higher-amplitude waveforms than all the other groups, whereas the other three groups did not show statistical differences from each other (see Fig. 4).

In a final experiment we entrained mice to L:D 18:6 [similar to the photoperiod used by Inagaki et al. (6)] and compared the rhythm waveforms with rhythms after L:D 16:8 and L:D 8:16. To better reveal rhythm differences, we did not smooth or detrend the bioluminescence data and we focused our comparisons on only the initial cycles in vitro. Data from all three photoperiods are shown in Fig. 5. For all three photoperiods and for both anterior and posterior slices there appears to be large between-slice variability in mPer2::luc bioluminescence on the initial cycle in vitro following tissue dissection. The larger variability among rhythms was present for 12–24 h in vitro, and then a more consistent rhythmic pattern emerged. This initial variability often includes an initial “spike” in bioluminescence within the first 1–3 h of placing the tissue in culture, followed by a second, usually smaller peak occurring ~3–5 h later. There was no consistent pattern for the timing or presence of the second peak relating to either photoperiod or SCN location. By 24–36 h in culture, most cultures showed a consistent rhythm that was more obviously circadian in nature and more similar in appearance to that observed in subsequent cycles. We saw no indication for double peaks or other obvious differences in rhythm profile after the initial variability.

**Comparison of period for bioluminescence and behavioral activity.** The period of Per2::luc rhythms in SCN slices did not vary with entrainment photoperiod or with location within the SCN. The mean period for anterior and posterior slices from L:D 16:8 was 24.60 \( \pm 0.13 \) and 24.54 \( \pm 0.13 \) h, respectively. The periods after L:D 8:16 were 24.89 \( \pm 0.11 \) and 24.63 \( \pm 0.09 \) h for anterior and posterior slices, respectively (Fig. 6). There was no significant effect of photoperiod or slice origin on the period of mPer2::luc expression in vitro [photoperiod effect: \( F(1, 34) = 2.4, P = 0.131 \); slice origin effect: \( F(1, 34) = 1.79, P = 0.190 \)].

For behavioral rhythms there was a significant effect of photoperiod on the period of activity onset. Animals from L:D 8:16 showed a longer period than those from L:D 16:8 \( [23.91 \pm 0.03 \text{ h (n = 10)} \) vs. \( 23.71 \pm 0.04 \text{ h (n = 9)} \); \( F(1, 17) = 14.87, P = 0.0013 \)]. The difference in average period in behavioral experiments versus in vitro experiments either may be attributed to the different mouse colonies used in the experiments or may be due to the in vivo/in vitro difference; however, the lack of an effect of photoperiod on period in vitro cannot be attributed to strain differences.

**DISCUSSION**

Our study demonstrated that photoperiod pretreatment altered SCN phase relative to activity onset and also altered several other rhythm parameters as measured in vitro with Per2::luc expression. First, we demonstrated that photoperiod pretreatment altered...
suggest that photoperiod responses are coded by changes in activity offset. A similar change in phase was also seen in the in vivo measurements from posterior slices from mice housed under short photoperiod. This increase in amplitude might indicate a tighter coupling of component oscillators, leading to a greater-amplitude population rhythm, or perhaps increased-amplitude cycles within individual neurons. Single-cell imaging experiments would be able to determine which of these explanations might be correct; one such study suggests that individual neurons do not show differences in amplitude of Per1-luc rhythms after different photoperiods but do show greater distribution of phase (13). Simulations suggest that very large changes in single-cell duration would be required for duration changes at the population level (16).

The waveforms were also analyzed for the period of time between the half-rise and the half-fall of the rhythm to search for predicted changes in duration in different photoperiods. Behavioral alpha measurements showed a difference of 4–5 h in activity length, while photoperiodic differences in bioluminescence showed a peak width difference of ~2 h, and this difference was observed only between two groups, with a smaller peak width in the posterior slices from the short photoperiod compared with the anterior slices in the long photoperiod. This provides some support to indicate that the Per2 measure reflects broadened peak duration as observed with electrophysiological measurements and measurements of per1 in vivo and in vitro, as well as per2 mRNA measurements in vivo. In mice, photoperiod effects on per2 mRNA measured in vivo are dramatic, with changes in both amplitude and duration (19). Effects of photoperiod on per1-luc waveform duration dissipate over several cycles in vitro (13).

Behavioral studies have demonstrated decreased-amplitude phase shift responses in mice from long photoperiods, an effect seen particularly in cycles with 14 or 16 h of light (21). This effect might arise from effective loss of subpopulations of per-expressing neurons from the larger coupled population of cells. This speculative hypothesis assumes that the population of per-expressing neurons that stay coupled to each other subserve the photic resetting, and that the magnitude of the phase shift induced by light is proportionate to the number of cells effectively coupled within this population. Our results support this hypothesis in that we observed decreased-amplitude Per2::Luc rhythms after long photoperiods with 16 h of light. We tentatively consider the measurements from Per2::Luc rhythms to be reflective of the summed activity of the coupled cells in the underlying oscillator, but future studies using other markers would be necessary to determine whether this is a valid interpretation. We interpret the decreased-amplitude Per2::Luc rhythm as resulting from loss of coupling of a subset of the population of cells, leaving fewer cells within the coupled group. The alternate interpretation of the decreased-amplitude Per2::Luc rhythm as resulting from decreased amplitude of each individual oscillator would predict increased phase shift amplitude (26), but this is not supported by available behavioral data (21). Our interpretation considers the altered phase-response curve as a network property and suggests that if we could measure the phase-response curve at a single-cell level it would not show changes due to photoperiod pretreatment.

Anterior sections of the SCN were more variable in expression, as can be seen in both representative data records and

![Diagram](http://ajpregu.physiology.org/)

Fig. 6. Circadian period of mPer2 peak expression and corresponding circadian period of behavioral activity rhythm after L:D 16:8 and L:D 8:16. Period of locomotor activity onset is shown for L:D 8:16 (gray hatched bar) and L:D 16:8 (open hatched bar) photoperiods. Period of locomotor activity is significantly different after the 2 photoperiods (ANOVA, P < 0.05). Period of Per2 peak expression is shown for posterior (P) and anterior (A) SCN for L:D 8:16 (gray bars) and L:D 16:8 (open bars). There was no significant effect of location within the SCN (anterior/posterior) or of prior photoperiod (L:D 8:16 or L:D 16:8) on period in vitro.
summary figures (Figs. 1–3). A similar increase in variability in anterior SCN sections with Per1-luc recordings in long photoperiods might be attributed to separate oscillators in the anterior SCN or flexibility in phase relationships and coupling among regions of the SCN (6, 13). It would be interesting to observe the response of these heterogeneous populations when the system is challenged with a light stimulus. We did not observe two peaks in Per2::Luc expression in rostral SCN after long-photoperiod pretreatment, as was reported by Inagaki et al. (6). Interestingly, they observed that after several days in vitro the bimodal peaks merged into a single peak, so it is possible that the Per2::Luc rhythm fuses more quickly than the per1-luc rhythms and that explains why we did not observe two peaks. It is also possible that our method for dissection and cutting coronal slices was not sufficient to reveal a parallel subpopulation of Per2::Luc cells, and future studies might better demonstrate this. We did see complex waveforms on the day of dissection, but this was unrelated to SCN region or photoperiod pretreatment and was not sustained into the first full cycle in vitro.

Period aftereffects (changes in free-running period induced after entrainment to long- or short-period light cycles) appear to be important for stabilizing entrainment (15). Using entrainment to T cycles (non-24-h period LD cycles), several studies have demonstrated that the SCN shows paradoxical period aftereffects, in that the SCN period in vitro is negatively correlated with the behavioral rhythm period (1, 11). In our hands, photoperiod pretreatment caused an aftereffect on behavioral rhythm period but had no effect on period of the SCN in vitro. Constant light can lengthen the period of SCN in vitro as well as the behavioral rhythm in vivo (4). Studies looking for evidence of period aftereffects in the isolated SCN give contradictory findings, and further experiments will be needed to clarify the reasons for these disparate findings.

Perspectives and Significance

These findings suggest that a robust outcome of exposure to varied photoperiod is the change in phase of the bulk of Per2-expressing cells relative to activity onset. The change in phase is such that it allows conservation of phase to activity offset, a finding mirrored in recent studies measuring mPer1 expression in vitro (6, 13). We suggest that this change in phase and the loss of coupling of a subpopulation of the light-responsive cells expressing circadian clock genes might help to explain alterations in photic sensitivity of circadian rhythms with seasonal changes. Changes in internal phase relationships among SCN cells might explain changes in magnitude of photic resetting responses.

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Present addresses: J. Stubblefield, Dept. of Biology, University of Virginia, Charlottesville, VA; D. E. Nelson Medtronic Neuromodulation Research, Medtronic Parkway NE, Minneapolis, MN 55432.

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