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Daniel Granados-Fuentes  
*Washington University in St. Louis*

Aaron J. Norris  
*Washington University School of Medicine in St. Louis*

Yarimar Carrasquillo  
*Washington University School of Medicine in St. Louis*

Jeanne M. Nerbonne  
*Washington University School of Medicine in St. Louis*

Erik D. Herzog  
*Washington University School of Medicine in St. Louis*

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I_A Channels Encoded by Kv1.4 and Kv4.2 Regulate Neuronal Firing in the Suprachiasmatic Nucleus and Circadian Rhythms in Locomotor Activity

Daniel Granados-Fuentes,1* Aaron J. Norris,2* Yamir Carrasquillo,2 Jeanne M. Nerbonne,2 and Erik D. Herzog1

1Department of Biology and 2Department of Molecular Biology and Pharmacology, School of Medicine, Washington University, St. Louis, Missouri 63130-4899

Neurons in the suprachiasmatic nucleus (SCN) display coordinated circadian changes in electrical activity that are critical for daily rhythms in physiology, metabolism, and behavior. SCN neurons depolarize spontaneously and fire repetitively during the day and hyperpolarize, drastically reducing firing rates, at night. To explore the hypothesis that rapidly activating and inactivating A-type (I_A) voltage-gated K^+ (Kv) channels are critical regulators of the excitability of SCN neurons, we examined locomotor activity and SCN firing in mice lacking Kv1.4 (Kv1.4^-/-), Kv4.2 (Kv4.2^-/-), or Kv4.3 (Kv4.3^-/-), the pore-forming (alpha) subunits of I_A channels. Mice lacking either Kv1.4 or Kv4.2 alpha subunits have markedly shorter (0.5 h) periods of locomotor activity than wild-type (WT) mice. In vitro extracellular multi-electrode recordings revealed that Kv1.4^-/- and Kv4.2^-/- SCN neurons display circadian rhythms in repetitive firing, but with shorter periods (0.5 h) than WT cells. In contrast, the periods of wheel-running activity in Kv4.3^-/- mice and firing in Kv4.3^-/- SCN neurons were indistinguishable from WT animals and neurons. Quantitative real-time PCR revealed that the transcripts encoding all three Kv channel alpha subunits, Kv1.4, Kv4.2, and Kv4.3, are expressed constitutively throughout the day and night in the SCN. Together, these results demonstrate that Kv1.4- and Kv4.2-encoded I_A channels regulate the intrinsic excitability of SCN neurons during the day and night and determine the period and amplitude of circadian rhythms in SCN neuron firing and locomotor behavior.

Introduction

The suprachiasmatic nucleus (SCN) acts as a master circadian pacemaker driving daily rhythms in physiology and behavior (Dibner et al., 2010; Welsh et al., 2010). SCN neurons undergo changes in membrane potential and repetitive firing with a near 24 h period (Schwartz and Zimmerman, 1991; Welsh et al., 1995; Colwell, 2011). During the day, SCN neurons depolarize and fire action potentials repetitively; during the night, however, SCN neurons hyperpolarize and rarely fire. These daily rhythms in membrane potential and spontaneous firing depend on (Liu et al., 1997; Herzog et al., 1998; Albus et al., 2002; Nakamura et al., 2002) and modulate a transcription-translation feedback loop (Lundkvist and Block, 2005; Nitabach et al., 2005; Brown and Piggins, 2007). Currently, the mechanisms linking the daily changes in gene expression and membrane excitability are poorly understood.

The resting and active membrane properties of SCN neurons are determined by the interaction of multiple ionic conductances that function at the resting potential (Pennartz et al., 2002; Häusser et al., 2004; Jackson et al., 2004). The spontaneous daytime depolarization in membrane potential of SCN neurons is accompanied by an increase in input resistance (de Jeu et al., 1998, 2002; Kuhlman and McMahon, 2004) suggesting that decreased subthreshold K^+ conductance(s) mediate the daytime depolarization and the increased firing of action potentials. Conversely, the nighttime hyperpolarization is associated with a decrease in input resistance, consistent with increased subthreshold K^+ conductance(s). In addition, injection of depolarizing current converts SCN neurons from the electrically quiet nighttime state to regular firing, further supporting the hypothesis that subthreshold K^+ channels are crucial regulators of the excitability of SCN neurons (Kuhlman and McMahon, 2004, 2006). Although specific roles for various K^+ currents in determining the excitability of SCN neurons have been proposed (Kononenko et al., 2008; Colwell, 2011), exploring these hypotheses directly has been hindered by a lack of knowledge about the channel proteins responsible for individual currents and the limited availability of selective channel blockers.

A-type (I_A) voltage-gated K^+ (Kv) channels activate and inactivate rapidly on membrane depolarization and, on hyperpolarization, recover rapidly from inactivation (Connor and...
Stevens, 1971a; Birnbaum et al., 2004; Jerg et al., 2005; Covarrubias et al., 2008). In many neurons, these properties impact repetitive firing rates (Connor and Stevens, 1971b; Kang et al., 2005; Yuan et al., 2005). In some cells, IA channels are active at subthreshold membrane potentials, influencing cell input resistances and excitability (de Jeu et al., 2002; Yuan et al., 2005). IA is readily detected in SCN neurons and has been suggested to function in the regulation of repetitive firing rates (Huang, 1993; Bousskila and Dudek, 1995; Alvado and Allen, 2008). Using mice harboring targeted disruptions in the genes encoding the voltage-gated K+ (Kv) channel pore-forming (α) subunits, Kcnq4 (Kv1.4-/-), Kcnq2 (Kv4.2-/-), or Kcnq3 (Kv4.3-/-) (Norris and Nerbonne, 2010), we directly tested the necessity of these α subunits in the generation of IA in SCN neurons and in regulating circadian rhythms in SCN neuron firing and locomotor behavior.

Materials and Methods

Animals. Mice were maintained on a C57BL/6 background in the Danforth and Medical School animal facilities at Washington University. The four genotypes of mice used in this study were wild-type (WT) mice and mice harboring targeted genetic disruptions of the Kcnd2 (Kv4.2-/-) (London et al., 1998), Kcnq2 (Kv4.2-/-) (Guo et al., 2005), or Kcnq3 (Kv4.3-/-) (Niwa et al., 2008) locus. All procedures were approved by the Animal Care and Use Committee of Washington University and conformed to US National Institutes of Health guidelines.

Behavioral recordings. Adult (8- to 10-week-old) Kv1.4-/- (n = 17), Kv4.2-/- (n = 17), Kv4.3-/- (n = 6), and WT (n = 19) male mice were housed individually in cages equipped with a running wheel in light-tight chambers illuminated with fluorescent bulbs (2.4 ± 0.5 × 10^10 photons/s m^-2; General Electric). Running-wheel activity was recorded in 6 min bins (ClockLab software; Actimetrics) for 5–10 d in a 12 h light (L)/dark (D) cycle (lights on at 7:00 A.M.), 11–12 d in constant (DD), 10–18 d in the LD cycle (lights on at 7:00 A.M.), 15–16 d in a 6 h delayed LD cycle (lights on at 1:00 P.M.), and finally for 15–17 d in a 6 h advanced LD cycle (lights on at 7:00 A.M.). The period of behavioral rhythmicity of each mouse was determined using χ^2 periodogram analysis (Sokolove and Bushell, 1978) from continuous recordings of 10 d in DD (ClockLab software). Rhythmicity was considered statistically significant if the χ^2 periodogram value exceeded the 99.9% confidence interval (Qp value). Additionally, the phase angle of entrainment in LD, number of days to re-entrain after shifts in the LD cycle, and total daily activity counts in DD were calculated for each mouse (ClockLab).

Cell culture and multi-electrode array recordings. SCNs were explanted from 3- to 7-d-old Kv1.4-/-, Kv4.2-/-, Kv4.3-/-, or WT mice housed in a 12 h LD cycle. Genotypes were confirmed by PCR of tail DNA. For dispersed cultures, 4–6 tissue punches (400 μm in diameter) containing the SCN, were obtained from 200-μm-thick coronal slices, and neurons were enzymatically dissociated using papain, as previously described (Herzog et al., 1998). Viable neurons were plated at a density of at least 10,000 neurons/mm^2 onto poly-d-lysine and laminin-coated multi-electrode arrays according to published methods (Aton et al., 2005) (60 10 μm diameter electrodes; Multichannel Systems). We maintained dispersed neuronal cultures in 1 ml of CO2-buffered DMEM (Sigma-Aldrich) medium with 10% fetal calf serum at 37°C with 5% CO2, for 2 weeks and then recorded extracellular action potentials for at least 5 d as described previously (Aton et al., 2005). Action potentials were digitized in real time (MC-Rack Software; Multichannel Systems) and discrimi-
nated off-line using principal component analysis (Offline Sorter; Plexon). Firing rates were binned in 10 min intervals (NeuroExplorer; Plexon).

Firing rate rhythms were evaluated by a Fast Fourier transformation as previously described (Aton et al., 2005). Data with a relative amplitude $/H_{11021}$ 0.2 defined neurons with a statistically significant circadian rhythm. Peak and trough firing rates were compared for the four genotypes. We used the Rayleigh test (Batschelet et al., 1981) to determine whether the times of peak firing for groups of SCN neurons were clustered or uniformly distributed.

Electrophysiological recordings from acute brain slices. Brain slices were prepared from 3- to 6-week-old WT, Kv1.4$^{-/-}$, Kv4.2$^{-/-}$, and Kv4.3$^{-/-}$ mice using standard procedures (Aton et al., 2005). Briefly, mice housed in a 12 h LD cycle, with lights on at 7:00 A.M., were deeply anesthetized with 1.25% Avertin (2,2,2-tribromoethanol and tert-amyl alcohol in 0.9% NaCl; 0.025 ml/g body weight) between 12:00 and 2:00 P.M. and then perfused transcardially with ice-cold cutting solution containing the following (in mM): 240 sucrose, 2.5 KCl, 1.25 NaH$_2$PO$_4$, 25 NaHCO$_3$, 0.5 CaCl$_2$, and 7 MgCl$_2$, saturated with 95% O$_2$/5% CO$_2$ before cooling. The brains were rapidly removed and placed in oxygenated ice-cold cutting solution. Coronal slices (350 $/H_{9262}$ m) containing the SCN were cut on a Leica VT1000 S vibrating blade microtome (Leica Microsystems). Slices were incubated in a holding chamber with oxygenated artificial CSF (ACSF) containing the following (in mM): 125 NaCl, 2.5 KCl, 1.25 NaH$_2$PO$_4$, 25 NaHCO$_3$, 0.5 CaCl$_2$, and 7 MgCl$_2$, saturated with 95% O$_2$/5% CO$_2$ before cooling. The brains were rapidly removed and placed in oxygenated ice-cold cutting solution. Coronal slices (350 $/H_{347}$ m) containing the SCN were cut on a Leica VT1000 S vibrating blade microtome (Leica Microsystems). Slices were incubated in a holding chamber with oxygenated artificial CSF (ACSF) containing the following (in mM): 125 NaCl, 2.5 KCl, 1.25 NaH$_2$PO$_4$, 25 NaHCO$_3$, 2 CaCl$_2$, 1 MgCl$_2$, and dextrose 25 at room temperature for at least 30 min before transfer to the recording chamber. Recordings were obtained from SCN neurons in the acute slice between 1:00 and 5:00 P.M.

Whole-cell voltage-clamp recordings were obtained at room temperature (22–24°C) from visually identified SCN neurons using differential interference contrast with infrared microscopy. Data were collected using a Multiclamp 700B patch-clamp amplifier interfaced with a Digidata 1332 and the pCLAMP 9 software (Molecular Devices) to a Gateway computer. Series resistances were compensated electronically by $/>90\%$. Signals were acquired at 20 kHz and filtered at 10 kHz before digitization and storage. The recording pipette solution contained the following (in mM): 130 KCl, 10 HEPES, 10 glu-
cose, 0.83 CaCl2, and 2.6 BAPTA, and 3 MgATP and 0.5 NaGTP were added the day of recording, pH 7.4 (300 mOsm). Tetraethylammonium (3 mM), CdCl2 (0.1 mM), and tetrodotoxin (100 nM) were added to the ACSF immediately before recordings. All reagents were from Sigma unless otherwise noted.

The rapidly activating and rapidly inactivating Kv current, \( I_{\text{A}} \), was isolated by a two-step voltage protocol, using previously described procedures (Norris and Nerbonne, 2010). Briefly, total whole-cell Kv currents were first evoked in response to 4 s depolarizing voltage steps to potentials between \(-70 \text{ mV} \) and \(-40 \text{ mV} \) (in 10 mV increments) from a holding potential of \(-70 \text{ mV} \). A prepulse paradigm that included a brief (60 ms) step to \(-20 \text{ mV} \) before the 4 s depolarizing voltage steps to potentials between \(-40 \text{ mV} \) and \(-40 \text{ mV} \) (in 10 mV increments) was then used. Off-line subtractions of the currents evoked with the prepulse from the currents evoked without the prepulse were performed to isolate \( I_{\text{A}} \).

The steady-state outward Kv current (ISS) was measured as the current remaining at the end of the 4 s depolarizing steps. Data were compiled and analyzed using ClampFit (Molecular Devices), Microsoft Excel, and Prism (GraphPad Software).

Quantitative real-time PCR. Eight-week-old C57BL/6 mice were housed 12 h LD cycle for a week and then switched to constant darkness for 48 h. Mice were killed at specific circadian times (CT0, 4, 8, 12, 16, and 20; \( n = 4 \) per CT) on the third day of constant darkness and SCN punches (400 \( \mu \text{m} \) in diameter) were collected. Total RNA was isolated from the SCN and RNA concentrations were determined by optical density measurements. The expression levels of genes encoding the Kv4.2 (\( Kcnd2 \)), Kv4.3 (\( Kcnd3 \)), and Kv1.4 (\( Kcna4 \)) subunits, as well as the endogenous control gene hypoxanthine guanine phosphoribosyl transferase (HPRT) were determined using Taqman-based real-time PCR in a two-step process as described previously (Yang et al., 2010); experiments were conducted on a 7900HT Sequence Detection System (Applied Biosystems). Data were analyzed using the threshold cycle relative quantification method and were normalized to the expression value for HPRT in the same sample (Schmittgen and Livak, 2008) and evaluated for rhythmicity using COSOPT (Abraham et al., 2005).

Figure 4. \( I_{\text{A}} \) densities are reduced in Kv4.2 \(-/-\) and Kv1.4 \(-/-\) SCN neurons. Aa–Da, Representative whole-cell Kv currents, recorded in response to voltage steps to potentials ranging from \(-40 \text{ mV} \) to \(+40 \text{ mV} \) (in 10 mV increments) from a holding potential of \(-70 \text{ mV} \) in WT (Aa), Kv4.2 \(-/-\) (Ba), Kv4.3 \(-/-\) (Ca), and Kv1.4 \(-/-\) (Da) SCN neurons, are displayed. In each cell, outward Kv currents evoked at the same test potentials were also recorded following a brief prepulse to \(-20 \text{ mV} \) to inactivate \( I_{\text{A}} \). The amplitudes of \( I_{\text{A}} \) in individual cells of each genotype were then obtained by digital off-line subtraction (a-b) of the recordings with the prepulse (b) from the recordings without the prepulse (a); the subtracted records are also shown on an unexpanded time scale to facilitate direct comparisons. Mean (±SEM) \( I_{\text{A}} \) (E) and steady-state Kv current (ISS) (F) in WT (\( n = 10 \)), Kv4.2 \(-/-\) (\( n = 20 \)), Kv4.3 \(-/-\) (\( n = 5 \)), and Kv1.4 \(-/-\) (\( n = 14 \)) SCN neurons are plotted as function of test potential. *Values indicated are significantly different at the \( p < 0.001 \).
mice are presented in Figure 1A. In constant darkness (DD), Kv1.4−/− and Kv4.2−/− mice displayed locomotor activity patterns with shorter circadian periods than WT or Kv4.3−/− mice (one-way ANOVA, $p < 0.001$; 23.8 ± 0.02 h for WT and 23.8 ± 0.04 h for Kv4.3−/−; 23.4 ± 0.07 h for Kv1.4−/− and 23.5 ± 0.07 h for Kv4.2−/−, mean ± SEM). Over 50% of the Kv1.4- or Kv4.2-deficient mice had periods shorter than WT mice (Fig. 1B). In addition, when housed in a 12 h LD cycle, Kv1.4−/− and Kv4.2−/− mice initiated daily wheel running ~0.5 h earlier (Fig. 1C,D; one-way ANOVA, $p < 0.04$) than WT mice.

The Kv1.4−/− and Kv4.2−/− mice also required fewer days to establish a stable phase relationship to the time of daily light onset following a 6 h advance in the light cycle (one-way ANOVA, $p < 0.004$; Fig. 2). It is also of interest to note that the Kv1.4−/− mice showed more total daily wheel running than WT (one-way ANOVA, $p < 0.003$), Kv4.2−/− or Kv4.3−/− mice, primarily due to an increase in the amount of time they were active each night (Figs. 1A,D, 2; one-way ANOVA, $p < 0.003$). In marked contrast to the Kv1.4−/− and the Kv4.2−/− mice, the circadian period, phase angle of entrainment, ability to adjust to shifts in light schedule, and the daily wheel-running activity of Kv4.3−/− mice were statistically indistinguishable from WT mice.

$I_\alpha$ densities are reduced in Kv1.4−/− and Kv4.2−/− mice

The observation that the alterations in the circadian patterns in wheel-running behavior of the Kv1.4−/− and Kv4.2−/− mice, compared with WT mice, were similar suggests that Kv1.4- and Kv4.2-encoded $I_\alpha$ channels play similar roles in determining the onset and the period of circadian locomotor rhythms. Quantitative real-time PCR (qRT-PCR) analyses of mRNA transcripts from SCN samples revealed that the Kv1.4, Kv4.2, and Kv4.3 transcripts were readily detected in the mouse SCN. In contrast with Per2, the expression levels of the three Kv $\alpha$ subunits did not vary significantly with CT (Fig. 3; COSOPT test, $p > 0.3$).

Whole-cell voltage-clamp recordings from SCN neurons in acute brain slices demonstrated that $I_\alpha$ was present in all SCN cells examined, consistent with previous reports (Bouskila and Dudek, 1995; Itri et al., 2010), and that the densities of the currents were similar among SCN neurons. Voltage-clamp recordings from Kv1.4−/− and Kv4.2−/− neurons revealed that the loss of Kv1.4 or Kv4.2 attenuated mean $I_\alpha$ densities (Student’s $t$ test, $p < 0.001$) by ~50% (Fig. 4). The mean $I_\alpha$ density in Kv4.3−/− SCN neurons, in contrast, was not significantly different from

The primers sequences used to detect transcript expression were as follows: Kv4.2 (KcnD2): 5′-TGAATCAGTTTGTGCATTAGTGAA and 5′-TTCGACCGCTCATCTGGAGG; Kv4.3 (KcnD3): 5′-GGCCACGCAGTCTCTTGT and 5′-CCGCTCGAATGTGCTATCATG; Kv4.1 (KcnA4): 5′-AGAGCCGGATGGAACCACCTCTGCTCAGT; and HPRT: 5′-TGAATCAGTTTGTGCTTATAAGG and 5′-TTCGACCGCTCATCTGGAGG.

Results

The period of locomotor activity is altered in Kv1.4−/− and Kv4.2−/− mice

Representative recordings of wheel-running activity per unit time (actograms) in WT, Kv1.4−/−, Kv4.2−/−, and Kv4.3−/− neurons displayed markedly shorter circadian firing periods than WT or Kv4.3−/− mice (one-way ANOVA, $p < 0.001$; 23.8 ± 0.02 h for WT and 23.8 ± 0.04 h for Kv4.3−/−; 23.4 ± 0.07 h for Kv1.4−/− and 23.5 ± 0.07 h for Kv4.2−/−, mean ± SEM). Over 50% of the Kv1.4- or Kv4.2-deficient mice had periods shorter than WT mice (Fig. 1B). In addition, when housed in a 12 h LD cycle, Kv1.4−/− and Kv4.2−/− mice initiated daily wheel running ~0.5 h earlier (Fig. 1C,D; one-way ANOVA, $p < 0.04$) than WT mice.

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Firing rates of Kv1.4 \(^{-/-}\) and Kv4.2 \(^{-/-}\) SCN neurons are increased and circadian periods are shorter

To determine whether the markedly shortened periods in wheel-running activity seen in mice lacking either Kv1.4 or Kv4.2 relate to changes in circadian rhythms in SCN neuron firing, we measured the functional consequences of disrupting \(I_A\) on the firing rates of isolated SCN neurons directly. Using multi-electrode arrays, we observed circadian patterns of electrical activity in SCN neurons from WT, Kv1.4 \(^{-/-}\), Kv4.2 \(^{-/-}\), and Kv4.3 \(^{-/-}\) mice (Fig. 5A), although the circadian periods of firing of Kv1.4 \(^{-/-}\) and Kv4.2 \(^{-/-}\) SCN neurons were significantly shorter (0.5 h) than in WT or Kv4.3 \(^{-/-}\) SCN neurons (Fig. 5A; one-way ANOVA, \(p < 0.03\)). The measured periods were 23.9 ± 0.1 h (mean ± SEM) for WT, 23.7 ± 0.2 h for Kv4.3 \(^{-/-}\), 23.4 ± 0.1 h for Kv1.4 \(^{-/-}\), and 22.7 ± 0.2 h for Kv4.2 \(^{-/-}\) neurons. The magnitude of this effect (0.5 h) is similar to the shortening of the period of locomotor activity seen in Kv1.4 \(^{-/-}\) and Kv4.2 \(^{-/-}\) mice (Fig. 1). Analyses of these data also revealed that the circadian periods of firing were shorter in only ~40% of the Kv1.4 \(^{-/-}\) or Kv4.2 \(^{-/-}\) SCN neurons, whereas the remaining cells had circadian periods similar to WT or Kv4.3 \(^{-/-}\) cells (Fig. 5B; Kolmogorov–Smirnov test, \(p < 0.02\)).

Kv1.4 \(^{-/-}\) or Kv4.2 \(^{-/-}\) SCN neurons also fired at higher frequencies during both the subjective night and day than either WT or Kv4.3 \(^{-/-}\) SCN neurons significantly (Fig. 5C; Kolmogorov–Smirnov test, \(p < 0.01\)). The grouping of the daily peaks in the firing rates among Kv1.4 \(^{-/-}\) or Kv4.2 \(^{-/-}\) SCN neurons, however, was not significantly different from WT neurons (Fig. 6; Rayleigh test, \(p > 0.05\)).

Discussion

Together, the findings here demonstrate that the Kv1.4 and Kv4.2 \(\alpha\) subunits encode \(I_A\) channels in the SCN that control the intrinsic excitability, set the circadian firing period, and modulate the firing rates of SCN neurons. The voltage-clamp experiments revealed that the loss of either Kv1.4 or Kv4.2 reduces \(I_A\) density in SCN neurons by ~50%. Although Kv4.3 mRNA is expressed in the SCN and the Kv4.3 protein can associate and form heteromultimeric channels with Kv4.2 (Guo et al., 2002) and has been shown to encode \(I_A\) channels in cortical pyramidal neurons (Norris and Nerbonne, 2010), the loss of Kv4.3 expression had no measureable effects on \(I_A\) densities, electrical activity, or circadian rhythms in SCN neurons.

Circadian locomotor activity is altered in Kv1.4 \(^{-/-}\) and Kv4.2 \(^{-/-}\) mice

The results presented here demonstrate that the loss of either Kv1.4- or Kv4.2-encoded \(I_A\) channels has very striking effects on the circadian period of behavioral rhythms in locomotor activity. Interestingly, loss of either Kv1.4 or Kv4.2 has a much larger effect on the circadian period than the loss of other genes, including some canonical clock genes, such as Clock (DeBruyne et al., 2007). In addition, the 0.5 h shortening of the period of circadian rhythms in locomotor activity of mice lacking either Kv1.4 or Kv4.2 contrasts with the recently reported 0.5 h lengthening of the period of circadian behavior in mice lacking one copy of Scn1a, which encodes the voltage-gated sodium channel \(\alpha\) subunit, Nav1.1 (Han et al., 2012). The results here also contrast markedly with the results of several previous studies conducted on mice with other channel deficiencies, including mice lacking the large conductance, Ca\(^{2+}\) and voltage-dependent, BK (Kcnma1); K\(^{+}\) channel subunit (Meredith et al., 2006); the voltage-gated Ca\(^{2+}\) channel subunit, Cav2.2 (Caenabl) (Beuckmann et al., 2003); or the K channel subunits, Kv3.1 and Kv3.2 (Kcnc1 and Kcnc2) (Kudo et al., 2011). In contrast with the findings here for Kv1.4 \(^{-/-}\) and Kv4.2 \(^{-/-}\) mice, negligible changes in circadian period were reported in BK-, Cav2.2-, or Kv3.1/Kv3.2-deficient mice, suggesting that Kv1.4- and Kv4.2-encoded \(I_A\) channels subserve unique and important roles in modulating the functioning of the SCN and in regulating circadian biology.

Kv1.4- and Kv4.2-encoded \(I_A\) channels regulate the intrinsic excitability of SCN neurons

The finding that both peak and trough firing rates were increased in Kv1.4 \(^{-/-}\) and Kv4.2 \(^{-/-}\) SCN neurons indicates that Kv1.4- and Kv4.2-encoded \(I_A\) channels play critical roles in regulating the intrinsic excitability of SCN neurons during the day and during the night, i.e., throughout the circadian cycle. In addition, the qRT-PCR analyses revealed that the expression levels of the transcripts encoding these subunits do not vary with CT, consistent with previously published in situ hybridization (Lein et al., 2007) and DNA microarray data (Panda et al., 2002; Kasukawa et al., 2011). It has, however, recently been reported that \(I_A\) density is higher during the day than at night in a subpopulation of SCN neurons (Itri et al., 2010). It is certainly possible that \(I_A\) densities are altered in SCN neurons as a result of circadian regulation of
the transcripts encoding critical accessory or regulatory proteins. Interestingly, and consistent with the hypothesis that accessory subunits could play a critical role, it was recently reported that the circadian regulation of Kv4.2-encoded I_{AHP} channels in mouse ventricular myocytes reflects circadian changes in the accessory Kv channel interacting protein, KChIP2, not in the Kv4.2 α subunit (Jeyaraj et al., 2012). Alternatively, post-transcriptional mechanisms, such as changes in Kv α subunit protein expression, localization and/or degradation, or altered interactions with other Kv α subunits, accessory subunits, or other regulatory proteins (Covarrubias et al., 2008; Norris et al., 2010) could mediate circadian changes in I_{AHP} densities in SCN neurons. Additional experiments aimed at exploring each of these possibilities directly will be of considerable interest.

As classically described, the properties of rapid activation and inactivation position I_{AHP} as a critical regulator of interspike interval and firing frequency (Connor and Stevens, 1971a,b). In addition, in some neurons, I_{AHP} has been shown to contribute to resting, subthreshold K^+ conductance, influencing input resistances, and excitability thresholds (de Jeu et al., 2002; Yuan et al., 2005). These unique properties, together with variable dendritic expression patterns, allow I_{AHP} channels to play important roles in modulating the responses to synaptic inputs and to influence synaptic integration and neuronal output properties (Birnbaum et al., 2004; Jerng et al., 2004). In spontaneously active neurons, like those in the SCN, therefore, I_{AHP} channels would be expected to function to regulate excitability and the initiation of firing by opposing membrane depolarizations, resulting from the closing/opening of other channels, as well as influence the voltage trajectory during the interspike interval, which will regulate repetitive firing rates (Connor and Stevens, 1971b; Kang et al., 2000; Kim et al., 2005; Yuan et al., 2005; Khaliq and Bean, 2008). Although I_{AHP} was previously hypothesized to be critical in regulating the transition from the silent night state to the spontaneously active day state (Kim and Dudek, 1993; Bouskila and Dudek, 1995), the results presented here demonstrate that Kv1.4- and Kv4.2-encoded I_{AHP} channels are instead critical in determining the threshold for firing of SCN neurons during the day and during the night. Other, yet to be identified subthreshold K^+ channels, therefore, must be involved in determining the transitions between the night and day patterns of electrical activity in the SCN.

The role of I_{AHP} in setting the circadian period of rhythmic firing seems likely to be intrinsic to individual SCN neurons. The experiments here demonstrated that dispersed Kv1.4⁻/⁻ and Kv4.2⁻/⁻ SCN neurons remain synchronized in the timing of daily maximum and minimum firing rates. In addition, Kv1.4⁻/⁻ and Kv4.2⁻/⁻ SCN neurons display a shortened circadian period in vitro that is very similar to the shortened period observed in locomotor behavior in Kv1.4⁻/⁻ and Kv4.2⁻/⁻ mice. The change in period observed at the cellular level supports the hypothesis that membrane excitability likely is an integral part of the core circadian clock (Nitabach et al., 2002, 2005; Lundkvist and Block, 2005). We conclude that I_{AHP} functions to reduce the intrinsic excitability of SCN neurons, thereby contributing to the lengthening of circadian periods of firing and behavior.

The results presented here also hint at a novel role for Kv1.4 outside of the SCN. The increased duration of daily locomotion of mice lacking Kv1.4, for example, could reflect alterations in sleep–wake cycles, specifically reductions in daily sleep duration. Because the durations in daily firing patterns in SCN neurons deficient for Kv1.4 were not significantly different from WT SCN neurons, the behavioral phenotype may result from the functioning of Kv1.4-encoded I_{AHP} channels outside the SCN. Together with previous studies linking Kv channel functioning to sleep in mice and in flies (Vyazovskiy et al., 2002; Cirelli et al., 2005; Douglas et al., 2007; Espinosa et al., 2008; Koh et al., 2008), the results here support the interesting possibility that modulation of Kv channels (and other conductances that regulate intrinsic neuronal excitability) may play critical roles in both circadian biology and sleep.

In summary, these results demonstrate that, in the SCN, Kv1.4 and Kv4.2, but not Kv4.3, are critical subunits for the generation of rapidly activating and inactivating K⁺ currents that regulate the excitability, firing rate, and, consequently, the circadian period of firing and locomotor behavior.

References


