Phosphorylation of a constitutive serine inhibits BK channel variants containing the alternate exon "SRKR"

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BK Ca\(^{2+}\)-activated K\(^+\) currents exhibit diverse properties across tissues. The functional variation in voltage- and Ca\(^{2+}\)-dependent gating underlying this diversity arises from multiple mechanisms, including alternate splicing of Kenmama1, the gene encoding the pore-forming (α) subunit of the BK channel, phosphorylation of α subunits, and inclusion of β subunits in channel complexes. To address the interplay of these mechanisms in the regulation of BK currents, two native splice variants, BK\(_{\text{const}}\) and BK\(_{\text{SRKR}}\), were cloned from a tissue that exhibits dynamic daily expression of BK channel, the central circadian pacemaker in the suprachiasmatic nucleus (SCN) of mouse hypothalamus. The BK\(_{\text{const}}\) and BK\(_{\text{SRKR}}\) variants differed by the inclusion of a four–amino acid alternate exon at splice site 1 (SRKR), which showed increased expression during the day. The functional properties of the variants were investigated in HEK293 cells using standard voltage-clamp protocols. Compared with BK\(_{\text{const}}\), BK\(_{\text{SRKR}}\) currents had a significantly right-shifted conductance–voltage (G–V) relationship across a range of Ca\(^{2+}\) concentrations, slower activation, and faster deactivation. These effects were dependent on the phosphorylation state of S642, a serine residue within the constitutive exon immediately preceding the SRKR insert. Coexpression of the neuronal β4 subunit slowed gating kinetics and shifted the G–V relationship in a Ca\(^{2+}\)-dependent manner, enhancing the functional differences between the variants. Next, using native action potential (AP) command waveforms recorded from SCN to elicit BK currents, we found that these splice variant differences persist under dynamic activation conditions in physiological ionic concentrations. AP-induced currents from BK\(_{\text{SRKR}}\) channels were significantly reduced compared with BK\(_{\text{const}}\), an effect that was maintained with coexpression of the β4 subunit but abolished by the mutation of S642. These results demonstrate a novel mechanism for reducing BK current activation under reconstituted physiological conditions, and further suggest that S642 is selectively phosphorylated in the presence of SRKR.

INTRODUCTION

The BK large-conductance calcium- and voltage-activated potassium channel (K\(_{\text{Ca1.1}},\) MaxiK) is gated by both membrane depolarization and intracellular Ca\(^{2+}\) (Butler et al., 1993). BK channels are tetramers of four pore-forming α subunits (Shen et al., 1994), encoded by the Kenmama1 (Slo1) gene (Butler et al., 1993; Meredith et al., 2004). BK α subunits are comprised of seven transmembrane domains (S0–S6) that form the channel pore and are responsible for voltage sensing and a cytoplasmic C-terminal domain that binds Ca\(^{2+}\) (Yuan et al., 2012). BK channels are expressed in a wide variety of excitatory and nonexcitable cells (Tseng-Crank et al., 1994) where they have heterogeneous physiological functions. In neurons, BK channels contribute to the fast afterhyperpolarization of the action potential (AP) (Storm, 1987), AP duration and frequency (Storm, 1987; Montgomery et al., 2013), and neurotransmitter release (Robitaille and Charlton, 1992; Hu et al., 2001). In smooth muscle, BK currents limit contraction and promote relaxation (Brayden and Nelson, 1992; Brenner et al., 2000b). In nonexcitable cells, BK channels regulate K\(^+\) homeostasis (Rieg et al., 2007) and hormone secretion (Petersen and Maruyama, 1984). Based on this wide array of cellular mechanisms, the BK channel has been implicated in the physiology of hearing (Fuchs and Evans, 1990; Pyott et al., 2007), circadian rhythmicity (Meredith et al., 2006), cardiovascular function (Sausbier et al., 2005), urination (Meredith et al., 2004), and locomotor function (Sausbier et al., 2004; Imlach et al., 2010).

Unlike many other K\(^+\) channels, the BK channel tetramer is encoded by a single gene. Given this, physiological diversity of BK currents is achieved through several mechanisms including formation of macromolecular complexes with auxiliary subunits (Orio et al., 2002; Yan and Aldrich, 2012), posttranslational modification of α subunits (Tian et al., 2008; Yan et al., 2008), and alternative splicing (Shipston, 2001). Alternative splicing of Kenmama1 transcripts has been proposed as a major mechanism driving the diversity of BK currents. Within the cytoplasmic C-terminal domain, four splice sites...
have been identified. Site 1 is near the end of the first regulator of K+ conductance (RCK) domain, containing either no insert or a four–amino acid alternate exon (SRKR) (Tseng-Crank et al., 1994). Site 2 is the stress axis–regulated exon (STREX) site, found within the linker between RCK1 and RCK2. This site can contain either no insert, a three–amino acid exon (IYF), a 61–amino acid exon (STREX1), the IYF and 61–amino acid exons (STREX2), or a 28–amino acid exon (Saito et al., 1997; Xie and Cobb, 1998; Chen et al., 2005). Site 3 is located near the high affinity Ca2+–binding site (the “Ca2+–bowl”) within RCK2 and can contain either no insert or a 27–amino acid alternate exon (Ha et al., 2000). Lastly, alternate exons at site 4 produce three distinct C termini (“VEDEC,” “VYR,” or “QEERL”) (Saito et al., 1997).

Functionally, inclusion of alternate exons at each site produces effects on current properties that range in magnitude and mechanism, with the STREX site (site 2) being the most widely characterized. Inclusion of STREX1 or STREX2 enhances BK currents by speeding activation, slowing deactivation, and shifting the voltage dependence of activation more negative compared with insertless variants (Xie and Cobb, 1998; Chen et al., 2005). Depending on the Ca2+ concentration and the background splice variant sequence, inclusion of STREX can left-shift the V1/2 up to 50 mV (Chen et al., 2005). Conversely, inclusion of the SRKR insert at site 1 was found to reduce BK currents by either right-shifting the V1/2 (Tseng-Crank et al., 1994) or by reducing voltage sensitivity (Rosenblatt et al., 1997). Inclusion of the Ca2+–bowl exon at site 3 does not have an appreciable effect on steady-state current properties; however, it does speed activation kinetics in a Ca2+-dependent manner (Ha et al., 2000). Finally, the C-terminal exons so far appear to regulate trafficking and cell surface localization for the channel rather than significantly affecting current properties (Kim et al., 2008; Singh et al., 2013).

In addition to alternate splicing, posttranslational modifications, such as phosphorylation, have also been shown to underlie some of the physiological diversity in BK currents. The BK channel has ~200 potentially phosphorylated serine and threonine residues. However, recent mass spectrometric analysis of rat brain BK protein detected only ~30 endogenously phosphorylated residues (Yan et al., 2008). Some of these phosphorylated residues were found within alternative exons at sites 2, 3, and 4 (Yan et al., 2008), suggesting that the combinatorial effects of exon-specific phosphorylation could further contribute to BK current properties. Consistent with this, inclusion of the STREX insert (site 2) changes the phosphorylation dependence of activation. Phosphorylation of S869 in channels lacking the STREX insert increases channel activity, whereas phosphorylation of a serine within the STREX exon (S4STREX) inhibits channel activity (Tian et al., 2001). Although the phosphorylation dependence of STREX has been the most comprehensively studied, the functional impact of phosphorylation at other alternate exons has not been addressed.

In this study, we cloned and characterized two novel Kenneth variants (BK0 and BKSRKR) from the mouse suprachiasmatic nucleus (SCN) of the hypothalamus, the brain’s circadian clock. We chose to identify BK channel sequences from SCN because (a) BK currents undergo dynamic transcriptional regulation in this tissue, suggesting that multiple splice variants may be expressed (Panda et al., 2002; Pitts et al., 2006); (b) distinct AP waveforms have been characterized in SCN neurons that are hypothesized to exert differential influence on intrinsic currents (Colwell, 2011; Montgomery and Meredith, 2012); and (c) SCN expresses BK modulatory β subunits (Montgomery and Meredith, 2012). The BK0 and BKSRKR variants were identical except for the presence of a four–amino acid insert (SRKR) at splice site 1. To determine the functional effects of the SRKR insert, we expressed each variant with and without the β4 accessory subunit in HEK293 cells, eliciting currents in response to standard voltage jumps and representative “day” and “night” SCN AP waveform commands. Next, using a combination of phosphatase treatment and site-directed mutagenesis, we addressed the role of two serine residues, predicted to be phosphorylated in BKSRKR, in generating the distinct current properties produced by the BK0 and BKSRKR variants. Lastly, using SCN AP commands and expression of β4 to reconstitute activation of BK currents under physiologically relevant conditions, we found that the differences in BK currents were expressed in response to the type of AP command, the splice variant expressed, the presence of the β4 subunit, and the phosphorylation status of S642.

**MATERIALS AND METHODS**

Cloning BK0 and BKSRKR variants from SCN

Total RNA was isolated from 10 pooled mouse SCNs harvested at zeitgeber time (ZT)6 (6 h after lights on), as described in Meredith et al. (2006). Total RNA was column purified (RNAeasy; Qiagen), and cDNA was synthesized from 10 µg of total RNA (Superscript II; Invitrogen) using a gene-specific reverse primer (5'-GGTGACCATCATTCTCCTCAAAG-3') and a common forward primer (5'-CATTCAAATCAAGCCCATGACCC-3') (Superscript II; Invitrogen). PCR amplifications were performed on 1 µl cDNA (Expand Taq; Roche). Products were gel purified, subcloned (pGEMTeasy; Promega), and sequenced to determine the presence of alternate exons at splice sites 1–4 (5’-GGAGACACTCATCTCTTCGACAAG-3’). PCR reactions amplifying the N-terminal (5’-ATGGCCTGGTGATGTTCTGGG-3’ and 5’-GGCCCAGGAAAGAAGTGTCACG-3’) or a single product containing splice sites 1–4 (5’-GAGTACAAGTCTGCCAACAG-3’ and 5’-CATTCAATATCGACCCCATGAGGTACC-3’) were performed on 10 µl cDNA (Expand Taq; Roche). Products were gel purified, subcloned (pGEMTeasy; Promega), and sequenced to determine the presence of alternate exons at splice sites 1–4. Two PCR products were obtained with unique cDNA sequences, BK0 and BKSRKR. A series of BK channel expression vectors containing the stepwise sequence additions and deletions of exons were introduced into mbr5 (Butler et al., 1993) in pcDNA3.1. In this study, mbr5 contained an N-terminal myc tag and EYFP (716 bp) inserted in the RCK2 domain at nucleotide position (nt) 2032 (GenBank...
accession no. KF530038; provided by R. Brenner, The University of Texas at San Antonio, San Antonio, TX). Construct 2 (C2; GenBank accession no. KF530044; Fig. S1) contained an insertion of an N-terminal myc tag followed by 195 bp of additional keratin 1 N-terminal sequence encoding the “MANG” alternate start site. The inserted sequence was synthesized as a minigene (Integrated DNA Technologies), digested with KpnI/AgeI, and subcloned into mbr5/pcDNA3.1 at KpnI and BoHII sites.

For all subsequent constructs, splice site insertion or deletions were performed using site-directed PCR mutagenesis (QuikChange; Agilent Technologies) according to the manufacturer’s protocol. For construct 3, three amino acids (IYF) were deleted from the STREX site by amplifying construct 2 with a primer containing a deletion of nts 2094–2102 (GenBank accession no. KF530036). For construct 4, 180 bp encoding the “VEDEC” C terminus was deleted by PCR from construct 3 (nts 4236–4415), and 21 bp encoding the “KRKMYR” C terminus was introduced in a second PCR amplification (GenBank accession no. KF530042). BK<sub>K</sub> was generated by the addition of 78 bp encoding the 27–amino acid Ca<sup>2+</sup>-bowl exon at nt 3564 in construct 4 (GenBank accession no. KF530040). BK<sub>S642A</sub> was generated by the addition of 12 bp encoding the SRKR exon at nt 1929 (GenBank accession no. KF530041).

**Quantitative PCR**

Total RNA was isolated from SCN tissue at 6 h after lights on (ZT6; “day”) and 7 h after lights off (ZT19; “night”) with the RNeasy mini kit (QiAGEN) as described previously (Montgomery and Meredith, 2012). Primers were as follows: SRKR, (F) 5’-GTTT-TTGAAGCTTAAGCTCCTGAT-3’ and (R) 5’-AGAGCCGAAG-CGGAAAGC-3’; exon 16, (F) 5’-GTTTGTAGAATGCTGTC-CTGAT-3’ and 5’-CTCTCTGTGGCAGACTTGTAC-3’. Real-time qPCR data were analyzed using the 2<sup>-ΔΔCT</sup> method (Livak and Schmittgen, 2001). SRKR and exon 16 Ct values for day were normalized to the night values.

**Predicted phosphorylation sites and mutagenesis of BK<sub>K</sub> and BK<sub>S642A</sub>**

Sequence analysis for potential phosphorylation sites was performed online using NetPhos 2.0 (Blom et al., 1999). Predicted consensus phosphorylation sites received a score >0.9.

Mutation of predicted phosphorylation sites S642 and S644 was performed using site-directed PCR mutagenesis (QuickChange; Agilent Technologies). BK<sub>K</sub>S642D contained the mutation AGC to GCC at nts 1929–1931 (GenBank accession no. KF530040). BK<sub>S642A</sub>S644A contained the mutation AGC to GCC at nts 1929–1935 (GenBank accession no. KF530041). All mutations were verified by sequencing.

**HEK cell patch-clamp electrophysiology**

HEK293T cells were transfected with BK expression constructs, with or without β4 in pcDNA3.1 (Wang et al., 2009). 1 μg DNA/35-mm dish was transfected for a subunits only, and 1 μg α plus 1.7 μg β4 was transfected for cotransfections, using Lipofectamine 2000 (Invitrogen). Cells were plated on glass coverslips 3–5 h later and recorded from 20 to 30 h after transfection. Patch-clamp recording was performed using the voltage-clamp mode in the inside-out patch configuration (Axopatch; Molecular Devices) using thin-walled borosilicate pipettes with resistances of 2–5 MΩ. Recordings were made at room temperature, and the data were acquired at 50 kHz and filtered at 10 kHz. Total [Ca<sup>2+</sup>], was calculated using Webmax C Standard software and the appropriate addition of CaCl<sub>2</sub> to achieve the indicated free Ca<sup>2+</sup>. For 0 Ca<sup>2+</sup> solutions, 5 mM EGTA was used, where free Ca<sup>2+</sup> is expected to be <1 nM under these conditions (Bao et al., 2002). For 100-μM Ca<sup>2+</sup> solutions, no EGTA was used. For inside-out patch recordings, the pipette (extracellular) solution was composed of (mM): 140 KMeSO<sub>3</sub>, 2 KCl, 2 MgCl<sub>2</sub>, and 20 HEPES, pH 7.3. The bath (intracellular) solution was composed of (mM): 140 KMeSO<sub>3</sub>, 2 KCl, 20 HEPES, and 5 HEDTA for 1 and 10 μM free Ca<sup>2+</sup>. To elicit BK currents using voltage-jump protocols, patches were stepped from holding potentials of −100 to −150 to +350 mV (in 10-mV increments) for 20 ms (α alone) or 50 ms (α + β4), stepping back to −90 mV for 10 ms (α alone) or 20 ms (α + β4) to generate tail currents. Conductance (G) was calculated from tail currents, normalized to the highest conductance calculated for each patch, and plotted against membrane potential to generate G-V curves. Boltzmann functions were fitted to curves in Origin 8.5 (OriginLab) using the equation: \( y = \frac{(A_1 - A_2)}{(1 + e^{\Delta \phi - \phi \Delta / kT})} + A_2 \). Single-exponential functions were fitted to current traces in pCLAMP (Molecular Devices) to determine the time constants of activation, deactivation, and inactivation, using the equation \( f(x) = A e^{-(x-t)/\tau} + C \). For dephosphorylation experiments, calf intestinal alkaline phosphatase (New England Biolabs, Inc.) was diluted to 10 U/ml in the intracellular bath solution. After recording baseline currents, phosphatase-containing solution (Alk P) was perfused into the bath, and posttreatment currents were recorded 5 min later.

**Activation of BK currents by AP commands**

AP command waveforms were originally recorded in whole-cell current-clamp mode from acute SCN slices during the day and night of the circadian cycle as described previously (Montgomery and Meredith, 2012). Day and night waveforms representative of the mean parameter values (see Results and Fig. 2A) were used to elicit BK currents in whole-cell recording mode. The pipette (intracellular) solution was identical to the intracellular solution used in inside-out recordings, except with free Ca<sup>2+</sup> adjusted to 50 μM. The bath (extracellular) solution consisted of (mM): 145 NaCl, 5 KCl, 2 CaCl<sub>2</sub>, 1 MgCl<sub>2</sub>, and 10 HEPES, with pH adjusted to 7.2 with NaOH. Cells were held at −100 mV. The applied voltage stimulation protocol consisted of a depolarizing prepulse to +160 mV for 20 ms, followed by either day or night physiological AP commands that included holding the cells at the appropriate physiological resting membrane potentials (−48.6 mV for day AP commands and −45.0 mV for night AP commands) for 50 ms before the actual AP commands (Fig. S3 A). Series resistance was compensated up to 60%.

For comparisons between splice variants, peak AP-elicited currents were normalized to the +160-mV prepulse (Fig. S3 B). For AP-induced BK currents, peak current was the difference between the baseline current at −100 mV and the maximum current observed in response to the AP command. Current half-width was the duration at 50% of the maximal current magnitude, measured between the current maximum and the afterhyperpolarization current. Charge transfer was calculated as the area of the normalized current from baseline (at resting membrane potential) to the maximum of the AP-induced current.

**Statistics**

Two-way ANOVAs with Bonferroni post-hoc analysis were used to determine statistical significance for differences between \( V_{1/2} \) values and day versus night AP-induced current parameters for the BK variants, as indicated in the figure legends. One-way ANOVA was used for comparisons between wild-type and mutant AP-elicited currents (Fig. 4 H), and \( t \) test was used to compare currents elicited by square voltage pulses in Fig. 4 G. Factorial ANOVAs with Bonferroni post-hoc analyses were used to determine statistical significance for \( \tau_{\text{act}} \) and \( \tau_{\text{deact}} \) between constructs elicited at different Ca<sup>2+</sup> across a range of voltages. Statistical significance was achieved if \( P \leq 0.05 \). Mean values in figures are given as ±SEM.
SRKR alters kinetic and steady-state BK current properties

Given the difference in abundance of SRKR-containing transcripts in day and night SCN tissue from the normal Kcnma1 expression pattern, we looked at the functional impact of this exon on BK currents. BK0 and BKSRKR were expressed in HEK293 cells, and macroscopic BK currents were recorded from inside-out patches in symmetrical K+ at 0, 1, 10, and 100 µM of intracellular Ca2+. Expression of both variants produced BK currents that activated with increasing depolarization and Ca2+ (Fig. 2 A). Differences in current properties were apparent at 0, 1, and 100 µM, where the BKSRKR G-V relationship was right-shifted compared with BK0, but not at 10 µM Ca2+ (Fig. 2 B). At 100 µM Ca2+, a condition where all of the high affinity Ca2+-binding sites are expected to be occupied, BKSRKR currents were harder to activate relative to BK0, with the V1/2 right-shifted by 19 mV compared with BK0 (Fig. 2 C). This effect was also seen at 1 µM Ca2+ (+35-mV right-shift), as well as in the absence of Ca2+.

Online supplemental material

Fig. S1 demonstrates that the addition of the alternate N terminus, deletion of residues IVF at site 2, and the addition of the alternate C terminus (RKEMVYR) have little effect on BK current properties. Fig. S2 demonstrates the functional differences produced by the addition of the Ca2+-bowl exon at site 3 (BKc3). Fig. S3 shows the prepulse voltage step used to normalize the AP-induced BK currents in relation to the AP voltage command and the BK currents elicited by each command to scale. Fig. S4 compares the amino acid sequences and secondary structures in the SRKR region from the two isoforms of the BK channel that have been crystallized to date. Table S1 contains the V1/2 values of all the constructs used in this study. The online supplemental material is available at http://www.jgp.org/cgi/content/full/jgp.201311072/DC1.

RESULTS

Identification of two novel BK channel splice variants

To investigate BK current properties produced by native BK channels, we amplified full-length Kcnma1 cDNAs from SCN, a tissue that exhibits dynamic expression of Kcnma1 transcripts (Panda et al., 2002; Pitts et al., 2006). Two novel variants were identified (BK0 and BKSRKR; Fig. 1 A), containing an alternate exon combination that differed at four splice sites from previously identified BK variants. Both variants contained an alternative translation start encoded by additional sequence at the extracellular N terminus of the channel (“MANG”; Fig. 1 A). In addition, both variants lacked any insert at the STREX site (site 2), contained the 27–amino acid Ca2+-bowl insert (site 3), and contained the same alternative C terminus, RKEMVYR (site 4). However, the variants differed from each other at splice site 1, located within RCK1, where BKSRKR contains a four–amino acid insert, SRKR, that is absent in BK0. Stepwise addition of each alternate sequence into a BK channel lacking inserts at splice sites 1 and 3, and replacements of exons at sites 2 and 4, revealed that the addition of the SRKR insert had the greatest impact on BK current properties (Figs. S1 and S2).

To determine whether the expression of BK transcripts containing the SRKR exon differed from the previously characterized nighttime-phased expression pattern of Kcnma1 in SCN (Meredith et al., 2006; Montgomery and Meredith, 2012), exon 16 (the constitutive exon immediately preceding SRKR) and SRKR levels were determined by quantitative RT-PCR from day and night SCNs (Fig. 1 B). The abundance of transcripts containing exon 16 was greater at night compared with day (Fig. 1 B), similar to other constitutive Kcnma1 exons (Panda et al., 2002; Pitts et al., 2006; not depicted). In contrast, the abundance of transcripts containing SRKR was higher during the day (Fig. 1 B), suggesting that SRKR-containing transcripts form a distinct type of BK channel that is regulated differentially with respect to circadian phase.
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at 10-µM Ca\(^{2+}\) concentrations (Fig. 2 F). Overall, at saturating (100 µM) Ca\(^{2+}\), the right-shifted V\(_{1/2}\) increased activation time constants, and decreased deactivation time constants would be expected to reduce native BK\(_{SRKR}\) currents compared with BK\(_{0}\). The same effect is expected at low Ca\(^{2+}\) (0 and 1 µM), where BK\(_{SRKR}\) currents were similarly right-shifted and slower to activate compared with BK\(_{0}\), but absent at intermediate Ca\(^{2+}\) (10 µM), where no significant differences in current properties were observed.

SRKR is required for phosphoregulation of serine 642

BK\(_{0}\) and BK\(_{SRKR}\) splice variants differ by only four amino acids, yet they have substantial differences in both steady-state and kinetic properties. Given the presence of a serine within the SRKR insert, we hypothesized that phosphorylation of this residue (S644; Fig. S4) could contribute to the difference in BK\(_{0}\) and BK\(_{SRKR}\) current properties. Analysis of SRKR and the surrounding sequence in BK\(_{SRKR}\) with a phosphorylation prediction (+49-mV right-shift). There was no change in the slope factor (voltage sensitivity) between BK\(_{0}\) and BK\(_{SRKR}\) G-Vs at each Ca\(^{2+}\) concentration. These data show that inclusion of the SRKR exon shifts the voltage dependence of activation to more positive values across a range of membrane potentials. Surprisingly, there was no significant difference in the V\(_{1/2}\) values of BK\(_{SRKR}\) and BK\(_{0}\) G-Vs at the intermediate Ca\(^{2+}\) concentration of 10 µM (Fig. 2, B and C).

There were also differences in the kinetics of activation and deactivation between BK\(_{0}\) and BK\(_{SRKR}\) currents. At 0, 1, and 100 µM Ca\(^{2+}\), the time constants of activation (τ\(_{act}\)) for BK\(_{SRKR}\) currents were increased compared with BK\(_{0}\) (Fig. 2 D). This slowing of activation was particularly pronounced at 100 µM Ca\(^{2+}\) at more negative voltages (Fig. 2 E). As with the steady-state G-V data, there was no difference in BK\(_{0}\) and BK\(_{SRKR}\) current activation time constants in the intermediate Ca\(^{2+}\) concentration of 10 µM. BK\(_{SRKR}\) currents showed decreased τ\(_{deact}\) at 100, 1, and 0 µM Ca\(^{2+}\) with no significant effect at 10-µM Ca\(^{2+}\) concentrations (Fig. 2 F). Overall, at saturating (100 µM) Ca\(^{2+}\), the right-shifted V\(_{1/2}\) increased activation time constants, and decreased deactivation time constants would be expected to reduce native BK\(_{SRKR}\) currents compared with BK\(_{0}\). The same effect is expected at low Ca\(^{2+}\) (0 and 1 µM), where BK\(_{SRKR}\) currents were similarly right-shifted and slower to activate compared with BK\(_{0}\), but absent at intermediate Ca\(^{2+}\) (10 µM), where no significant differences in current properties were observed.

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Functional effects of SRKR exon on BK currents

algorithm (Blom et al., 1999) indicated a high probability for S644 phosphorylation (NetPhos score of 0.991). S642, a constitutive residue preceding the SRKR insert (Fig. S4), also had a high probability score for phosphorylation (NetPhos score of 0.994). In contrast, in the BK0 sequence, which lacked SRKR, the constitutive S642 had a significantly lower probability of phosphorylation (0.557). These results predicted that phosphorylation of S642 and/or S644 may occur in BKSRKR, but not in BK0, channels.

First, to determine if net phosphorylation of BKSRKR could explain the differences in current properties compared with BK0, we applied alkaline phosphatase to the intracellular side of macropatches. At 100 µM Ca2+, the BKSRKR G-V curve was right-shifted compared with BK0. In the presence of phosphatase, the BKSRKR V1/2 was left-shifted by 14 mV, making it similar to that of BK0 and significantly different from that of BKSRKR (Fig. 3, A and B). This result suggested that dephosphorylation increased BKSRKR channel activity to match that of BK0. In contrast, the application of phosphatase had no effect on BK0 currents (Fig. 3, A and B), confirming that the SRKR insert is required to render the currents susceptible to phosphatase. These experimental results are consistent with the high phosphorylation probability

![Figure 3. BK0 and BK SRKR current properties in patches treated with phosphatase or from phospho-site mutants. (A) G-V relationship of BK0 and BKSRKR at 100 µM Ca2+ before and after the application of calf intestinal alkaline phosphatase (Alk P). Alk P caused a leftward shift of the BKSRKR G-V, but not of BK0 G-V. (B) Plot of V1/2 versus Ca2+, illustrating that dephosphorylation of BKSRKR produces a BK0-like phenotype in the voltage dependence of activation. After treatment with phosphatase, the V1/2 was significantly left-shifted with BKSRKR at 1 µM Ca2+ (204–155 mV; P = 0.01; n = 8; two-way ANOVA) and at 100 µM Ca2+ (8 to 6 mV; P = 0.03; n = 8; two-way ANOVA), but not with BK0 at either 1 µM Ca2+ (169–156 mV; P = 0.25; n = 8; two-way ANOVA) or 100 µM Ca2+ (11 vs. 7 mV; P = 0.53; n = 8; two-way ANOVA), making the phosphatase-treated BKSRKR not significantly different from BK0 (P = 0.7; n = 8). Symbols are as in A. (C) G-V relationship at 100 µM Ca2+ of specific phospho-mutants used to determine the site of differential phospho-regulation of BKSRKR. Phospho-mutant BKSRKR S644A, the S in SRKR, had no effect on G-V relationship, whereas phospho-mimetic BKSRKR S642D, found within a constitutive portion of the channel, was found to be necessary and sufficient to induce that change in current properties. (D) Plot of V1/2 versus Ca2+ for BK constructs shown in C. At 1 and 100 µM Ca2+, the V1/2 values cluster into two groups, with those of BKSRKR, BKSRKR-S644A, and BK0-S642D further right-shifted than those of BK0, BKSRKR-S642A, and BKSRKR-S642A/S644A. Symbols are as in C. At 1 µM Ca2+, there were no significant differences in the V1/2 values between BKSRKR and BKSRKR-S644A (204 and 206 mV; P = 0.29; n = 8–12; two-way ANOVA), between BKSRKR-S642A and BK0 (166 and 169 mV; P = 0.29; n = 8–10; two-way ANOVA), between BKSRKR-S644A and BK0 (167 vs. 169 mV; P = 0.25; n = 8–15; two-way ANOVA). At 100 µM Ca2+, there were no significant differences in the V1/2 values between BKSRKR and BKSRKR-S644A (6 and 8 mV; P = 0.3; n = 8–12; two-way ANOVA), between BKSRKR-S642A and BK0 (V1/2 values both −11 mV; P = 0.6; n = 8; two-way ANOVA), between BKSRKR-S644A and BK0 (11 vs. 8 mV; P = 0.4; n = 8; two-way ANOVA), or between BKSRKR-S642A/S644A and BK0 (−15 vs. −11 mV; P = 0.6; n = 8; two-way ANOVA). (E and F) Plots of τact (E) and τdeact (F) versus voltage for BK0, BKSRKR, and BKSRKR-S642A at 100 µM Ca2+. Activation kinetics of BKSRKR-S642A were not significantly different to those of BK0 (P = 0.34; n = 8) but were significantly slowed compared with BKSRKR (P = 0.02; n = 8–9; factorial ANOVA). Similarly, deactivation kinetics of BKSRKR-S642A were not significantly different to those of BK0 (P = 0.49; n = 8; factorial ANOVA) but were significantly slowed compared with BKSRKR (P < 0.01; n = 8–9; factorial ANOVA).
score for the BKSRKR variant and low probability score for the BK0 variant.

Next, to determine if a serine residue was required within the SRKR insert, we mutated S644 to an alanine. We hypothesized that currents produced from BKSRKR-S644A mutants would resemble both BK0 and the phosphatase-treated BKSRKR currents. Surprisingly, BKSRKR-S644A showed no difference in the G-V relationship compared with wild-type BKSRKR (Fig. 3, C and D). Furthermore, phosphatase treatment of BKSRKR-S644A still left-shifted the G-V curve to match that of BK0 (not depicted). Collectively, these results suggested that dephosphorylation is occurring at a residue other than S644 to produce the observed functional effects.

Next, we focused on S642, a constitutive serine present in both BK0 and BKSRKR variants. We hypothesized that in the absence of SRKR, S642 would be resistant to phosphorylation. Conversely, the presence of the SRKR insert may render S642 permissive to phosphorylation, leading to the observed right-shifted G-V curve for BKSRKR currents. To test this prediction, S642 was mutated to alanine. Confirming the role of this serine in the functional current differences, the G-V curve of BKSRKR-S642A matched that of BK0 (Fig. 3, C and D). Furthermore, the introduction of the converse phosphomimetic mutation into BK0, BK0-S642D, produced a right-shifted G-V curve that matched that of BKSRKR (Fig. 3, C and D). These results identify S642 as the critical residue underlying the differences in BKSRKR current properties compared with BK0 and suggest that phosphorylation of S642 decreases channel activity.

Mutating both serine residues to alanine in BKSRKR (BKSRKR-S642A/S644A) resulted in G-V curves that matched those of BK0 (Fig. 3, C and D), and no further left-shift of the G-V was produced by phosphatase treatment of this mutant (not depicted). These data confirm that no additional phosphorylation sites (sensitive to this phosphatase) were involved in the G-V differences between BK0 and BKSRKR. Quantitatively similar results were obtained when the phosphatase and mutagenesis experiments were repeated at 1 mM Ca2+ (Fig. 3 D). However, at 10 mM Ca2+, in which no functional differences in the currents produced by BK0 and BKSRKR were observed (Fig. 2), neither phosphatase treatment nor mutagenesis altered the G-V curves (Fig. 3, B and D).

In addition to steady-state current properties, mutation of S642 also impacted current kinetics. We found that the time constants of both activation and deactivation of BKSRKR-S642A were similar to those of BK0 (Fig. 3, E and F). Thus, both the steady-state and kinetics properties of BKSRKR lacking the S642 phosphorylation site resemble those of BK0.

SRKR reduces BK currents elicited by AP commands

Much of the previously published work on BK current properties produced by specific variants has examined currents in response to standard square voltage-pulse protocols. Few studies have addressed the properties of BK currents elicited by a more physiologically relevant stimulus, such as an AP. Therefore, we recorded BK currents elicited from two distinct types of AP waveforms (day and night), based on recordings from SCN neurons where BK0 and BKSRKR are endogenously expressed. These experiments examined whether splice variant–based differences in BK currents were still apparent in response to physiologically relevant activation. We also examined currents elicited under these conditions co-expressed with β4, the major neuronal auxiliary subunit, which is known to be expressed within the SCN (Montgomery and Meredith, 2012).

The representative day and night AP waveforms used as a voltage command for BK current activation were obtained from whole-cell recordings of spontaneously firing neurons during the peak (day) and trough (night) of neural activity (Montgomery and Meredith, 2012). Day and night AP waveforms differed significantly in several parameters (Montgomery and Meredith, 2012), including peak depolarization (14.9 mV for day compared with −2.2 mV for night), half-width (2.1 vs. 3.0 ms, respectively), and baseline membrane potential (−48.6 vs. −45.0 mV, respectively). These distinct AP properties were expected to exert a differential drive on BK activation, conditions that might alter the expression of the differences observed under square voltage-jump conditions between BK0 and BKSRKR currents.

Day and night AP command waveforms (Fig. 4 A) were used to elicit currents in physiological ion concentrations (low internal K+ and high external Na+). Fig. 4 B shows AP-elicited currents from BK0 and BKSRKR, elicited from day or night AP command waveforms at 50 µM Ca2+, mimicking the high local Ca2+ concentration experienced by BK channels in vivo coupled with voltage-gated Ca2+ channels (Fakler and Adelman, 2008). In this configuration, the time course of the BK currents generally followed the depolarizing phase of the AP command waveforms, mirroring the differences seen between the day and night AP properties. To directly compare BK0 and BKSRKR properties across cells expressing different levels of each channel variant, the AP-elicited currents were normalized to a maximally activating depolarizing prepulse step (Fig. S3). The mean AP-induced peak currents from both variants were very low, ranging from 2 to 8% of maximal current, depending on the variant and AP command waveform. Both BK0 and BKSRKR currents elicited by day AP commands had significantly increased peak current amplitudes and decreased half-widths compared with currents elicited by night AP commands (Fig. 4, D and F). There was no significant difference in charge transfer elicited by day and night AP commands (Fig. 4, E and F). These data show that the day and night AP command waveforms produced differing drives on BK current activation under standardized conditions.

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Next, we compared the properties of BK₀ and BKSRKR currents to each other, using either day or night AP commands (Fig. 4 C). BK₀ and BKSRKR currents elicited by the same stimulus, either the day or night AP command waveform, showed different properties (Fig. 4 C). BKSRKR had significantly decreased peak currents (Fig. 4 D), decreased charge transfer (Fig. 4 E), and increased current half-width (Fig. 4 F) compared with BK₀. These results are consistent with our findings using standard voltage jumps (Fig. 2 B). The reduced BKSRKR currents compared with BK₀ elicited by AP commands could be caused by differences in the steady-state properties of the channels, kinetic properties, or a combination of both. To determine whether reduced steady-state activation could explain the differences between BK₀ and BKSRKR AP-induced currents, we used a square pulse of 40 ms to depolarize cells to +14.9 mV, the peak voltage achieved using the day AP command. We found no significant difference between BK₀ and BKSRKR currents under these conditions (Fig. 4 G), indicating that differences in steady-state parameters are not responsible for the distinct AP-induced BK₀ and BKSRKR current properties. This suggests that kinetic differences account for the major differences in currents produced by the two variants elicited by AP commands.

Because S642 is critical for expression of the differences between BK₀ and BKSRKR currents under standard voltage-jump protocols, we next sought to determine whether removal of the phosphorylation site S642 in the BKSRKR variant would render channels “BK₀-like” in their responses to AP command waveforms. BKSRKR-S642A currents elicited by the day AP command were next compared with BK₀, BKSRKR-S642A, and BKSRKR currents elicited by day or night AP commands (Fig. 4 C). BK₀ and BKSRKR currents elicited by the same stimulus, either the day or night AP command waveform, showed different properties (Fig. 4 C). BKSRKR had significantly decreased peak currents (Fig. 4 D), decreased charge transfer (Fig. 4 E), and increased current half-width (Fig. 4 F) compared with BK₀. These results are consistent with our findings using standard voltage jumps (Fig. 2 B). The reduced BKSRKR currents compared with BK₀ elicited by AP commands could be caused by differences in the steady-state properties of the channels, kinetic properties, or a combination of both. To determine whether reduced steady-state activation could explain the differences between BK₀ and BKSRKR AP-induced currents, we used a square pulse of 40 ms to depolarize cells to +14.9 mV, the peak voltage achieved using the day AP command. We found no significant difference between BK₀ and BKSRKR currents under these conditions (Fig. 4 G), indicating that differences in steady-state parameters are not responsible for the distinct AP-induced BK₀ and BKSRKR current properties. This suggests that kinetic differences account for the major differences in currents produced by the two variants elicited by AP commands.

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significantly larger than wild-type BKSRKR currents, and were similar to BK0 peak currents (Fig. 4 H). This result suggests that phosphorylation of S642 in BKSRKR channels would be expected to alter BK currents activated during the AP, opening the possibility that the SRKR splice site could play a role in influencing neural activity.

β4 enhances the functional differences between BK0 and BKSRKR

In vivo, BK current properties are modified by association with accessory β subunits. In the SCN (Montgomery and Meredith, 2012) and the brain as a whole, β4 is the predominant BK modulatory subunit (Behrens et al., 2000; Brenner et al., 2000a). Coexpression of β4 with the mbr5 BK channel has been shown to exert complex effects, decreasing Ca2+ sensitivity at low Ca2+ and increasing Ca2+ sensitivity at high Ca2+, with little effect at 10 µM Ca2+ (Behrens et al., 2000; Brenner et al., 2000a; Lippiat et al., 2003; Wang et al., 2006). β4 also influences channel kinetics, slowing rates of both activation and deactivation (Behrens et al., 2000; Brenner et al., 2000a). To examine the influence of β4 on each splice variant, and to determine whether β4 affects the functional differences observed between the splice variants, we coexpressed β4 with either BK0 or BKSRKR. First, currents were recorded from inside-out patches in symmetrical K+ solutions with standard square voltage jumps (Fig. 5 A), and G-V relationships and activation and deactivation kinetics were examined (Fig. 5, B–G). Similar to the previously reported effects on other BK channel variants, coexpression of β4 slowed activation of both BK0 and BKSRKR at both 10 and 100 µM Ca2+ (Fig. 5, A and B). At 10 µM Ca2+, β4 slowed the activation time constants of BK0 and BKSRKR to a comparable degree. Thus, the activation time constants at 10 µM Ca2+ were similar between the variants with and without β4 (Figs. 2 D and 5 B).

In contrast, at 100 µM Ca2+, although β4 slowed the activation of both BK0 and BKSRKR, the relative differences in activation kinetics between the variants in significantly larger than wild-type BKSRKR currents, and were similar to BK0 peak currents (Fig. 4 H). This result suggests that phosphorylation of S642 in BKSRKR channels would be expected to alter BK currents activated during the AP, opening the possibility that the SRKR splice site could play a role in influencing neural activity.

Figure 5. BK0 and BKSRKR currents from variants coexpressed with β4. (A) BK0 and BKSRKR currents were elicited with voltage steps from −100 to 200 mV with 10 µM Ca2+. Current traces were normalized to Imax elicited from either the α subunit alone (black) or coexpressed with β4 (gray traces). (B and C) Plot of τact (B) or τdeact (C) versus voltage for BK0 and BKSRKR coexpressed with β4 at 10 and 100 µM Ca2+. Symbols are as in C. Although no difference is found in kinetics between the two variants at 10 µM Ca2+, BKSRKR was significantly (P < 0.01; factorial ANOVA) slower to activate and faster to deactivate (P < 0.01) when coexpressed with β4 at 100 µM Ca2+. (D and E) G-V relationships of BK0 and BKSRKR coexpressed with β4 at 10 µM Ca2+ (D) and 100 µM Ca2+ (E). V1/2 values were significantly different between BK0 and BKSRKR at both 10 µM Ca2+ (151 vs. 94 mV; P < 0.01; n = 8; two-way ANOVA) and at 100 µM Ca2+ (15 vs. −23 mV; P < 0.01; n = 8; two-way ANOVA). (F) Plot of ΔV1/2 versus Ca2+ for each splice variant caused by β4 expression (V1/2[αβ] − V1/2[α]). Increasing Ca2+ reduces the V1/2 shift caused by β4 for both variants, albeit to differing extents. (G) Plot of ΔV1/2 versus Ca2+ between variants (V1/2[BKSRKR] − V1/2[BK0]). The addition of β4 increases the difference in V1/2 between splice variants.
presence of β4 were increased at positive voltages (Fig. 5 B) compared with activation kinetics in the absence of β4 (Fig. 2 E). The net effect was a larger divergence in activation time constants between the variants when β4 was coexpressed with 100 µM Ca²⁺.

β4 also slowed deactivation of both splice variants at 100 µM Ca²⁺, while still maintaining the faster deactivation of BKSRKR compared with BK₀ at more depolarized potentials (Fig. 5 C). In the absence of β4, differences in BK₀ and BKSRKR deactivation were most apparent at potentials hyperpolarized to approximately –100 mV (Fig. 2 B). However, in the presence of β4, increasing voltage caused a widening divergence of deactivation time constants between BK₀ and BKSRKR (Fig. 5 C). At 10 µM Ca²⁺, β4 also slowed deactivation of both variants approximately fourfold. As with activation time constants, the deactivation time constants of the two variants at 10 µM Ca²⁺ in the absence of β4 were similar (Fig. 2 F), and β4 coexpression did not introduce any further relative differences between BK₀ and BKSRKR.

Although β4 coexpression did not affect the differences in kinetics of activation or deactivation between the variants at 10 µM Ca²⁺, it did influence the G-V relationship. At 10 µM Ca²⁺, the presence of β4 right-shifted the V₁/₂ values for BK₀ and BKSRKR by 30 and 84 mV, respectively (Fig. 5, D and F, and Table S1). Furthermore, although in the absence of β4 at 10 µM Ca²⁺, no difference was seen between the G-V curves of BK₀ and BKSRKR (Fig. 2, B and E), coexpression with β4 now produced a 57-mV difference in V₁/₂ values (Fig. 5 G).

In contrast to 10 µM Ca²⁺, at 100 µM Ca²⁺, β4 coexpression was found to have much reduced, but opposing, effects on the G-V curves. For BK₀, the addition of β4 left-shifted the V₁/₂ by 12 mV; however, in the case of BKSRKR, β4 caused a 7-mV right-shift (Fig. 5, E and F, and Table S1). Coexpression of β4 increased the differences in V₁/₂ between the variants, although to a lesser extent than at 10 µM Ca²⁺ (Fig. 5 G). These results demonstrate that β4 coexpression acts to increase the functional differences between the steady-state properties of the splice variants at 10 and 100 µM Ca²⁺.

We next examined the consequence of β4 coexpression on the splice variant responses to day and night SCN AP waveforms. Whole-cell currents were recorded with physiological solutions (low internal K⁺ and high external Na⁺) at 50 µM of internal Ca²⁺. In response to night AP commands, β4 coexpression led to a significant twofold increase in BK₀ peak current compared with BK₀ alone (Fig. 6 B). This effect was consistent with the left-shifted G-V from BK₀/β4 currents at a similar high Ca²⁺ (Fig. 5 E). In contrast, BKSRKR peak currents elicited by night AP commands were not significantly enhanced by coexpression of β4, agreeing with the modestly right-shifted G-V for BKSRKR/β4. In addition, β4 had no significant effect on either BK₀ or BKSRKR currents elicited by day AP commands (Fig. 6, A and B). These results show that β4 differentially affects the BK₀ and BKSRKR splice variants under distinct stimulus conditions, an effect that is consistent with results obtained in symmetrical K⁺ conditions (Fig. 5).

Because BKSRKR peak currents were reduced compared with BK₀ in response to both day and night AP commands without β4, we compared the magnitude of the decrease in BKSRKR peak current in the presence of β4 (Fig. 6). In response to the night AP command without β4, BKSRKR currents were decreased by 2.5-fold compared with BK₀. In the presence of β4, this difference widened to threefold (Fig. 6 B). In contrast, the day AP command produced a significant 2.8-fold reduction in BKSRKR current compared with BK₀ in the absence of β4, which was blunted to a 2.1-fold reduction by the addition of β4 (Fig. 6 A). Furthermore, in the presence of β4, BKSRKR peak currents were no longer statistically different to BK₀ because of an increase in the variability with β4 coexpression. Collectively, these results suggest that β4 plays a greater role in modulating the differences between BK₀ and BKSRKR currents in response to night AP commands than in response to day AP commands.

**Discussion**

In this study, we identified two BK channel splice variants containing novel combinations of alternate exons

![Figure 6](image-url)
(BK₀ and BK₅₉₆₆), which differed from each other by a four-amino acid insert (Fig. 1). We determined that the functional differences between BK₀ and BK₅₉₆₆ currents were caused by the SRKR exon (Fig. 2). Inclusion of the SRKR exon (BK₅₉₆₆) resulted in a large reduction of BK channel activity, demonstrated by a rightward shift in G-V relationships, slower activation rates, and faster deactivation rates. These effects were abrogated by dephosphorylation only when SRKR was present and, conversely, could be reproduced in the absence of SRKR by a phosphomimetic mutation (S642D) (Fig. 3). Overall, the changes in current properties caused by SRKR were larger than those caused by the inclusion of other alternate exons at sites 2, 3, and 4 (Figs. S1 and S2, and Table S1). When BK₅₉₆₆ was examined in a reconstituted physiological system activated by neuronal AP commands in saturating Ca²⁺, we found that even with the slower rate of depolarization and the much shorter duration of AP commands compared with voltage jumps, BK₅₉₆₆ still had reduced activity compared with BK₀. This inhibition was manifested as a reduction in peak current, charge transfer, and current half-width during the AP stimulus compared with BK₀ and was eliminated by mutation of S642 (Fig. 4). Collectively, the central finding of this study establishes the context-dependent phosphorylation of S642 in the presence of the SRKR exon as a novel means for reducing BK activation during an AP stimulus.

The functional significance of SRKR inclusion in BK transcripts parallels the known physiological role for BK currents in regulating neural activity in the SCN. BK currents primarily regulate firing frequency at night in SCN neurons, correlating with the window of higher total BK channel expression and larger BK currents in wild-type SCNs (Panda et al., 2002; Meredith et al., 2006; Montgomery et al., 2013). Yet BK currents have little impact on firing rate during the day, despite the presence of a baseline BK current during this time in wild-type SCN neurons (Meredith et al., 2006; Pitts et al., 2006). Our data suggest that part of the reduced influence for BK currents on SCN neuronal activity during the day could be mediated by an increase in SRKR-containing transcripts, leading to decreased AP-induced BK currents.

To further understand how the changes in BK₅₉₆₆ current properties lead to a reduction in AP-induced current, we considered the effect of SRKR on both kinetic and steady-state parameters, finding alterations in both compared with BK₀. Mutation of S642 to alanine (BK₅₉₆₆-S642A) converted BK₅₉₆₆ steady-state and kinetic properties to be more “BK₀-like” (Fig. 3, C–F). However, the reduction in BK₅₉₆₆ current compared with BK₀ was eliminated when a step command to the same peak voltage applied during the AP command was used (Fig. 4 G), suggesting that the reduced BK₅₉₆₆ current elicited by APs may be largely based on kinetic differences between the variants. Consistent with this, BK₅₉₆₆ currents exhibited slowed activation and faster deactivation at high Ca²⁺ (Fig. 2), factors that could be expected to prominently reduce BK₅₉₆₆ current activation in response to a brief AP stimulus compared with BK₀. Interestingly, when β₄ was coexpressed, which further slows both activation and deactivation of BK₅₉₆₆, only the night AP recapitulated the significant difference in peak current between the variants (Fig. 6). In this case, the slower rise time of the night AP compared with the day AP (Fig. 4 A) may facilitate expression of the intrinsic difference between BK₀ and BK₅₉₆₆. Collectively, these data strongly support the idea that the expression of BK₀ and BK₅₉₆₆ is expected to impact excitability in vivo.

Variants with and without the SRKR insert have been previously cloned from human, chick, and turtle tissues (Tseng-Crank et al., 1994; Rosenblatt et al., 1997; Jones et al., 1999). The suppressive effect of SRKR on BK currents in this study is consistent with some data from more limited investigations of the SRKR insert. In the human BK variants, SRKR right-shifted the V₅/₂ of ~10 mV at 2.4, 75, and 405 μM Ca²⁺ (Tseng-Crank et al., 1994). In another study, BK variants with and without SRKR isolated from the chick cochlea showed no significant differences in V₅/₂ values, but the SRKR-containing clone had reduced voltage sensitivity, indicated by a shallower G-V curve (Rosenblatt et al., 1997). In turtle cochlear variants, the presence of SRKR could either speed or slow activation and left- or right-shift G-V curves, depending on the exons at other splice sites (Jones et al., 1999). Comparing these data to our results, we observed larger V₅/₂ right-shifts of 20–40 mV and no effect on the G-V slope. These differences may be attributable to the novel combination of exons contained within BK₀ and BK₅₉₆₆ splice variants that may interact to modulate the functional effects of SRKR. For example, the human variant also contained the STREX insert, the chick variant contained an additional eight-residue insert in the RCK1 domain, and both lacked the 27-residue Ca²⁺-bowl insert at splice site 3 (Tseng-Crank et al., 1994; Rosenblatt et al., 1997). In our study, the addition of the Ca²⁺-bowl exon also right-shifted the V₅/₂ (Fig. S2). This exon is present in both BK₀ and BK₅₉₆₆ variants and may enhance the right-shifting effect of SRKR in BK₅₉₆₆. Such functional interactions between alternative exons have been described previously in BK channels cloned from Caenorhabditis elegans (Johnson et al., 2011). For example, insertions at a splice site within the RCK1 domain can modulate Ca²⁺ sensitivity and activation kinetics, but only when insertions are present at further splice sites within the RCK1–RCK2 linker (Johnson et al., 2011).

Mechanistically, how does inclusion of the SRKR exon regulate phosphorylation? Consistent with our results showing that the dephosphorylation-dependent functional differences at S642 cannot be detected without SRKR present, insertion of SRKR likely alters the...
structure of the local protein environment to render S642 susceptible to phosphorylation. Examination of the available crystal structures of BK channels supports this idea (Fig. S4). The structure of the cytosolic domain of a human BK channel variant (Wu et al., 2010) contains the homologous residue to S642 but lacks the SRKR insert. In this structure, S642 is positioned at the extreme N terminus of a β strand immediately after a short five-residue loop toward the C-terminal end of the RCK1 domain. The only other isofrom for which there is a crystal structure is a zebrafish isofrom that contains the SRKR insert (Yuan et al., 2012). In this structure, the atomic coordinates corresponding to S642 and several surrounding residues, including the SR portion of SRKR, were not obtained. However, the structure suggests that S642 resides within an inter β-strand loop; therefore, the presence of SRKR shifts S642 from within a β-strand region to a loop (Fig. S4). The shift of S642 into this loop could better position it for accessibility to kinases for phosphorylation. In addition, SRKR insertion would distance S642 from the subsequent hydrophobic residues ILI, the proximity of which reduces the predicted phosphorylation score in NetPhos 2.0. Interestingly, of the 30 phospho-residues detected in rat brain lysates (Yan et al., 2008), no peptide fragment containing S642 was detected at all (phosphorylated or not), presumably because of its small size, leaving open the investigation for BKSRKR variants phosphorylated at S642 in native tissues.

How might SRKR and a phosphate group at S642 act to reduce channel activity? In other studies, phosphorylation has been shown to either increase or decrease BK channel activity, depending on the kinase, residue position, and channel variant (Reinhart et al., 1991; Ling et al., 2000; Tian et al., 2001; Yan et al., 2008). In the case of S642 and SRKR, these residues are not located in the vicinity of any of the divergent ion-binding sites, nor are they close to the transmembrane regions that form the voltage sensors or the pore. In the absence of Ca\(^{2+}\), the rightward V\(_{1/2}\) shift of BKSRKR compared with BK\(_{0}\) indicates that phosphorylation of S642 reduces BK channel activation via steps that are Ca\(^{2+}\) independent, for example, those involved in intrinsic pore opening or the coupling of voltage-sensor movement to pore opening. However, direct effects on voltage-sensor movement are unlikely caused by the parallel slopes of the G-V curves, which indicate similar voltage sensitivities of activation.

In the presence of Ca\(^{2+}\), SRKR also decreases channel activity, at 1 and 100 µM, but not at 10 µM Ca\(^{2+}\). Thus, as well as the Ca\(^{2+}\)-independent effect described above, SRKR also has Ca\(^{2+}\)-dependent effects. Changes in several of the many parameters that define Ca\(^{2+}\)- and voltage-dependent gating could underlie these effects. Although the precise mechanism is not addressed in the experiments performed in this study, one potential explanation may be that BK\(_{SRKR}\) gating is more sensitive to Ca\(^{2+}\) than BK\(_{0}\) gating. Supporting this idea, the decrease in V\(_{1/2}\) with increasing Ca\(^{2+}\) is greater for BK\(_{SRKR}\) compared with BK\(_{0}\). From zero Ca\(^{2+}\), the V\(_{1/2}\) shifts are 45, 188, and 241 mV for BK\(_{SRKR}\), compared with 29, 142, and 211 mV for BK\(_{0}\) at 1, 10, and 100 µM Ca\(^{2+}\) (Fig. 2C and Table S1). Therefore, SRKR would be expected to inhibit gating via a Ca\(^{2+}\)-independent mechanism but conversely increase the ability of Ca\(^{2+}\) to gate the channel. At 10 µM Ca\(^{2+}\), the lack of difference in the V\(_{1/2}\) of BK\(_{SRKR}\) compared with BK\(_{0}\) could be explained if the enhanced Ca\(^{2+}\)-dependent gating overrode the decreased Ca\(^{2+}\)-independent gating of BK\(_{SRKR}\). This possibility is suggested by the observation that the range where no differences were observed was in the V\(_{1/2}\) between BK\(_{0}\) and BK\(_{SRKR}\) (>1 and <100 µM) is near the Ca\(^{2+}\) affinities estimated for both the RCK1 and the Ca\(^{2+}\)-bowl sites (~4 µM; Bao et al., 2002). Thus, under conditions where the differences in the ability of Ca\(^{2+}\) to gate the two variants are expected to be small (in the absence, at low, and at saturating Ca\(^{2+}\)), the suppressive effect of SRKR would prevail, while the balance between the two gating processes would be altered at intermediate Ca\(^{2+}\) (10 µM).

In addition to the complex effects resulting from the addition of the SRKR exon, we found that coexpression of thepredominant brain BK channel accessory subunit, β4, had a range of effects on BK\(_{0}\) and BK\(_{SRKR}\). As has been reported previously for other BK channel isoforms, β4 slowed both activation and deactivation kinetics (Behrens et al., 2000; Brenner et al., 2000a). In addition, β4 altered BK\(_{0}\) and BK\(_{SRKR}\) G-V relationships in a Ca\(^{2+}\)-dependent manner. Ca\(^{2+}\)-dependent changes in β4 effects have been observed previously for other BK channel variants (Brenner et al., 2000a, Lippiat et al., 2003; Ha et al., 2004; Wang et al., 2006, 2009; Petrik and Brenner, 2007). At low Ca\(^{2+}\), β4 inhibits BK currents, whereas at high Ca\(^{2+}\), it increases BK currents. A qualitatively similar process is occurring with BK\(_{0}\) and BK\(_{SRKR}\), with a large rightward shift in G-V curves at 10 µM Ca\(^{2+}\) and small rightward (BK\(_{SRKR}\)) or leftward (BK\(_{0}\)) shifts at 100 µM Ca\(^{2+}\). Previous work has shown that β4 acts to decrease channel activation by increasing the energetic barrier for intrinsic pore opening and decreasing coupling between the voltage sensors and Ca\(^{2+}\) binding, and conversely, β4 acts to increase channel activation by increasing the coupling between Ca\(^{2+}\) binding and pore opening, and stabilizing voltage-sensor activation (Wang et al., 2006; Contreras et al., 2012). Given the broadly similar relationships between V\(_{1/2}\) changes and Ca\(^{2+}\) from previous work and our data, it is reasonable to think that β4 is impacting BK\(_{0}\) and BK\(_{SRKR}\) function via the same mechanisms, with the differences between the variants attributable to shifts in the Ca\(^{2+}\) concentration at which β4 switches its net effect on the G-V relationship. This Ca\(^{2+}\) concentration would be expected to be higher for BK\(_{SRKR}\) than BK\(_{0}\). However, the
relative interaction of B4's effects on Ca2+ dependence, activation, and deactivation kinetics remains to be determined for the observed differences between BKα and BKSRKR AP-induced currents.

We wish to thank Hyunjin Choi for assistance with cloning BK splice variants, Jessica Lu for assistance with GenBank sequence submissions, and Man-Kyo Chung for comments on the manuscript.

This work was supported by grants from National Heart, Lung, and Blood Institute (R01-HL102758), National Institute of Diabetes and Digestive Kidney Diseases (R21-DK089337), National Science Foundation (IOS-0956257), and The American Physiological Society’s Ryuji Ueno award, sponsored by the S&R Foundation (all to A.L. Meredith). J.P. Whitt was supported by National Institute of Arthritis and Musculoskeletal and Skin Diseases training grant T32-AR007592. All authors declare no conflict of interest.

Sharona E. Gordon served as editor.

Submitted: 31 July 2013
Accepted: 11 October 2013

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