Structural determinants of Kv β 1.3-induced channel inactivation: a hairpin modulated by PIP₂

Niels Decher^{1,2,5,*}, Teresa Gonzalez^{1,4,5},
Anne Kathrin Streit², Frank B Sachse¹,
Vijay Renigunta², Malle Soom³,
Stefan H Heinemann³, Jürgen Daut²
and Michael C Sanguinetti¹

¹Nora Eccles Harrison Cardiovascular Research and Training Institute, University of Utah, Salt Lake City, UT, USA, ²Institute for Physiology and Pathophysiology, Department of Cell Physiology, University of Marburg, Marburg, Germany, ³Department of Biophysics, Center for Molecular Biomedicine, Friedrich Schiller University Jena, Jena, Germany and ⁴Instituto de Investigaciones Biomédicas 'Alberto Sols' CSIC/UAM, Departamento de Modelos Experimentales de Enfermedades Humanas, Madrid, Spain

Inactivation of voltage-gated Kv1 channels can be altered by Kv β subunits, which block the ion-conducting pore to induce a rapid ('N-type') inactivation. Here, we investigate the mechanisms and structural basis of Kv β 1.3 interaction with the pore domain of Kv1.5 channels. Inactivation induced by Kv β 1.3 was antagonized by intracellular PIP₂. Mutations of R5 or T6 in Kv β 1.3 enhanced Kv1.5 inactivation and markedly reduced the effects of PIP₂. R5C or T6C Kv β 1.3 also exhibited diminished binding of PIP₂ compared with wild-type channels in an *in vitro* lipid-binding assay. Further, scanning mutagenesis of the N terminus of Kv β 1.3 revealed that mutations of L2 and A3 eliminated N-type inactivation. Double-mutant cycle analysis indicates that R5 interacts with A501 and T480 of Kv1.5, residues located deep within the pore of the channel. These interactions indicate that Kv β 1.3, in contrast to Kv β 1.1, assumes a hairpin structure to inactivate Kv1 channels. Taken together, our findings indicate that inactivation of Kv1.5 is mediated by an equilibrium binding of the N terminus of Kv β 1.3 between phosphoinositides (PIPs) and the inner pore region of the channel.

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Introduction

Voltage-gated potassium (Kv) channels are essential for regulating resting membrane potential, repolarization of action potentials, pacemaking and neurotransmitter release. Kv channels are tetrameric complexes formed by coassembly

of four identical or homologous α -subunits. Rapid N-type inactivation of Kv1 channels can result from binding of a single N-terminal hydrophobic, 'inactivation ball' peptide of an α -subunit to the inner pore region of the channel complex (Hoshi *et al*, 1990). The inactivation ball of *Shaker* B (Kv1.1 of *Drosophila*) α -subunits is a random coil in aqueous solution (Lee *et al*, 1993), but forms a β -hairpin structure when exposed to a more hydrophobic environment (Lee *et al*, 1993; Fernandez-Ballester *et al*, 1995). There may be variation in how inactivation ball peptides interact with the inner pore as the NMR structures of the inactivation balls of Kv1.4 and Kv3.4 α -subunits are clearly different (Antz *et al*, 1997).

An alternative structural basis of N-type inactivation of Kv1 channels has been described. Fast inactivation can also be mediated by the N terminus of a Kv β subunit (Rettig *et al*, 1994; Heinemann *et al*, 1996) that is tethered to the T1 domain of a Kv1 α -subunit. For example, Kv β 1, Kv β 2 and Kv β 3 subunits alter the activation and inactivation gating of Kv1.5 channels (Leicher *et al*, 1998). The inactivation of Kv1 channels is diversified by alternative splicing of the Kv β 1 gene, resulting in the isoforms Kv β 1.1, Kv β 1.2 and Kv β 1.3. The N terminus of Kv β 1 subunits was proposed to enter the pore of a Kv1 channel as an extended peptide (Zhou *et al*, 2001). In contrast, the N-terminal ball peptides of Kv α -subunits were proposed to form a compact hairpin structure that binds to the inner vestibule to occlude the pore (Antz *et al*, 1997; Antz and Fakler, 1998). As illustrated by comparison of the N-terminal regions of two Kv α and three Kv β subunits in Figure 1A, there is no apparent sequence conservation for inactivation ball peptides.

Mutations in the N terminus of Kv β or Kv1 subunits can prevent their ability to inactivate Kv channels. For example, deletion of 10 amino acids from the N terminus of Kv β 1.3 (Uebele *et al*, 1998) causes a loss of function as does the L7E mutation in *Shaker* B α -subunits (Hoshi *et al*, 1990). Cysteine residues at position 7 of Kv β 1.1 (Rettig *et al*, 1994), position 6 of Kv3.4 (Stephens and Robertson, 1995) or position 13 of Kv1.4 (Ruppersberg *et al*, 1991) confer a redox sensitivity to channel inactivation. The loss of function by L7E or L7R in *Shaker* B (Hoshi *et al*, 1990) can be mimicked by phosphorylation of Y8 that prevents formation of a functional hairpin structure (Encinar *et al*, 2002). In addition, N-type inactivation of Kv1.5/Kv β 1.3 channels is modulated by protein kinase C (Kwak *et al*, 1999) and inactivation of Kv1.1/Kv β 1.1 is antagonized by intracellular Ca²⁺ (Jow *et al*, 2004). However, the molecular mechanisms and structural basis of Kv α –Kv β interactions that mediate these effects are poorly understood.

N-type inactivation of Kv3.4 alone or inactivation of Kv1.1 mediated by Kv β 1.1 are antagonized by PIP₂ (Oliver *et al*, 2004). For Kv3.4, binding of PIP₂ to residues R13 and K14 of the N terminus appears to mediate this effect (Oliver *et al*, 2004). Although all three Kv β 1 isoforms introduce N-type inactivation, they differ in inactivation kinetics, intracellular

*Corresponding author. Institute of Physiology and Pathophysiology, Philipps-University Marburg, Deutschhausstraße 1–2, Marburg, Hessen 35037, Germany. Tel.: +49 642 128 621 48; Fax: +49 642 128 689 60; E-mail: decher@staff.uni-marburg.de

⁵These authors contributed equally to this work

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Figure 1 N-type inactivation of Kv1.5 by Kv β 1.3. (A) Alignment of the N termini of Kv β isoforms and of N-type inactivating Kv3.4 and Shaker channels. (B) Kv1.5 currents during short and long voltage steps to +70 mV, illustrating slow time course of C-type inactivation. (C) Superimposed current traces in response to depolarizations applied in 10-mV increments to test potentials ranging from –70 to +70 mV for Kv1.5 alone, co-expressed with Kv β 1.3 or with a Kv β 1.3, which lacks the N-terminal amino acids 2–10.

modulation and expression pattern. This diversity plus cellular regulation helps to tune K⁺ channels to serve specific function.

We recently identified residues in the pore region of Kv1.5 that interact with Kv β 1.3 (Decher *et al*, 2005). Blockade of Kv1.5 by drugs such as S0100176 and bupivacaine can be modified by Kv β 1.3. Accordingly, site-directed mutagenesis studies revealed that the binding sites for drugs and Kv β 1.3 partially overlap (Gonzalez *et al*, 2002; Decher *et al*, 2004, 2005). In the present study, we used a mutagenesis approach to identify the residues of Kv β 1.3 and Kv1.5 that interact with one another to mediate fast inactivation. We also examined the structural basis for inhibition of Kv β 1.3-mediated inactivation by PIP₂. Taken together, our findings indicate that when dissociated from PIP₂, the N terminus of Kv β 1.3 forms a hairpin structure and reaches deep into the central cavity of the Kv1.5 channel to cause inactivation. This binding mode of Kv β 1.3 differs from that found earlier for Kv β 1.1, indicating a Kv β 1 isoform-specific interaction in the pore cavity.

Results

Identification of residues important for Kv β 1.3 function using cysteine- and alanine-scanning mutagenesis

Wild-type (WT) Kv1.5 channels activate rapidly and exhibit almost no inactivation when cells are depolarized for 200 ms (Figure 1B, left panel). Longer pulses cause channels to inactivate by a slow ‘C-type’ mechanism that results in an ~20% decay of current amplitude during 1.5 s depolarizations to +70 mV (Figure 1B, right panel). At this potential, inactivation is monoexponential with a time constant of ~0.8 s. Co-expression of Kv1.5 with Kv β 1.3 subunits introduces an additional, rapid component (‘N-type’) of inactivation (Figure 1C). As reported previously (Uebele *et al*, 1998), Kv β subunit-induced fast inactivation is eliminated when

Kv β 1.3 is truncated by the removal of residues 2–10 (Kv β 1.3 Δ 2–10; Figure 1C).

To assess the importance of specific residues in the N terminus of Kv β 1.3 for N-type inactivation, we made individual mutations of residues 2–11 of Kv β 1.3 to alanine or cysteine and co-expressed these mutant subunits with Kv1.5 subunits. Alanine residues were substituted with cysteine or valine. Substitution of native residues with alanine or valine introduces or retains hydrophobicity without disturbing helical structure, whereas substitution with cysteine introduces or retains hydrophilicity. In addition, cysteine residues can be subjected to oxidizing conditions to favour crosslinking with another cysteine residue. Representative currents recorded in oocytes co-expressing WT Kv1.5 plus mutant Kv β 1.3 subunits are depicted in Figure 2A and B. Mutations at positions 2 and 3 of Kv β 1.3 (L2A/C and A3V/C) led to a complete loss of N-type inactivation (Figure 2A–D). A similar, but less pronounced, reduction of N-type inactivation was observed for A4C, G7C and A8V mutants. In contrast, mutations of R5, T6 and G10 of Kv β 1.3 increased inactivation of Kv1.5 channels (Figure 2A and B). The effects of all the Kv β 1.3 mutations on inactivation are summarized in Figure 2C and D. In addition, the inactivation of channels with cysteine substitutions was quantified by their fast and slow time constants (τ_{inact}) during a 1.5-s pulse to +70 mV in Figure 2E. In the presence of Kv β 1.3, the inactivation of Kv1.5 channels was bi-exponential. With the exceptions of L2C and A3C, cysteine mutant Kv β 1.3 subunits introduced fast inactivation (Figure 2E, lower panel). Acceleration of slow inactivation was especially pronounced for R5C and T6C Kv β 1.3 (Figure 2E, lower panel). The more pronounced steady-state inactivation of R5C and T6C (Figure 2A and B) was not caused by a marked increase of the fast component of inactivation (Figure 2E, upper panel).

Kv β 1.3 mutations change inactivation kinetics independent of intracellular Ca²⁺

Rapid inactivation of Kv1.1 by Kv β 1.1 is antagonized by intracellular Ca²⁺. This Ca²⁺-sensitivity is mediated by the N terminus of Kv β 1.1 (Jow *et al*, 2004), but the molecular determinants of Ca²⁺-binding are unknown. The mutation-induced changes in the rate of inactivation could potentially result from an altered Ca²⁺-sensitivity of the Kv β 1.3 N terminus. Application of the Ca²⁺ ionophore ionomycin (10 μ M) for 3 min removed rapid inactivation of Kv1.1/Kv β 1.1 channels (Figure 3A). However, this effect was not observed when either Kv1.5 (Figure 3B) or Kv1.1 (Figure 3C) was co-expressed with Kv β 1.3 subunits. Thus, alternative splicing of Kv β 1 can alter its Ca²⁺-sensitivity.

Mutant Kv β 1.3 subunits that disrupt inactivation retain ability to alter voltage-dependent gating of Kv1.5 channels

We reported earlier that although mutation of specific residues in the S6 domain of Kv1.5 could disrupt N-type inactivation, these mutations did not alter the ability of Kv β 1.3 to cause shifts in the voltage dependence of channel gating (Decher *et al*, 2005). This finding suggests that WT Kv β 1.3 can bind to and affect Kv1.5 gating without blocking the pore. Can mutant Kv β 1.3 subunits that no longer induce fast N-type inactivation still cause shifts in the gating of Kv1.5? This question was addressed by comparing the voltage

Figure 2 Scanning mutagenesis of the Kvβ1.3 N terminus. Superimposed currents elicited by depolarizations applied in 10-mV increments to test potentials ranging from -70 to $+70$ mV for Kv1.5 co-expressed with Kvβ1.3 containing either (A) alanine or (B) cysteine mutations as indicated. (C, D) Relative inactivation plotted as a ratio of steady-state current after 1.5 s (I_{ss}) to peak current (I_{max}) for alanine/valine or cysteine point mutations of the Kvβ1.3 N terminus. A value of 1.0 indicates no inactivation; a value of 0 indicates complete inactivation. (E) Kinetics of inactivation for Kv1.5 and Kv1.5/Kvβ1.3 channel currents determined at $+70$ mV. Labels indicate cysteine mutations in Kvβ1.3. Upper panel: relative contribution of fast (A_f) and slow (A_s) components of inactivation. Lower panel: time constants of inactivation. For (C–E), $*P < 0.05$; $**P < 0.005$ compared with Kv1.5 plus wild-type Kvβ1.3 ($n = 4–13$).

Figure 3 Ca^{2+} -sensitivity of Kvβ1.1 versus Kvβ1.3. Currents were recorded at $+70$ mV under control conditions and after the addition of $10 \mu M$ ionomycin. (A) Ionomycin prevents N-type inactivation of Kv1.1 by Kvβ1.1. Elevation of intracellular $[Ca^{2+}]$ does not prevent Kvβ1.3-induced N-type inactivation of Kv1.5 (B) or Kv1.1 (C).

dependence of activation and inactivation of Kv1.5 when co-expressed with WT and mutant Kvβ1.3 subunits. WT subunits shifted the voltage required for half-maximal activation by -15 mV and the voltage dependence of inactivation by -11 mV (Figure 4A and B). Mutant Kvβ1.3 subunits retained their ability to cause negative shifts in the half-points of activation and inactivation, albeit to a variable degree (Figure 4A and B). These findings suggest that point mutations in the N terminus of Kvβ1.3, including those that eliminated N-type inactivation, did not disrupt co-assembly of Kvβ1.3 with the Kv1.5 channel.

Interaction of PIP₂ with R5 of Kvβ1.3

The most pronounced gain of Kvβ1.3-induced inactivation was observed after mutation of R5 or T6 to cysteine or alanine. To further explore the role of charge at position 5 in Kvβ1.3, R5 was substituted with another basic (K), a neutral (Q) or an acidic (E) amino acid. Introduction of neutral or negatively charged residues at position 5 accelerated Kvβ1.3-induced fast inactivation, without increasing the relative contribution of the fast component (Figure 5A and B). In contrast, the conservative exchange R5K did not alter inactivation (Figure 5B, lower panel). All of the R5 mutant Kvβ1.3 subunits retained their ability to cause negative shifts in the half-points of activation and inactivation (Figure 5C).

PIP₂ antagonizes N-type inactivation of Kv3.4 by binding to its N-terminal residues R13 and K14 and immobilizing the inactivation ball (Oliver *et al*, 2004). PIP₂ also removes fast inactivation of Kv1.1 channels by immobilization of Kvβ1.1 (Oliver *et al*, 2004). We tested whether the R5C or T6C mutations in Kvβ1.3 altered the binding affinity of PIP₂ for the Kvβ1.3 N terminus (residues 1–33). A lipid-binding assay was performed using N-terminal Kvβ1.3 GST fusion proteins and fluorescently labelled liposomes with incorporated PI(4,5)P₂, as described earlier (Soom *et al*, 2001). The PIP₂-binding affinity (expressed as relative arbitrary units) of R5C or T6C N-terminal Kvβ1.3 fusion proteins was reduced compared with WT fusion proteins (Figure 5D). Specifically, the fluorescence signal was reduced by 90% for R5C and 41% for T6C Kvβ1.3. The reduction in PIP₂-binding affinity for the

Figure 4 Shifts in voltage dependence of Kv1.5 gating induced by co-expression with wild-type and mutant Kv β 1.3 subunits. (A) Half-point ($V_{1/2}$) for activation (left panels) and inactivation (right panels) of current determined for oocytes expressing Kv1.5 channels alone or when co-expressed with Kv β 1.3 subunits with alanine/valine substitutions of indicated residues. (B) Same analysis for cysteine substitutions. * $P < 0.05$; ** $P < 0.005$ compared with Kv1.5 alone ($n = 4-11$).

T6C mutant might be caused by impairment of electrostatic PIP₂ interaction with the neighbouring residue R5.

PIP₂ (10 μ M) eliminated Kv β 1.3-induced N-type fast inactivation of Kv1.5 measured in inside-out macro-patches from *Xenopus* oocytes (Figure 5E, left panel and 5F). In contrast, PIP₂ only partially attenuated the fast channel inactivation mediated by R5C Kv β 1.3 (Figure 5E, right panel and 5F). PIP₂ antagonized Kv1.5 inactivation by Kv β 1.3 in a concentration-dependent manner (Figure 5G). Attenuation of Kv β 1.3-induced inactivation was also observed for 10 μ M PIP and PIP₃ (Figure 5H). In contrast, inactivation was only partially antagonized by these phosphoinositides (PIPs) when Kv1.5 was co-expressed with R5C Kv β 1.3 (Figure 5I). Thus, the sensitivity of Kv1.5 channels to Kv β 1.3 is modulated by PIPs, and R5 is a key residue for this functional modulation.

Endogenous PIP₂ modulates Kv1.5/Kv β 1.3 channels

Next, we tested whether a depletion of endogenous PIP₂ can modulate inactivation of the Kv1.5/Kv β 1.3 channel complex. Kv1.5, Kv β 1.3 and the Gq-coupled α_{1A} -adrenoreceptor were co-expressed in *Xenopus* oocytes. We have shown earlier that activation of α_{1A} -adrenoreceptors by methoxamine triggers Gq-coupled inhibition of the K⁺ channel TASK-1 in oocytes (Putzke *et al*, 2007). Stimulation of the α_{1A} -receptor with 1 μ M methoxamine accelerated inactivation and led to a small but significant reduction in steady-state current amplitude of the Kv1.5/Kv β 1.3 channel complex. Currents were reduced by $10.5 \pm 1.9\%$ ($n = 8$). However, receptor stimulation might not be sufficient to globally deplete PIP₂ from the plasma membrane of an *Xenopus* oocyte, especially if the channel complex and receptors are not adequately colocalized in the cell membrane, an argument used to explain why stimulation of several Gq-coupled receptors (bradykinin BK2, muscarinic M1, TrkA) did not cause the expected shift in the voltage dependence of HCN channel activation (Pian *et al*, 2007).

The Kv1.5/Kv β 1.3 channel complex expressed in *Xenopus* oocytes has a more pronounced inactivation when recorded from an inside-out macropatch (Figure 5E, left panel) as compared with two-electrode voltage-clamp recordings (Figure 1C, middle panel). I_{ss}/I_{max} was significantly decreased from 0.40 ± 0.02 (Figure 2C) to 0.24 ± 0.04 (Figure 5G) in an excised patch. This effect might be partially explained by PIP₂ depletion from the patch. Therefore, we performed inside-out macropatches from *Xenopus* oocytes and applied poly-lysine (25 μ g/ml) to the inside of the

patch to deplete PIPs from the membrane (Oliver *et al*, 2004). Poly-lysine enhanced the extent of steady-state inactivation, decreasing the I_{ss}/I_{max} from 26.0 ± 5.0 to $10.5 \pm 4.3\%$ (Figure 5J). Taken together, these findings indicate that endogenous PIPs are important determinants of the inactivation kinetics of the Kv1.5/Kv β 1.3 channel complexes.

Co-expression of mutant Kv1.5 and Kv β 1.3 subunits

In an attempt to determine the structural basis of Kv β 1.3 interaction with the S6 domain of Kv1.5, single cysteine mutations were introduced into each subunit. Our previous alanine scan of the S6 domain (Decher *et al*, 2005) identified V505, I508, V512 and V516 in Kv1.5 as important for interaction with Kv β 1.3. Here, these S6 residues (and A501) were individually substituted with cysteine and co-expressed with Kv β 1.3 subunits containing single cysteine substitutions of L2–T6. Potential physical interaction between cysteine residues in the α - and β -subunits was assayed by changes in the extent of current inactivation at +70 mV (Figure 6).

N-type inactivation was eliminated when L2C Kv β 1.3 was co-expressed with WT Kv1.5 or mutant Kv1.5 channels with cysteine residues in pore-facing positions (Figures 2B and 6A). Co-expression of L2C Kv β 1.3 with I508C Kv1.5 slowed C-type inactivation, whereas C-type inactivation was enhanced when L2C Kv β 1.3 was co-expressed with V512C Kv1.5 (Figure 6A). For A3C Kv β 1.3, the strongest changes in inactivation were observed by mutating residues V505, I508 and V512 in Kv1.5 (Figure 6B). For A4C Kv β 1.3, the extent of inactivation was changed by co-expression with Kv1.5 subunits carrying mutations at position A501, V505 or I508 (Figure 6C). The pronounced inactivation observed after co-expression of R5C Kv β 1.3 with WT Kv1.5 was significantly reduced by the mutation A501C (Figure 6D). A501 is located in the S6 segment close to the inner pore helix. The strong inactivation of Kv1.5 channels by T6C Kv β 1.3 was antagonized by cysteine substitution of A501, V505 and I508 of Kv1.5 (Figure 6E). Taken together, these data suggest that R5 and T6 of Kv β 1.3 interact with residues located in the upper S6 segment of Kv1.5, whereas L2 and A3 apparently interact with residues in the middle part of the S6 segment.

Mutations located deep in the pore of Kv1.5 reduce R5C Kv β 1.3 gain of function

To investigate further whether R5 of Kv β 1.3 interacts with specific residues in the upper region in S6 of Kv1.5, we co-expressed R5C Kv β 1.3 with Kv1.5 subunits harbouring single

Figure 5 Phosphoinositide-binding of Kv β 1.3 prevents N-type inactivation. (A) Superimposed current traces in response to depolarizations applied in 10-mV increments to test potentials ranging from -70 to $+70$ for Kv1.5 co-expressed with Kv β 1.3 subunits with indicated mutations of R5. (B) Kinetics of inactivation for Kv1.5 and Kv1.5/Kv β 1.3 channel currents determined at $+70$ mV ($n=4-13$). Labels indicate mutations of R5 in Kv β 1.3. Upper panel: relative contribution of fast (A_f) and slow (A_s) components of inactivation. Lower panel: time constants of inactivation. (C) Half-points ($V_{1/2}$) for activation (left panel) and inactivation (right panel) of current determined for oocytes expressing Kv1.5 channels alone or when co-expressed with WT or R5 mutant Kv β 1.3 subunits ($n=4-13$). For (B, C), $*P<0.05$; $**P<0.005$ compared with Kv1.5 alone. $^{##}P<0.005$ compared with Kv1.5 + Kv β 1.3 WT. (D) *In vitro* PIP₂-binding assay of N-terminal Kv β 1.3 fragments. Average-binding data of GST alone and GST-Kv β 1.3 fusion proteins (amino acids 1-33) of WT and mutant subunits. Relative arbitrary units of PIP₂-binding are shown for liposomes containing 5 mol% PI(4,5)P₂. $**P<0.01$. (E) Fast inactivation of Kv1.5/Kv β 1.3 channels is removed by application of 10μ M PIP₂ to the cytosolic side of an inside-out macropatch from *Xenopus* oocytes (left panel). Here, 10μ M PIP₂ only partially antagonizes inactivation introduced by R5C Kv β 1.3 (right panel). (F) R5C mutation in Kv β 1.3 blunts the slowing of fast inactivation induced by a 15 s application of PIP₂ to inside-out macro-patches ($n=6-8$). (G-J) Effect of PIPs on relative inactivation of current in inside-out macropatches plotted as a ratio of steady-state current after 1.5 s (I_{ss}) to peak current (I_{max}). For all experiments $n=4-7$. (G) [PIP₂]-dependent reduction of inactivation of Kv1.5/Kv β 1.3 channel current. (H) At 10μ M, PIP, PIP₂ and PIP₃ have equivalent effects on WT Kv1.5/Kv β 1.3 channel inactivation. (I) At 10μ M, PIP, PIP₂ and PIP₃ have similar but reduced effects on Kv1.5/R5C Kv β 1.3 channel inactivation. (J) Poly-lysine (polyK, 25μ g/ml) applied to cytosolic side of membrane enhances inactivation of Kv1.5/Kv β 1.3 channels ($n=6$). For (G-J) $*P<0.05$ compared with Kv1.5 alone by Student's *t*-test.

alanine mutations. In addition, R5C Kv β 1.3 was paired with T480A or A501V Kv1.5 subunits (Figure 7A). These co-expression studies confirmed the putative interaction between R5 and A501, identified with the A501C mutation

(Figure 6D) and revealed potential interactions of R5 with V505 and T480 (Figure 7A). We also studied several double alanine mutations in S6 (V505A/I508A, V505A/V512A and I508A/V512A). The double mutation V505A/I508A

Figure 6 Co-expression of mutant Kv1.5 and Kv β 1.3 subunits. Upper panels are plots of current inactivation (expressed as % reduction in peak current) induced by a 1.5 s pulse to +70 mV. Lower panels show superimposed currents elicited by shorter (200 ms) pulses to potentials ranging from -70 to +70 mV for channels that altered inactivation the most. Possible interacting residues are indicated by arrows (for example, L2C of Kv β 1.3 with I508C Kv1.5 in (A)). For all experiments $n = 4-13$.

Figure 7 Arginine 5 of Kv β 1.3 interacts with residues near the selectivity filter of Kv1.5. (A) Traces for mutant channel currents elicited by pulses to potentials ranging from -70 to +70 mV, applied in 10-mV increments. Labels indicate point mutations of pore-facing residues in Kv1.5. All were co-expressed with R5C Kv β 1.3. Mutations A501V and T480A strongly reduce inactivation by R5C Kv β 1.3. (B) Current traces for channels with indicated double mutations of Kv1.5 and co-expressed with R5C Kv β 1.3. Scale bars represent 0.1 μ A and 100 ms, respectively. (C) Inactivation (expressed as % reduction in peak current for 200 ms pulse to +70 mV) of WT and indicated mutant Kv1.5/R5C Kv β 1.3 channels. The ability of R5C Kv β 1.3 to induce inactivation is blunted when residues located high in the inner cavity of Kv1.5 are mutated. For all experiments, $n = 4-13$.

eliminated N-type inactivation induced by R5C (Figure 7B), whereas the other double mutations had no discernable effect on fast inactivation (Figure 7B and C). Thus, the enhanced N-type inactivation induced by R5C Kv β 1.3 was only attenuated by single-residue mutations when they were located in the upper S6 segment or the pore helix (e.g. V505, A501 and T480; Figure 7A and C). The reduction in inactivation was the more pronounced the deeper the mutations were located in the pore cavity (Figure 7C).

Co-expression of WT Kv1.5 with Kv β 1.3 cysteine mutants under oxidizing conditions

Exposure of oocytes to the membrane-permeable oxidizing agent, *t*-butyl HO₂ (100 μ M) had no significant effect on WT Kv1.5/Kv β 1.3 channel inactivation (Figure 8A). However, oxidation of channels formed by co-expression of Kv1.5 with R5C or T6C Kv β 1.3 subunits caused ~20-fold increase in current magnitude, when analysed at the end of a 1.5 s pulse to +70 mV (Figure 8B and C). Oocytes co-expressing Kv1.5 and L2C or A3C Kv β 1.3 were also relatively unaffected by oxidation, whereas A4C Kv β 1.3 had an intermediate sensitivity (Figure 8C). Redox sensitivity was correlated with the extent of inactivation induced by the specific mutation in Kv β 1.3. Unfortunately, these effects prevented cross-linking experiments as an aid to identify specific residue interactions between Kv β 1.3 and Kv1.5. However, the prevention of R5C-mediated channel inactivation by *t*-butyl HO₂ were utilized to gain further evidence that this residue is located deep within the pore. When 100 μ M *t*-butyl HO₂ was applied to Kv1.5/R5C Kv β 1.3 channels that were inactivated by a prolonged pulse to +40 mV, only partial relief of inactivation was achieved (Figure 8D). Inactivation was nearly completely removed when the cell was subsequently held at -80 mV for 3 min before recording another voltage step to +40 mV (Figure 8D, right panel). Summary data for multiple cells ($n = 8$) are plotted in Figure 8E. The oxidation-induced alteration of R5C channel gating does not require

Figure 8 Effects of oxidation on WT and cysteine mutant Kv1.5/Kv β 1.3 channels. (A) *t*-Butyl HO₂ (100 μ M) has no effect on WT Kv1.5/Kv β 1.3 current elicited by 1 s pulse to +70 mV. Currents were normalized to the peak amplitude before the addition of *t*-butyl HO₂. (B) Normalized currents of Kv1.5/R5C Kv β 1.3 before and after the application of 100 μ M *t*-butyl HO₂. (C) Change in Kv1.5/Kv β 1.3 channel current magnitude induced by 100 μ M *t*-butyl HO₂. Mutations of Kv β 1.3 are indicated on X-axis. (D, E) State-dependent modification of inactivation by oxidation. Kv1.5/R5C Kv β 1.3 current was first recorded with a voltage step to +40 mV (D, left panel). Subsequently, the oocyte was held at +40 mV while the cell chamber was perfused with 100 μ M *t*-butyl HO₂ (D, middle panel). The oocyte was then clamped to -80 mV for 3 min before another voltage step to +40 mV was recorded (D, right panel). (E) Change in current amplitude induced by oxidation when oocytes were held at a potential of +40 mV (*) and for the same cells after holding at -80 mV for 3 min (#). **P* < 0.05. (F) Voltage steps to +40 mV were repeated every 30 s. After stabilization of the current, the oocyte was held at -80 mV without pulsing while the chamber was perfused with 100 μ M *t*-butyl HO₂ for 8 min. The oocyte was then pulsed again to +40 mV. Insets show current traces before the application of 100 μ M *t*-butyl HO₂ and with the first voltage step to +40 mV following the 8 min pulse-free period. Scale bars represent 0.5 μ A and 50 ms, respectively. Current inactivation was recovered after washout of *t*-butyl HO₂. (G) Double-mutant cycle analysis of channels with indicated mutations in Kv1.5 and Kv β 1.3 subunits.

pulsing. When oocytes were held at -80 mV during perfusion with *t*-butyl HO₂, currents were maximally increased during the first pulse to +40 mV, and this effect was fully reversible upon washout of *t*-butyl HO₂ from the bath solution (Figure 8F). The relatively rapid reversal indicates that the C5 in Kv β 1.3 was probably oxidized to a sulphinic or sulphonic acid (Claiborne *et al*, 2001; Poole *et al*, 2004), rather than forming a disulphide bridge with another Cys in the same or another Kv β 1.3 subunit. These findings suggest that when Kv β 1.3 subunit is bound to the channel pore, it is protected from the oxidizing agent.

Double-mutant cycle analysis of Kv1.5-Kv β 1.3 interactions

The experiments summarized in Figures 6D and E, and 7A-C predict that R5 and T6 of Kv β 1.3 interact with residues in the upper S6 segment, near the selectivity filter of Kv1.5. In contrast, for Kv β 1.1 and Kv1.4 (Zhou *et al*, 2001), this interaction would not be possible because residue 5 interacts with a valine residue equivalent to V516 that is located in the lower S6 segment (Zhou *et al*, 2001). To identify residues of Kv1.5 that potentially interact with R5 and T6, we performed a double-mutant cycle analysis. The *K_d* values for single

mutations (α or β subunit) and double mutations (α and β subunits) were calculated to test whether the effects of mutations were coupled. The apparent K_d values were calculated based on the time constant for the onset of inactivation and the steady-state value (% inactivation; see Materials and methods). Figure 8G summarizes the analysis for the co-expressions that resulted in functional channel activity. Surprisingly, no strong deviation from unity for Ω was observed for R5C and T6C in combination with A501C, despite the effects observed on the steady-state current (Figure 6D and E). In addition, only small deviations from unity for Ω were observed for R5C co-expressed with V505A, although the extent of inactivation was altered (Figure 7A). The highest Ω values were for R5C in combination with

T480A or A501V. These data, together with the results shown in Figures 6 and 7, suggest that Kv β 1.3 binds to the pore of the channel with R5 near the selectivity filter. In this conformation, the side chain of R5 might be able to reach A501 of the upper S6 segment, which is located in a side pocket close to the pore helix.

Model of the Kv β 1.3-binding mode in the pore of Kv1.5 channels

Our data suggest that R5 of Kv β 1.3 can reach deep into the inner cavity of Kv1.5. Our observations are difficult to reconcile with a linear Kv β 1.3 structure as proposed for interaction of Kv β 1.1 with Kv1.1 (Zhou *et al*, 2001). The Kv1.5 residues proposed to interact with Kv β 1.3 are

Figure 9 Structural model of Kv β 1.3 bound to the pore of Kv1.5 channels. (A) Amino-acid sequence of the Kv1.5 pore-forming region. Residues that may interact with Kv β 1.3 based on an earlier site-directed mutagenesis study (Decher *et al*, 2005) are depicted in bold. (B) Structure of the N-terminal region (residues 1–11) of Kv β 1.3. (C) Kv β 1.3 docked into the Kv1.5 pore homology model showing a single subunit. Kv β 1.3 side chains are shown as ball and stick models and residues of the Kv β 1.3-binding site in Kv1.5 are depicted with van der Waals surfaces. The symbol ' indicates the end of long side chains. (D) Kv β 1.3 docked into the Kv1.5 pore homology model showing two subunits. (E) Kv β 1.3 hairpin bound to Kv1.5. Two of the four channel subunits in stereo-view are depicted.

Figure 10 Tryptophan substitutions of R5, T6, G7 and G10. Currents shown were elicited by 200 ms pulses to test potentials ranging from -70 to $+70$ mV from a holding potential of -80 mV. Peak current amplitudes were reduced by $78.8 \pm 3.1\%$ ($n=8$) for R5W, by $86.1 \pm 2.8\%$ for T6W ($n=9$), by $12.5 \pm 1.8\%$ for G7W ($n=10$) and by $60.7 \pm 2.4\%$ for G10W ($n=9$).

highlighted in Figure 9A. The energy-optimized model of the first 11 residues of the Kv β 1.3 N terminus is shown in Figure 9B. The side chain of R5 points towards A3 leading to a compact hairpin structure that would easily fit into the inner cavity of the Kv1.5 pore. This Kv β 1.3 structure was manually positioned within the confines of the Kv1.5 central cavity before calculating energy-minimized binding poses. Figure 9C illustrates the docking of Kv β 1.3 with a single Kv1.5 subunit. The residues in Kv1.5 described earlier as important for interaction with Kv β 1.3 (Decher *et al*, 2005) are highlighted with van der Waals surfaces. Figure 9D depicts the docking of Kv β 1.3 with two subunits, displaying important Kv1.5 residues as ball and stick model. A stereo-view of the docking with two Kv1.5 subunits is shown in Figure 9E. In the docking shown, the backbone of the Kv β 1.3 hairpin at position R5 and the residues T6 are in close proximity ($2.74 \text{ \AA}$) to T480 of the selectivity filter.

Next, we tested whether bulky side-chains at key residues in the N terminus of Kv β 1.3 affect inactivation. Introducing a tryptophan at positions R5 and T6 (at the tip of the proposed hairpin) enhanced inactivation (Figure 10A) as observed for other substitutions of these residues, consistent with the backbone of R5, and not its bulky side chain interacting with the selectivity filter. Kv β 1.3 has two Gly residues located at positions 7 and 10. Mutation of G10 to Ala or Cys (Figure 2) or Trp (Figure 10B) did not reduce the ability of Kv β 1.3 to induce inactivation. In contrast, although mutation of G7 to Ala had no functional consequence (Figure 2A), substitution with Cys significantly reduced inactivation (Figure 2B). Mutation of G7 to a much bulkier and hydrophobic Trp completely eliminated inactivation (Figure 10B), indicating the requirement for a small residue in this position located near the start of the hairpin loop.

Discussion

Occlusion of the central cavity by an inactivation peptide is the mechanism of rapid, N-type inactivation of Kv channels (Hoshi *et al*, 1990). Depending on the specific Kv channel, the

inactivation peptide can either be the N terminus of the Kv α -subunit or a separate, tethered Kv β subunit. Considering their common function, the N-terminal regions of Kv1.4, Kv3.4 or *Shaker* B α -subunits and the three Kv β 1 subunit isoforms have a surprisingly low sequence homology. NMR structures of Kv1.4 and Kv3.4 indicated earlier that Kv α inactivation peptides can adopt different tertiary structures. Using systematic site-directed mutagenesis, we studied the mode of binding of Kv β 1.3 subunits to Kv1.5 channels. Comparing earlier work with our new findings suggests that the mode of binding of Kv β 1.x subunits to Kv channels exhibit significant variability. We also found that Kv β 1 isoforms are differentially modulated by Ca^{2+} and PIP_2 .

We have identified an arginine residue (R5) located in the proximal N terminus of Kv β 1.3 subunits as a likely binding site for intracellular PIP_2 . Binding of PIPs to R5 prevents N-type inactivation mediated by Kv β 1.3. Although Kv β 1.1 is also sensitive to PIP_2 , the first 10 amino acids of this subunit do not include an arginine residue. Thus, the PIP_2 sensor of Kv β 1.1 remains to be discovered. In our lipid-binding assay, the N terminus of Kv β 1.3 binds PIP_2 with high affinity. For the N terminus of Kv β 1.3, we observed a strong PIP_2 -binding signal with 5 mol% of PIP_2 . With the same assay, addition of 10 and 35 mol% PIP_2 was required for significant binding to the Kv3.4 and Kv1.4 N termini (Oliver *et al*, 2004). In addition, we were able to show that a single residue substitution in the Kv β 1.3 N terminus can almost completely abolish PIP_2 -binding. When bound to PIP_2 , Kv β 1.3 may be positioned near the channel pore, but incapable of blocking the channel. This putative resting state might correlate with the pre-bound or pre-blocking state (O'), as was proposed earlier for Kv β 1 subunits (Zhou *et al*, 2001). Binding of Kv β 1.3 to the O' state might induce shifts in the voltage dependence of steady-state activation and C-type inactivation, even for mutant forms of Kv β 1.3 that are no longer capable of inducing N-type inactivation. The modulation of N-type inactivation in native Kv1.x–Kv β 1.3 complexes by PIP_2 might be important for the fine-tuning of neuronal excitability. As a result, fluctuations in intracellular PIP_2 levels due to Gq-coupled receptor stimulation might be relevant for the inactivation of K^+ channels and thus, for electrical signalling in the brain.

The variation in the amino-acid sequence of the proximal N termini also determines the different redox sensitivities of Kv β 1.1 and Kv β 1.3. Normally, Kv β 1.3 subunits are redox insensitive. However, we found that a single cysteine residue introduced at any position between amino acids 3–11 is sufficient to confer redox sensitivity to Kv β 1.3. Also in contrast to Kv β 1.1, we found that Kv β 1.3 was not sensitive to increased intracellular Ca^{2+} concentrations. Thus, an important physiological consequence of N-terminal splicing of the Kv β 1 gene might be the generation of rapidly inactivating channel complexes with different sensitivities to redox potential and intracellular Ca^{2+} .

We propose that Kv β 1.3 binds to the pore of Kv1.5 channels as a hairpin-like structure, similar to the N-terminal inactivation particles of Kv1.4 and Kv3.4 α -subunits (Antz *et al*, 1997). This is in contrast to Kv β 1.1, which was reported to bind to the central cavity of the Kv1 channel as a linear peptide (Zhou *et al*, 2001). For Kv β 1.1, interactions of residue 5 (Ile) were observed with sites in the distal S6 segment of Kv1.4, three helix turns distal to the PVP motif (Zhou *et al*,

2001). The interaction of R5 and T6 from Kvβ1.3 with the S6 segment residues high in the inner cavity and residues near the selectivity filter of Kv1.5 is only plausible if Kvβ1.3 blocks the channel as a small hairpin, as in the energy-minimized conformation illustrated in our model. The narrowing of the pore by the four S6 segments near the PVP motif with a diameter of 0.9–1.0 nm suggests that Kvβ1.3 can enter the inner cavity configured as a small hairpin. In addition, this hairpin structure is smaller than the N-terminal ball domains that were proposed earlier for the Kv1.4 and Kv3.4 N termini (Antz *et al*, 1997). On the basis of the crystal structures available, these inactivation balls are too large to pass the PVP barrier and enter the inner cavity. Accordingly, these N-terminal ball domains might bind more distally in the S6 segments and block the pore as 'shallow plugs' (Antz *et al*, 1997).

Mutation of R5 in Kvβ1.3 to E, C, A, Q and W accelerated the Kv1.5 channel inactivation. Thus, the acceleration of inactivation by R5 mutations is independent of the size and charge of the residue introduced. Together with our PIP₂-binding assay, these findings suggest that PIP₂ immobilizes Kvβ1.3 and prevents it from entering the central cavity to induce N-type inactivation. Our model predicts that the backbone of the hairpin, near R5, interacts with the selectivity filter. This is in good agreement with our observation that the nature of the side chain introduced at position 5 was not relevant for the blocking efficiency of the hairpin.

N-terminal splicing of Kvβ1 produces the Ca²⁺-insensitive Kvβ1.3 isoform that retains the ability to induce Kv1 channel inactivation. We propose that the N terminus of Kvβ1.3 exists in a pre-blocking state when PIPs located in the lipid membrane bind to R5. We further propose that when Kvβ1.3 dissociates from PIPs, it assumes a hairpin structure that can enter the central cavity of an open Kv1.5 channel to induce N-type inactivation.

Materials and methods

Molecular biology

Kv1.5 cDNA in the pSGEM oocyte expression vector and the methods of site-directed mutagenesis were described earlier (Decher *et al*, 2004). The Kv1.5 sequence (NM_002234) has an N terminus with two additional residues compared with an earlier database entry (M60451). This results in a shift of the amino acid numbering of +2 when compared with older literature. Restriction mapping and DNA sequencing were used to confirm the presence of the desired mutation and the lack of extra mutations in the PCR-amplified segment. Complementary RNA (cRNA) for injection into oocytes was prepared with T7 Capscribe (Roche) after linearization with *NheI*. The Kvβ1.3 construct in a modified pSP64T vector was described previously (England *et al*, 1995) and cRNA was made with SP6 Capscribe (Roche) after linearization with *EcoRI*. The quality and quantity of cRNA were determined by gel electrophoresis and UV spectroscopy.

Lipid-binding assays

For the lipid-binding assay, the nucleotide sequence encoding amino acids 1–33 of WT Kvβ1.3 and mutants R5C and T6C were subcloned with *EcoRI*–*Sall* into the pGEX4T-1 vector (Amersham Pharmacia Biotech) to produce an in-frame GST fusion protein. Proteins and liposomes were prepared and assayed as described (Soom *et al*, 2001). Briefly, GST, GST-fused Kvβ1.3 (residues 1–33), Kvβ1.3 (residues 1–33) R5C and Kvβ1.3 (residues 1–33) T6C were overexpressed in *Escherichia coli* strain BL-21 Codon Plus and immobilized on GSH Sepharose according to the manufacturer's instructions (Amersham Pharmacia Biotech). Mixed liposomes were prepared from PI(4,5)P₂, phosphatidylcholine (PC), phosphatidylethanolamine (PE), cholesterol (ChS) and rhodamine-PE (Rh-PE) to obtain a lipid composition of 5 mol% PI(4,5)P₂. The PE, ChS and Rh-PE contents were always 50, 32 and 1 mol%, respectively. Immobilized GST proteins (0.01 mM) were incubated with liposomes with subsequent washing. Binding of liposomes to immobilized proteins was quantified by fluorescence measurement using excitation/emission wavelengths of 390/590 nm (cutoff at 570 nm). The data were corrected by subtracting the fluorescence of control liposomes without PI(4,5)P₂ from the values obtained in assays with liposomes containing PI(4,5)P₂ and normalized to the binding of GST-fused Kvβ1.3 WT peptide. Results are presented as mean ± s.e.m. of three parallel experiments.

tidylethanolamine (PE), cholesterol (ChS) and rhodamine-PE (Rh-PE) to obtain a lipid composition of 5 mol% PI(4,5)P₂. The PE, ChS and Rh-PE contents were always 50, 32 and 1 mol%, respectively. Immobilized GST proteins (0.01 mM) were incubated with liposomes with subsequent washing. Binding of liposomes to immobilized proteins was quantified by fluorescence measurement using excitation/emission wavelengths of 390/590 nm (cutoff at 570 nm). The data were corrected by subtracting the fluorescence of control liposomes without PI(4,5)P₂ from the values obtained in assays with liposomes containing PI(4,5)P₂ and normalized to the binding of GST-fused Kvβ1.3 WT peptide. Results are presented as mean ± s.e.m. of three parallel experiments.

Two-electrode voltage-clamp

Stage IV and V *Xenopus laevis* oocytes were isolated and injected with cRNA encoding WT or mutant Kv1.5 and Kvβ1.3 subunits as described earlier (Decher *et al*, 2004). Oocytes were cultured in Barth's solution supplemented with 50 µg/ml gentamycin and 1 mM pyruvate at 18°C for 1–3 days before use. Barth's solution contained (in mM): 88 NaCl, 1 KCl, 0.4 CaCl₂, 0.33 Ca(NO₃)₂, 1 MgSO₄, 2.4 NaHCO₃, 10 HEPES (pH 7.4 with NaOH). For voltage-clamp experiments, oocytes were bathed in a modified ND96 solution containing (in mM): 96 NaCl, 4 KCl, 1 MgCl₂, 1 CaCl₂, 5 HEPES (pH 7.6 with NaOH). Currents were recorded at room temperature (23–25°C) with standard two-microelectrode voltage-clamp techniques (Stuhmer, 1992). The holding potential was –80 mV. The interpulse interval for all voltage-clamp protocols was 10 s or longer to allow for full recovery from inactivation between pulses. The standard protocol to obtain current–voltage (*I*–*V*) relationships and activation curves consisted of 200 ms or 1.5 s pulses that were applied in 10-mV increments between –70 and +70 mV, followed by a repolarizing step to –40 mV.

The voltage dependence of the Kv1.5 channel activation (with or without co-expression with Kvβ1.3) was determined from tail current analyses at –40 mV. The resulting relationship was fit to a Boltzmann equation (equation (1)) to obtain the half-point ($V_{1/2\text{act}}$) and slope factor (k_{act}).

$$I_n = \frac{1}{1 + \exp[(V_{1/2\text{act}} - V_t)/k_{\text{act}}]} \quad (1)$$

The voltage dependence of Kv1.5 inactivation was determined by using a two-pulse protocol. A prepulse of 1 s was applied to potentials ranging from –90 to +70 mV and was immediately followed by a 200 ms test pulse to +70 mV. The relative amplitude of peak current during the test pulse was plotted as a function of the prepulse voltage and the relationship fit to a Boltzmann function to obtain the $V_{1/2\text{inact}}$ for inactivation. Other voltage pulse protocols are described in the Results and figure legends. Data are expressed as mean ± s.e.m. (n = number of oocytes).

Excised macropatches from *Xenopus* oocytes

Recordings from inside-out macropatches were performed as described previously (Oliver *et al*, 2004). Pipettes (0.2–0.4 MΩ) were filled with extracellular solution (mM): 115 NaCl, 5 KCl, 10 HEPES and 1 CaCl₂ (pH 7.2 with NaOH). Intracellular solution contained (mM): 100 KCl, 10 EGTA and 10 HEPES (pH 7.2 with KOH). A hypertonic solution used to shrink oocytes and facilitate removal of the vitelline membrane contained (mM): 200 K-aspartate, 20 KCl, 1 MgCl₂, 10 EGTA and 10 HEPES (pH 7.4 with KOH).

Double-mutant cycle analysis

The double-mutant cycle parameter Ω (equation (2)) was calculated to quantify the degree of coupling between two mutations, as described previously (Hidalgo and MacKinnon, 1995; Gulbis *et al*, 2000).

$$\Omega = \frac{K_d^{\text{WT+WT}} \times K_d^{\text{mut+mut}}}{K_d^{\text{WT+mut}} \times K_d^{\text{mut+WT}}} \quad (2)$$

A value of Ω greater than unity indicates that the effects of two mutations are coupled. For Ω values smaller than 1, the reciprocal was taken to facilitate the display of changes from unity, as described previously (Hidalgo and MacKinnon, 1995). The K_d values were obtained from the apparent rate constants for binding

(α) and unbinding (β), according to:

$$K_d = \frac{\alpha}{\beta}$$

The α and β values were obtained from the inactivation time constant (τ_{in}) and the steady-state probability (ss) at the end of 1.5-s pulses. For the double-cycle analysis, only the predominant (fast) component of the inactivation was analysed:

$$\tau_{in} = \frac{1}{\alpha + \beta}$$

$$ss = \frac{\beta}{\alpha + \beta}$$

Molecular modelling and docking

Molecular modelling was based on the crystal structure of the mammalian Shaker Kv1.2 potassium channel subunit complex (PDB 2A79) and the sequence of potassium channel subunit Kv1.5

(NP_037104). A hairpin-shaped structural model of the first 11 residues of the Kvβ1.3 N terminus (AAD37855) was built and energy-optimized (Figure 9B). Interactions of the Kvβ1.3 N terminus and Kv1.5 were determined by molecular docking. The docking was initiated from random configurations ($n=200$) of the Kvβ1.3 N terminus in the cytoplasmic cavity of the Kv1.5 tetramer and performed by using simulated annealing techniques with an initial temperature of 500 K and a final temperature of 300 K. Residues of Kv1.5 facing the pore region and the Kvβ1.3 N terminus were flexible during the optimization procedure. Modelling and docking were performed with the Insight II modules Homology, Builder and Docking (version 8.2; Accelrys, San Diego, CA).

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