Voltage Clamp Fluorimetry Reveals a Novel Outer Pore Instability in a Mammalian Voltage-gated Potassium Channel

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INTRODUCTION

Voltage-gated potassium (Kv) channel gating involves complex structural rearrangements that regulate the ability of channels to conduct K⁺ ions. Fluorescence-based approaches provide a powerful technique to directly report structural dynamics underlying these gating processes in Shaker Kv channels. Here, we apply voltage clamp fluorimetry, for the first time, to study voltage sensor motions in mammalian Kv1.5 channels. Despite the homology between Kv1.5 and the Shaker channel, attaching TMRM or PyMPO fluorescent probes to substituted cysteine residues in the S3–S4 linker of Kv1.5 (M394C-V401C) revealed unique and unusual fluorescence signals. Whereas the fluorescence during voltage sensor movement in Shaker channels was monoexponential and occurred with a similar time course to ionic current activation, the fluorescence report of Kv1.5 voltage sensor motions was transient with a prominent rapidly dequenching component that, with TMRM at A397C (equivalent to Shaker A359C), represented 36 ± 3% of the total signal and occurred with a τ of 3.4 ± 0.6 ms at +60 mV (n = 4). Using a number of approaches, including 4-AP drug block and the ILT triple mutation, which dissociate channel opening from voltage sensor movement, we demonstrate that the unique dequenching component of fluorescence is associated with channel opening. By regulating the outer pore structure using raised (99 mM) external K⁺ to stabilize the conducting configuration of the selectivity filter, or the mutations W472F (equivalent to Shaker W434F) and H463G to stabilize the nonconducting (P-type inactivated) configuration of the selectivity filter, we show that the dequenching of fluorescence reflects rapid structural events at the selectivity filter gate rather than the intracellular pore gate.

Fluorescence-based methods have allowed direct observation of ion channel gating conformations. Using voltage clamp fluorimetry (VCF) and the archetypal Shaker Kv channel, simultaneous recording of ionic current along with changes in fluorescence emission from tetramethylrhodamine-5-maleimide (TMRM) attached in the S3–S4 linker has shown that the fluorescence changes provide a faithful report of the time and voltage dependence of voltage sensor movements (Mannuzzu et al., 1996; Cha and Bezanilla, 1997; Pathak et al., 2007). Two sites in particular in the S3–S4 linker, M356 and A359, have proven particularly useful in fluorimetric studies. Similar fluorescence reports of voltage sensor dynamics have also been described in hSkM1 Na⁺ channels, hERG, eag and BK Ca₂⁺ channels, and HCN channels (Cha et al., 1999; Smith and Yellen, 2002; Bannister et al., 2005; Savalli et al., 2006; Bruening-Wright et al., 2007).

In addition to the observation of voltage sensor motions, some of these fluorescence studies report additional conformational changes that occur after channel activation. These rearrangements reflect transitions that are associated with opening of the channel and/or collapse of the outer pore during inactivation. TMRM attached within the S3–S4 linker of the Shaker channel detects not only the independent movement of the voltage sensors, but also conformational rearrangements.
that are coupled with the concerted opening of the intracellular pore gate (Pathak et al., 2005). A number of sites within the S3–S4 linker also report the slow collapse of the outer pore that occurs during P/C-type inactivation (Loots and Isacoff, 1998, 2000; Gandhi et al., 2006). In hERG channels, slow TMRM fluorescence was reported to occur upon depolarization. Using 4-aminopyridine (4-AP) and the ILT triple mutation (V407I/I410L/S414T) to uncouple voltage sensor activation from opening of the intracellular pore gate, we demonstrate that this rapid unquenching of the fluorescence signal is associated with channel opening. Furthermore, by raising the external K+ concentration, or introducing the W472F and H463G mutations, or the outer pore mutation, R487V, which all regulate or immobilize the outer pore gate while sparing normal functioning of the intracellular pore gate, we show that the unique fluorescence component measured in Kv1.5 channels is associated with gating at the selectivity filter.

MATERIALS AND METHODS

Molecular Biology

The vector pEXO was used to express Kv1.5 and Shaker Δ46-46 channels in Xenopus laevis oocytes. Cysteine residues were introduced at specific sites (M394C to V401C in Kv1.5, A359C in Shaker) in the S3–S4 linker for TMRM labeling and the only externally accessible cysteine residue, found in the S1–S2 linker, was replaced with a valine residue (C268V in Kv1.5, C245V in Shaker) to prevent nonspecific dye labeling. The deletion mutation Δ46-46 was introduced into Shaker channels to remove fast inactivation (Hoshi et al., 1990). The W472F mutation (equivalent to W434F in Shaker channels) was introduced to permanently inactivate channels (Perozo et al., 1993; Yang et al., 1997). To dissociate voltage sensor movement from channel opening, the ILT triple mutation (V407I, I410L, and S414T) was introduced in the S4 domain (Smith-Maxwell et al., 1998a, b). The R487V outer pore mutation, which all regulate or immobilize the outer pore gate while sparing normal functioning of the intracellular pore gate, was shown to dissociate voltage sensor movement from channel opening.

Oocyte Preparation

Xenopus laevis oocytes were prepared and isolated as previously (Claydon et al., 2007a). Following removal of the follicular layer, oocytes were injected with cRNA and allowed to recover for at least 30 minutes in Barth’s solution supplemented with 0.1% BSA. The injection was performed using a micromanipulator (Eppendorf Micromanipulator) and a glass capillary (World Precision Instruments). The oocytes were then transferred to a recording chamber and allowed to equilibrate for at least 1 hour before the start of the measurements. The recording chamber was maintained at 18°C and continuously perfused with physiological saline (115 mM NaCl, 2.5 mM KCl, 1.8 mM CaCl2, 1 mM MgCl2, 5 mM HEPES, pH 7.4). The oocytes were held at −80 mV and voltage-clamped using a patch clamp amplifier (Axopatch 200B, Axon Instruments). The data were acquired using pClamp 9 software (Axon Instruments) and analyzed using Clampfit 9.2 software (Axon Instruments). The G-V relations of Kv1.5 WT and Kv1.5 C268V were significantly different (p < 0.01, unpaired t-test). Note that error bars generally fall within the plotted points in C and F.

Figure 1. TMRM fluorescence from Kv1.5 WT and Kv1.5 C268V channels. (A, B, D, and E) Ionic current (A and D) and fluorescence (B and E) traces recorded from oocytes expressing Kv1.5 WT (A and B) or Kv1.5 C268V (D and E) channels labeled with TMRM. Voltage clamp pulses were applied from −80 to +60 mV in 10-mV increments (100 ms duration) from a holding potential of −80 mV (only +60 mV traces are shown for clarity). (C and F) Mean G-V relations for Kv1.5 WT (C; n = 9) and Kv1.5 C268V (F; n = 4). V1/2 and k values for the G-V relation of Kv1.5 channels were 1.2 ± 1.3 and 12.5 ± 1.0 mV, respectively. The V1/2 and k values for the G-V relations of Kv1.5 WT and Kv1.5 C268V were −5.7 ± 1.0 and 10.8 ± 0.8 mV, respectively. The V1/2 values for the G-V relations of Kv1.5 WT and Kv1.5 C268V are significantly different (p < 0.01 (unpaired t-test). Note that error bars generally fall within the plotted points in C and F.
Data Analysis

Conductance–voltage (G-V) and fluorescence–voltage (F-V) relations were fitted with a single Boltzmann function: \( y = \frac{1}{1 + \exp \left( \frac{V - V_{1/2}}{k} \right)} \), where \( y \) is the conductance or fluorescence amplitude normalized with respect to maximal conductance or fluorescence amplitude, \( V_{1/2} \) is the half-activation potential, \( V \) is the test voltage, and \( k \) is the slope factor.

RESULTS

The Fluorescence Report of Voltage Sensor Movement in Kv1.5 Channels Is Unique

Kv1 channels have an extracellular cysteine residue in the S1–S2 linker that is a potential site for TMRM labeling and that may fluoresce in addition to any site that is engineered in the S3–S4 linker for gating studies. Control experiments in Fig. 1 showed that neither wild-type Kv1.5 (Fig. 1, A–C), nor Kv1.5 C268V (Fig. 1, D–F), showed a fluorescence report upon depolarization despite large expressed current amplitudes (Fig. 1, A and D). The G-V relations are shown in Fig. 1 (C and F) and the \( V_{1/2} \) and \( k \) values were 1.2 ± 1.3 and 12.5 ± 0.8 mV for wild type and −5.6 ± 1.0 and 10.8 ± 0.8 mV for C268V. All subsequent fluorescence experiments were performed using the Kv1.5 C268V mutation in addition to a fluorophore site as stated.

Dynamic changes in the fluorescence report from TMRM attached within the S3–S4 linker at A359C in Shaker channels provide a faithful report of the time and...
voltage dependence of voltage sensor motions induced by changes in the transmembrane potential (Mannuzzu et al., 1996; Cha and Bezanilla, 1997; Pathak et al., 2007). Fig. 2 (A–C) shows a characteristic example of the fluorescence reports from Shaker A359C during 100-ms voltage clamp pulses to a range of potentials. Simultaneous recordings of ionic current (Fig. 2 A) and TMRM fluorescence (Fig. 2 B) show that the fluorophore in Shaker reports a monoexponential quenching upon depolarization that occurs with a time course ($\tau = 2.4 \pm 0.5$ ms at +60 mV) that is similar to, but faster than, that of the activation of ionic current ($\tau = 4.5 \pm 1.0$ ms at +60 mV; $n = 4$). The conformational changes reported by the fluorophore display a voltage dependence ($V_{1/2} = -40.9 \pm 1.9$ mV) that is $\sim 25$ mV left shifted from the G-V relation ($V_{1/2} = -16.2 \pm 1.3$ mV; Fig. 2 C), as is expected for movement of the voltage sensors in advance of channel opening. Similar monoexponential reports of voltage sensor movement that precede at least one component of ionic current activation kinetics have also been reported in hERG, eag, BKCa, and HCN channels (Smith and Yellen, 2002; Bannister et al., 2005; Savalli et al., 2006; Bruening-Wright et al., 2007). Many of the clearest recordings in the above studies have come from Shaker A359C or the equivalent residues to those in other channels (e.g., L520C in hERG channels; Smith and Yellen, 2002).

In Fig. 2 (D–F), we show, for the first time, the fluorescence report of voltage sensor movement in a mammalian Kv1 channel, Kv1.5. The fluorescence report is from TMRM attached at A397C, which is the equivalent S3–S4 linker site to A359C in Shaker channels. In contrast to the monoexponential time course of the fluorescence quenching from Shaker (Fig. 2 B) and other Kv channels (Smith and Yellen, 2002; Bannister et al., 2005; Savalli et al., 2006), TMRM attached at Kv1.5 A397C...
reported a unique profile of fluorescence change (Fig. 2 E). Upon depolarization, the fluorescence quenching deflection from Kv1.5 A397C was rapid, but transient with a prominent dequenching component that reached a steady-state level by the end of the 100-ms voltage pulse. The recovering component of fluorescence was only observable at relatively depolarized potentials and became faster and larger with stronger depolarizations. Another difference from Shaker A359C (Fig. 2 C) was that the F-V relation of Kv1.5 A397C fluorescence, taken from the peak of the fluorescence deflections, was not significantly left-shifted ($V_{1/2} = 1.9 \pm 1.9$ mV) from the G-V relation ($V_{1/2} = 7.3 \pm 1.6$ mV; Fig. 2 F). Furthermore, the voltage dependence of the dequenching component amplitude was actually right-shifted ($V_{1/2} = 51.0 \pm 2.4$ mV) from the G-V relation (Fig. 2 F). These observations suggest that at least some component of the fluorescence deflection from TMRM attached at Kv1.5 A397C may be coupled in some way with channel opening.

**Spectroscopic Mapping of Kv1.5 Voltage Sensor Dynamics—Discrete vs. Global Rearrangements**

The different report of fluorophore environmental change observed in Kv1.5 (Fig. 2, D–F) may suggest that very different voltage sensor conformational changes occur in these channels than in Shaker channels. However, a fluorescence scan of the S3–S4 linker of Kv1.5 (Fig. 3) reveals that the complex structural changes are only reported by TMRM at two sites, A397C and M394C. This cannot be accounted for by differences in ionic current waveforms (Fig. 3 A) or changes in the G-V relationships with the different linker mutants (Fig. 3 C), since these all appear similar. Both A397C and M394C report a voltage-dependent acceleration of the dequenching component and an increase in its contribution to the total fluorescence change upon depolarization (Fig. 4). Other sites in the scan do not report the rapid component of fluorescence and produce deflections that are, in general, similar to those described from the equivalent sites in Shaker channels (Gandhi et al., 2000; Pathak et al., 2007). This suggests that Kv1.5 voltage sensor movement per se is not dissimilar to that of Shaker channels. Furthermore, given that the functional data suggests that the C-terminal end of the S3–S4 linker likely adopts an α-helical structure, placing M394C and A397C on the same side of the helix, (Li-Smerin et al., 2000; Gonzalez et al., 2001; Li-Smerin and Swartz, 2001), and at least M394C and A397C sense the same microenvironment changes, these data suggest that the fluorescence changes represent local protein rearrangements rather than global reconfigurations. Since M394C and A397C come close to the pore domain upon depolarization (Gandhi et al., 2000; Elinder et al., 2001), we hypothesized that TMRM attached at M394C or A397C might be reporting on an additional structural rearrangement that is associated with opening of the pore.

In support of this, the time constant data in Fig. 4 A show a reasonably good match between the time constants of channel activation (filled symbols) and the fast component of fluorescence dequenching, especially for M394C (open triangles).

A similar fluorescence profile of channel gating was reported by another fluorophore, PyMPO (Fig. 5), that has different physical and spectral properties to TMRM. Ionic currents (Fig. 5 A) and the G-V relationship (Fig. 5 C), which had a $V_{1/2}$ and $k$ of $2.3 \pm 1.9$ and $17.8 \pm 1.3$ mV, respectively, were relatively unaffected by labeling with PyMPO. The fluorescence reports were very similar to those seen with TMRM, and included a transient component that appeared at around 0 mV, a potential at which significant numbers of channels opened (Fig. 5 B). It was noted that PyMPO fluorescence recordings bleached more rapidly during clamp pulses, and perfect baseline correction was not possible when full F-V relations were obtained. This explains the baseline drift seen in recordings at 0 and +30 mV in Fig. 5 B. With TMRM, the time constant of the fluorescence dequenching at +60 mV was $3.4 \pm 0.6$ ms and it represented $36 \pm 3\%$ of the total signal (Fig. 4; $n = 4$). The corresponding values with PyMPO were similar (Fig. 5 D); the time
constant was $2.1 \pm 0.5$ ms and it represented $25 \pm 8\%$ of the total signal ($n = 3$), suggesting that the fluorescence changes observed are a faithful report of protein conformational changes. Upon repolarization, the return of both the TMRM and PyMPO fluorescence signals was rapid (see also Fig. 2 E), and showed no evidence of return of the rapid component of fluorescence. This suggests that the motions represented by the dequenching of fluorescence upon depolarization are slow to recover upon repolarization.

The Dequenching Component of Fluorescence Is Associated with Transitions to the Open State

As a first test of this hypothesis, we compared the effect of the holding potential on the fluorescence report from Kv1.5 A397C channels (Fig. 6). If the transient dequenching component of fluorescence is associated with late transitions to the open state rather than voltage sensor transitions early in the activation pathway, the fluorescence report from channels that open from hyperpolarized potentials (i.e., $-120$ mV) should be the same as that from channels opening from preactivated states (i.e., $-40$ mV). Data in Fig. 6 A shows ionic currents and fluorescence signals from TMRM attached to A397C recorded during a test pulse applied immediately after a conditioning pulse to either $-120$ or $-40$ mV. The ionic current records show that channel opening was more rapid following a conditioning pulse to $-40$ mV, which collects channels in preactivated states, than following a pulse to $-120$ mV, which collects channels in early closed states. Consistent with this, the fluorescence signal recorded following a prepulse to $-40$ mV dequenched more rapidly than following a pulse to $-120$ mV. However, the unique fluorescence profile that we observed in Kv1.5 channels was maintained following a prepulse to $-40$ mV, and the relative amplitude of the rapid fluorescence dequenching was unchanged by alterations in the holding potential (Fig. 6 B).

As a further test of this hypothesis, we investigated the effects of preventing opening on the fluorescence report from Kv1.5 A397C channel. We measured the fluorescence report in the presence of $10$ mM 4-AP or the ILT triple mutation (V407I, I410L, S414T), both of which dissociate voltage sensor movement from channel opening by stabilizing channels in preopen closed states (McCormack et al., 1994; Smith-Maxwell et al., 1998a, b; del Camino et al., 2005; Pathak et al., 2005). $10$ mM 4-AP reduced the ionic current by about two thirds and abolished the rapid component of fluorescence (Fig. 7, A and B), reverting the fluorescence phenotype to that reported by Shaker A359C (compare the fluorescence trace in the presence of 4-AP with that in Fig. 2 B). In addition, 4-AP restored the left-shifted voltage dependence of the F-V relation (the $V_{1/2}$ of the F-V was $\sim 41$ mV left shifted from that of the G-V; Fig. 7 C). In a similar manner, at potentials (e.g., $+60$ mV) that stabilize ILT mutant channels in the activated-not-open state (del Camino et al., 2005), where little ionic current is
observed (Fig. 7 D), the dequenching component of fluorescence from Kv1.5 ILT A397C channels was abolished and the Shaker-like monoexponential report of voltage sensor movement was rescued (Fig. 7 E). The F-V relation in ILT mutant channels was also left shifted from the G-V relation (the $V_{1/2}$ of the F-V was $-41.6 \pm 1.8$ mV; Fig. 7 F). These data clearly demonstrate that the dequenching component of fluorescence from A397C is associated with channel opening. Furthermore, these results show that the depolarized position of the F-V relation in Fig. 2 F does not reflect the true voltage dependence of voltage sensor movement, because it is influenced by an event that TMRM detects upon channel opening.

The Dequenching Component of Fluorescence Is Associated with Structural Events at the Selectivity Filter Gate

The observation in Fig. 2 E that the fluorescence signal does not report the return of the dequenching component upon repolarization suggests that the protein motions reported upon are relatively slow to recover. To investigate the time course of the recovery, we performed a double pulse protocol in which we applied a 75-ms test pulse to $+60$ mV at increasing intervals following a 15-ms conditioning pulse to $+60$ mV (Fig. 8 A). The dequenching phase of fluorescence was relatively slow to recover as shown in Fig. 8 B. After 7.5 ms repolarization, the amplitude of dequenching in the test pulse was significantly reduced, but increased with increasing rest interval duration. The time dependence of the return of the rapid component is plotted in Fig. 8 C, and the time constant for the return was $20.7 \pm 3.7$ ms ($n = 3$). This is significantly slower than the time constant of intracellular pore gate closure during deactivation in Kv1.5 channels, which is $\sim 5$ ms at these potentials (Chen et al., 1997), but since the experiments in Fig. 8 were performed in 3 mM extracellular $K^+$, it was not possible to resolve the time course of the decay of the tail currents directly at $-80$ mV. However, the time course of recovery of the instantaneous ionic current level from the same experiments is a reasonable index of channel closing, indicated by the dotted line in Fig. 8 A. The mean data show a recovery time constant of $10.7 \pm 0.8$ ms ($n = 3$), which is significantly faster than the recovery of fluorescence ($P < 0.05$).

The slow return of the fluorescence upon repolarization suggests that although associated with channel opening, the dequenching component of fluorescence does not reflect gating of the intracellular pore. We also observed that the recovery of this phase was dependent upon the conditioning pulse duration (Fig. 9). Following longer conditioning pulses ($15$ ms; Fig. 9 A) that allowed significant channel opening, the dequenching component was almost absent in the first test pulse, whereas following shorter conditioning pulses ($5$ ms; Fig. 9 B) that restricted channel opening, there was a robust component in the first test pulse. These data further demonstrate that the slowly recovering fluorescence dequenching is related to channel opening. Based on these data, we considered that the rapid fluorescence changes might reflect structural events occurring at the outer pore gate upon opening.

The Dequenching Component of Fluorescence Does Not Reflect Gating at the Intracellular Pore Gate

The observation in Fig. 2 E that the fluorescence signal does not report the return of the dequenching component upon repolarization suggests that the protein motions reported upon are relatively slow to recover. To investigate the time course of the recovery, we performed a double pulse protocol in which we applied a 75-ms test pulse to $+60$ mV at increasing intervals following a 15-ms conditioning pulse to $+60$ mV (Fig. 8 A). The dequenching phase of fluorescence was relatively slow to recover as shown in Fig. 8 B. After 7.5 ms repolarization, the amplitude of dequenching in the test pulse was significantly reduced, but increased with increasing rest interval duration. The time dependence of the return of the rapid component is plotted in Fig. 8 C, and the time constant for the return was $20.7 \pm 3.7$ ms ($n = 3$). This is significantly slower than the time constant of intracellular pore gate closure during deactivation in Kv1.5 channels, which is $\sim 5$ ms at these potentials (Chen et al., 1997), but since the experiments in Fig. 8 were performed in 3 mM extracellular $K^+$, it was not possible to resolve the time course of the decay of the tail currents directly at $-80$ mV. However, the time course of recovery of the instantaneous ionic current level from the same experiments is a reasonable index of channel closing, indicated by the dotted line in Fig. 8 A. The mean data show a recovery time constant of $10.7 \pm 0.8$ ms ($n = 3$), which is significantly faster than the recovery of fluorescence ($P < 0.05$).

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Figure 6. The dequenching of fluorescence is associated with late transitions in the activation pathway. (A) Ionic currents and fluorescence signals recorded from TMRM attached at A397C during a 100-ms test pulse to $+60$ mV following a 100-ms conditioning pulse to either $-120$ or $-40$ mV. The dequenching component of fluorescence remains robust in channels that activate from preactivated states, suggesting that the dequenching component does not reflect voltage sensor transitions early in the activation pathway. (B) Mean values for the time constant and relative amplitude of the dequenching component of fluorescence during a test pulse to $+60$ mV from a range of prepulse potentials ($n = 3$).
also prevents selectivity filter gating events, including inactivation (Castle et al., 1994; Claydon et al., 2007a). In the ILT experiments it was difficult to assess whether the function of the selectivity filter inactivation gate of ILT channels is still intact at the strongly depolarized potentials (+150 mV) required to drive channels into the open state, as activation was still slow at the maximum potentials studied in the experiments. Therefore, further experiments were performed using interventions and mutations known to affect the inactivation process or modulate outer pore gating in Kv1 channels, while measuring the fluorescence report from A397C.

There is strong evidence for a K⁺ modulatory site within the outer pore of Kv channels that, when occupied, prevents collapse of the selectivity filter, acting with a “foot-in-the-door” mechanism (Pardo et al., 1992; Lopez-Barneo et al., 1993; Baukrowitz and Yellen, 1995; Rasmusson et al., 1995; Kiss and Korn, 1998). In some Kv1 channels, including Kv1.5 (Fedida et al., 1999), this regulation does not appear to be as strong as in Shaker channels, but raising the extracellular K⁺ did significantly modulate the fluorescence report (Fig. 10). The ionic current amplitudes and time course are relatively unchanged at +60 mV, despite the reduction in driving force, as reported before for some Kv1 channels (Pardo et al., 1992). The fluorescence signals in Fig. 10 B show that raising the external K⁺ concentration to 99 mM abolished the dequenching component and restored more of a Shaker-like fluorescence report of voltage sensor movement. The fluorescence report is not identical to that in Shaker (Fig. 2), though, and appears to show both a fast and slow component of quenching, but with 99 mM K⁺ the left-shifted F-V relation of voltage sensor movement (V_{1/2} = −36.4 ± 1.2 mV) becomes apparent (Fig. 10 C; n = 3). Clearly, K⁺ regulation at an outer pore site is able to modulate the fluorescence environment change detected at A397C, and allow detection of voltage sensor movement.

To test outer pore regulation in another way, we introduced the mutation W472F, which is the equivalent to W434F in the Shaker channel, and which immobilizes the selectivity filter gate in a permanently nonconducting state in both Shaker (Perozo et al., 1993; Yang et al., 1997) and Kv1.5 channels (Wang et al., 1999; Wang and Fedida, 2001). Importantly, this mutation is not thought to immobilize the intracellular activation gate, as shown previously (Starkus et al., 1998). The tracings in Fig. 10 D show that although no ionic current was seen with this mutant, large fluorescence deflections were observed. In addition, the transient fluorescence change and thus structural events reported by the fluorescence at A397C were abolished. Taken together, these data strongly suggest that the dequenching component of fluorescence reported by TMRM attached at Kv1.5 A397C reflects rapid structural events at the selectivity filter gate that occur upon opening.

Closure of the selectivity filter gate underlies P/C-type inactivation in Kv channels (De Biasi et al., 1993;
Lopez-Barneo et al., 1993; Yellen, 1998; Kiss et al., 1999; Kurata and Fedida, 2006; Cordero-Morales et al., 2007. Although the time course of the fluorescence dequenching in Fig. 2 E is very rapid (3.4 ms at +60 mV) compared with the time course of P/C-type inactivation, which occurs in the order of seconds, we considered whether the novel fluorescence report from TMRM attached at A397C might reflect selectivity filter rearrangements that are associated in some way with inactivation. To test this, we examined the effects on the rapid component of fluorescence in Kv1.5 channels of the introduction of the outer pore mutation R487V (equivalent to T449V in Shaker), which is known to inhibit P/C-type inactivation in Shaker (Lopez-Barneo et al., 1993), and to abolish inactivation in Na+ -conducting Kv1.5 channels (Wang et al., 2000a,b). Data in Fig. 11 (A and B) show a typical ionic current and fluorescence signal recorded from TMRM attached at A397C in Kv1.5 R487V mutant channels during a 100-ms voltage clamp pulse to +60 mV. The tracing clearly shows that the dequenching component of fluorescence is maintained in the R487V mutation.

Other mutations in Shaker channels that modulate slow inactivation include E418 and V463 (Hoshi et al., 1991; Larsson and Elinder, 2000), but unfortunately we were unable to obtain current expression or fluorescence recordings in oocytes from the equivalent residues E456Q or E456C in Kv1.5 (see also Eduljee et al., 2007), or the equivalent residues to V463 in Shaker. However, another residue that modulates P/C-type inactivation in Kv1.5 is the His in the pore turret (Kehl et al., 2002). This mutation accelerated P/C-type inactivation as seen in the current record in Fig. 11 C, and also resulted in very low current expression levels, which was almost certainly due to both accelerated open-state and closed-state inactivation of the large population of the channels that do not pass current, as reported from our group and others previously (Jäger and Grissmer, 2001; Kehl et al., 2002; Zhang et al., 2005). This conclusion is supported by the robust fluorescence report from these channels (Fig. 11 D). This mutation completely removed the rapid dequenching phase of fluorescence at all potentials and showed a slow second phase of fluorescence quenching that matches the accelerated current inactivation seen in Fig. 11 C. The result suggests that collapse or immobilization of the selectivity filter gate before channel opening in H463G, as also seen in W472F (Fig. 10 D), prevents the structural change that causes the rapid fluorescence dequenching and allows unimpeded detection of voltage sensor movement.

**DISCUSSION**

**Kv1.5 Fluorescence Recording from A397C**

In this description of the first fluorescence reports of the conformational rearrangements associated with gating of a mammalian Kv1 channel, we observed complex fluorescence changes in the report of voltage sensor movements in Kv1.5 channels. Unlike in the Shaker channel, the rapid fluorescence quenching associated with activation of the voltage sensors upon depolarization was followed by a rapid dequenching of the fluorescence signal (Fig. 2). Furthermore, this fluorescence
profile was reported by two different fluorophore probes, TMRM and PyMPO (Figs. 2 and 5). This complex report of environmental changes occurring around the fluorophore attached at A397C (and M394C) upon depolarization (Figs. 2–5) suggests unique and complex motions of the voltage sensor domains in Kv1.5 channels that, at face value, are quite different from those reported in Shaker channels (Mannuzzi et al., 1996; Cha and Bezanilla, 1997). However, only sites that face the pore domain (A397C and M394C) report the complex changes; many of the sites that do not report the rapid dequenching of fluorescence have F-V relations that are well left shifted from the G-V relations (the $V_{1/2}$ values for the F-V relations of S395C, L396C, and L399C were 64, 12, and 48 mV left shifted from the corresponding $V_{1/2}$ values for the G-V relations) as would be predicted for a faithful report of Shaker-like voltage sensor movement (Fig. 3). Furthermore, prevention of the dequenching component of fluorescence from TMRM at A397C (Fig. 7, Fig. 10 D, and Fig. 11 D) unmasks a monophasic Shaker-like report of voltage sensor movement. Taken together, these observations suggest that although the fluorescence report from Kv1.5 channels is unique and unusual, voltage sensor movement per se in Kv1.5 channels is not dramatically different from that in the Shaker channel.

Clearly however, the data from Fig. 4 demonstrate that the fluorescence signal from TMRM attached to the S3–S4 linker reflects a voltage-dependent channel rearrangement that occurs rapidly at depolarized potentials ($\tau = 3.4$ ms at $+60$ mV) and is relatively slow to recover ($\tau = 21$ ms; Fig. 8 C). The observation that a
close proximity with the outer pore S5-P linker of its adjacent subunit upon depolarization (Blaustein et al., 2000; Gandhi et al., 2000; Larsson and Elinder, 2000; Li-Smerin et al., 2000; Loots and Isacoff, 2000; Ortega-Saenz et al., 2000; Elinder et al., 2001; Laine et al., 2003; Long et al., 2005b), and is supported by previous observations of structural changes reported by sites within the S3–S4 linker that are associated with opening of the intracellular gate and/or inactivation of the outer pore (Loots and Isacoff, 1998, 2000; Gandhi et al., 2000; Smith and Yellen, 2002; Pathak et al., 2005; Savalli et al., 2006; Claydon et al., 2007a, b).

The W434F mutation in \textit{Shaker} channels (equivalent to W472F in Kv1.5) permanently immobilizes the selectivity filter gate in the P-type inactivated conformation (Perozo et al., 1993; Yang et al., 1997). However, in the absence of K\(^+\), channels are capable of conducting Na\(^+\) (Starkus et al., 1998; Wang and Fedida, 2001). This demonstrates the preservation of intracellular pore gate functionality in W434F/W472F mutant \textit{Shaker} and Kv1.5 channels despite the effects at the selectivity filter gate. Our observation in Fig. 10 D that the W472F mutation abolishes the rapid dequenching component of fluorescence in Kv1.5 channels suggests therefore that the secondary dequenching represents a gating event taking place at the selectivity filter gate rather than the intracellular pore gate, since it is the outer pore gate and not the inner pore gate that is immobilized in this channel. The rapid Kv1.5 fluorescence signal does not reflect channel opening per se, but events at the selectivity filter that occur rapidly upon opening. Because W472F channels are thought to be P-type inactivated at rest (which is why they are nonconducting to K\(^+\)), as are the majority of H463G channels (Fig. 11, C and D), these observations raise the possibility that the rapid dequenching event is related to the onset of P-type inactivation in the channels.

The Secondary Dequenching Phase of Fluorescence from Kv1.5 A397C

We presented a series of experiments that strongly suggest that the secondary phase of fluorescence from TMRM-labeled Kv1.5 channels upon depolarization is a report of a gating event that is associated with channel opening (Fig. 2 F and Figs. 4, 6, and 7). By manipulating the holding potential, we demonstrated that labeled channels continued to display a robust secondary dequenching component when activated from preopening potentials (Fig. 6). This suggests that this phase of fluorescence reflects transitions late in the activation pathway that are associated with concerted opening of the channel, or even later events. This is further supported by evidence that binding of 4-AP or introduction of the ILT triple mutations, which both stabilize channels in the activated-not-open state upon depolarization without greatly altering early activation transitions (McCormack et al., 1994; Smith-Maxwell et al., 1998a,b; Armstrong and Loboda, 2001; Loboda and Armstrong, 2001; del Camino et al., 2005), abolished the dequenching of fluorescence (Fig. 7). This conclusion is consistent with electrophysiological and crystallographic data, which demonstrate that each voltage sensor comes into close proximity with the outer pore S5-P linker of its adjacent subunit upon depolarization (Blaustein et al., 2000; Gandhi et al., 2000; Larsson and Elinder, 2000; Li-Smerin et al., 2000; Loots and Isacoff, 2000; Ortega-Saenz et al., 2000; Elinder et al., 2001; Laine et al., 2003; Long et al., 2005b), and is supported by previous observations of structural changes reported by sites within the S3–S4 linker that are associated with opening of the intracellular gate and/or inactivation of the outer pore (Loots and Isacoff, 1998, 2000; Gandhi et al., 2000; Smith and Yellen, 2002; Pathak et al., 2005; Savalli et al., 2006; Claydon et al., 2007a,b).
An Outer Pore Regulatory Event upon Opening, or More Specifically P-type Inactivation? Secondary fluorescence deflections associated with rearrangements of the outer pore that occur in addition to activation of the voltage sensors upon depolarization have been previously reported from the equivalent site to A397C (A359C) in Shaker channels (Loots and Isacoff, 1998, 2000; Cha and Bezanilla, 1998; Claydon et al., 2007a,b). However, the rapid phase of fluorescence dequenching that we report in the present study from Kv1.5 channels is three orders of magnitude faster than the slow secondary movement associated with P/C-type inactivation reported by the fluorescence signal from Shaker A359C, and is only reflected by one side of the S3–S4 linker helix, whereas inactivation rearrangements in Shaker are sensed by all faces of the S3–S4 linker helix (Figs. 2 and 3) (Loots and Isacoff, 1998, 2000; Claydon et al., 2007b). As well, fluorescence events associated with P/C-type inactivation reported by Shaker A359C or Kv1.5 A397C (Claydon et al., 2007b) usually result in a slow phase of fluorescence quenching, as also seen in the present study in Kv1.5 A397C H463G (Fig. 11 D), rather than a rapid phase of fluorescence dequenching. Finally, and importantly, there is no obvious rapid activation of ionic current evident in Kv1.5 current traces with the time course of the rapid fluorescence event (e.g., Fig. 2 D). These discrepancies suggest that the rapid dequenching component reported by TMRM and PyMPO attached in Kv1.5 is not simply a report of outer pore movements associated with channel inactivation.

Although we observed that increased K+ at an outer pore regulatory site, and P-type immobilization of the selectivity filter gate in the mutation W472F, and H463G prevented the fluorescence dequenching (Figs. 10 and 11), the inability of the outer pore mutation R487V (Lopez-Barneo et al., 1993), which inhibits inactivation, to prevent this rapid component of fluorescence (Fig. 11 A) supports the idea that the rapid fluorescence dequenching does not represent the movements associated with P/C-type inactivation in this channel. This last observation must be interpreted with care, as in Kv1.5, the mutation R487V inhibits some events associated with P/C-type inactivation, but not all of them (Fedida et al., 1999; Wang et al., 2000b). Thus, the event may reflect a preinactivation transition that is not prevented by the outer pore mutation R487V. Alternatively, the observations that raising external K+ and not the R487V mutation prevented the secondary fluorescence signal (Fig. 10, A–C, and Fig. 11) may also suggest that the rapid dequenching of fluorescence reflects dynamics associated with occupancy of the selectivity filter K+ modulatory site. It is well known that Kv channel selectivity filter occupancy is modulated by a K+ binding site within the outer pore (Pardo et al., 1992; Lopez-Barneo et al., 1993; Baurkowitz and Yellen, 1995; Rasmussen et al., 1995; Kiss and Korn, 1998). Our fluorescence data suggest that K+ exit from its modulatory site within the selectivity filter is associated with local rearrangements that are sensed by the fluorophore attached within the S3–S4 linker region. However, although raised external K+ caused an appropriate hyperpolarizing shift of the Fv relation (Fig. 10 B), the fluorescence report in raised extracellular K+ appeared biphasic and was not entirely Shaker-like, so it is not entirely clear at this time whether or not K+ is specifically inhibiting the dequenching event.

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