

heat-inactivated fetal calf serum (FCS). On the night before transfection, cells were plated at 2×10^5 cells per well. Cells in each well were transfected with 2 μ g of pMTV-Luc, 2 μ g of either MR_{WT} or MR_{L810}, and 0.05 μ g of pSV2. Transfection was performed with cationic liposomes (Lipofectamine-Plus; Life Technologies) after which cells were incubated in DMEM supplemented with 10% FCS for 24 hours. Cells were washed, and then serum-free DMEM containing the test steroid was added. The cells were incubated for an additional 16 hours before assay. Luciferase and β -Gal activities were measured as described (22). Luciferase activity was normalized to β -Gal activity to correct for transfection efficiency and is expressed as a percentage of the MR_{WT} activity at 1 nM aldosterone. All results are the mean of at least nine independent transfections. For heterodimer experiments, the absolute levels of activity induced by aldosterone with MR_{WT} and MR_{L810} were indistinguishable, and multiple independent preparations of both plasmids yielded indistinguishable results in heterodimer experiments.

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15. The MR_{WT} and MR_{L810} models were generated from the PR crystal structure (13) with the Turbo Frodo molecular graphics program (http://afmb.cnrs-mrs.fr/TURBO_FRODO). Residues abutting the ligand were altered to match the MR sequence (G773A, M810S, Y941F, and M960L). In addition, the E774C substitution was incorporated to reflect the rotational freedom G774 confers on helix 3 in MR. The geometry was regularized with Turbo Frodo. Coordinates for aldosterone (23) were superimposed upon progesterone. Parameters for aldosterone were generated with HIC-Up (24). The 21-hydroxyl group was rotated to form a favorable interaction with the N770 side chain (14).

16. COS-7 cells were transfected in 100-mm plates with 10 μ g of receptor plasmid by using Lipofectamine 2000 (Life Technologies). On the day after transfection, serum-free media was substituted, and the cells were grown for an additional 24 hours. Cells were harvested in 40 mM tris-HCl (pH 7.5), 150 mM NaCl, 1 mM EDTA and lysed by freeze-thaw treatment in hypotonic buffer containing 10 mM tris-HCl (pH 7.8), 10 mM NaCl, 1 mM EDTA, 10 mM Na₂MoO₄, 5 mM dithiothreitol, antipain (5 μ g/ml), leupeptin (5 μ g/ml), chymostatin (5 μ g/ml), pepstatin A (5 μ g/ml), and 500 μ M phenylmethylsulfonyl fluoride. After centrifugation at 15,000g for 15 min, extracts were adjusted to 100 mM NaCl and 5% glycerol (binding buffer). Extracts were incubated overnight with [³H]aldosterone and competitor steroid at 0°C in a total volume of 200 μ l, and then incubated with 100 μ l of a 50% slurry of hydroxyapatite in binding buffer. Samples were centrifuged, washed twice in

binding buffer, and then resuspended in ethanol and prepared for scintillation counting. The value for 100% binding was determined by subtracting the number of counts per minute bound in the presence of 500-fold excess of unlabeled aldosterone from the counts bound in the absence of competitor. Nonspecific binding was determined with a 500-fold excess of unlabeled aldosterone. No specific binding was seen in mock-transfected cells.

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Structure of the Cytoplasmic β Subunit—T1 Assembly of Voltage-Dependent K⁺ Channels

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The structure of the cytoplasmic assembly of voltage-dependent K⁺ channels was solved by x-ray crystallography at 2.1 angstrom resolution. The assembly includes the cytoplasmic (T1) domain of the integral membrane α subunit together with the oxidoreductase β subunit in a fourfold symmetric T1 β ₄ complex. An electrophysiological assay showed that this complex is oriented with four T1 domains facing the transmembrane pore and four β subunits facing the cytoplasm. The transmembrane pore communicates with the cytoplasm through lateral, negatively charged openings above the T1 β ₄ complex. The inactivation peptides of voltage-dependent K⁺ channels reach their site of action by entering these openings.

The β subunit of voltage-dependent K⁺ channels is a tetramer of oxidoreductase proteins arranged with fourfold rotational symmetry like the integral membrane α subunits

(1). Each oxidoreductase protein contains an active site with catalytic residues and an NADPH (the reduced form of nicotinamide adenine dinucleotide phosphate) cofactor, but the specific substrate is unknown and the biological function of the β subunit remains a mystery.

Studies of K⁺ channel biosynthesis have shown that α and β subunits coassemble in the endoplasmic reticulum and remain together as a permanent complex (2, 3). The idea that a large macromolecular assembly is attached to the intracellular face of voltage-dependent K⁺ channels has important impli-

cations for channel regulation, but it also raises the question of how the transmembrane pore opens to the cytoplasm. This issue of pore access first arose when the T1 domain, an about 100-amino acid structure on the intracellular side of the first membrane-spanning segment of the α subunit, was found to form a tetrameric ring with a narrow, positively charged central pore (4). The small T1 pore diameter and positive charge are inconsistent with functional measurements showing that organic cations such as tetraethylammonium enter the transmembrane pore (5). Even a peptide segment from the channel itself (inactivation peptide) is thought to enter the pore to produce inactivation (6, 7). How can entry of these large molecules be reconciled in the setting of a narrow T1 pore? By analyzing the structure and function of the cytoplasmic interface, we resolve this apparent inconsistency and show how the T1 tetramer forms a docking platform for the β subunit without obstructing the transmembrane pathway.

It has been postulated that the intracellular T1 domain interacts with the β subunit (8, 9). We reinforced this idea by showing that removal of the T1 domain, but not the K⁺ channel's COOH-terminus, disrupts β subunit association (see below).

Coexpression of the rat β 2 subunit (10) (residues 36 to 367) and rat Kv1.1 (11) T1 domain (residues 1 to 135) in Sf9 cells yielded a stable complex. The complex was purified and crystallized, and its structure was determined by molecular replacement with the β subunit structure as a search model (12). The final model was refined to an R_{free}

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of 0.229 (40 to 2.1 Å) (12). Packing of subunits within the crystal defined an unambiguous interface through which the β subunit and T1 tetramers interact: The large flat surface of β engages four prominent loops (contact loops) extending from the NH_2 -terminal side of the T1 tetramer (Fig. 1).

Comparison of the isolated T1 and β subunit structural models with their counterparts in the complex reveals only small differences. One of these is a slight tilt of the T1 domain subunits with respect to the central four-fold axis (4). The tilt results in a separation of the subunits at the COOH-terminal side, widening the aperture, and a constriction at the NH_2 -terminal side where T1 contacts the β subunit.

To confirm this complex in vivo, we studied the interaction of α and β subunits in *Xenopus* oocytes by measuring inactivation, which refers to occlusion of the pore by an inactivation peptide (Fig. 2A) (6, 10). The inactivation peptide is effective when attached to the NH_2 -terminus of either the α or β subunits (Fig. 2B), apparently because the attachment sites are very near each other in the $\text{T1}_4\beta_4$ complex (Fig. 1C). Thus, if the α subunit does not have its own inactivation peptide, trans-inactivation caused by a β subunit indicates α - β subunit association. Following a described method (10), a Kv1.4 K^+ channel without an inactivation gate (Kv1.4-IR) was coexpressed with the $\beta 2$ subunit containing an inactivation gate ($\beta 12$ chimera) (Fig. 2) (13). The T1 domain of Kv1.4 is 85% identical to that of

Kv1.1 used in the structure determination.

When the T1 domain is removed from the Kv1.4-IR channel by means of genetic deletion (13) (residues 2 to 278), inactivation is not observed in the presence of the $\beta 12$ chimera (Fig. 2C), implying that the T1 domain is necessary for β subunit association. The occurrence of inactivation when the inactivation peptide is on the channel's own NH_2 -terminus, even in the absence of a T1 domain, demonstrates that the inactivation process itself does not require a T1 domain [Fig. 2C; see also (14)]. Thus, the T1 domain does not participate directly in inactivation but serves to hold the β subunit in place.

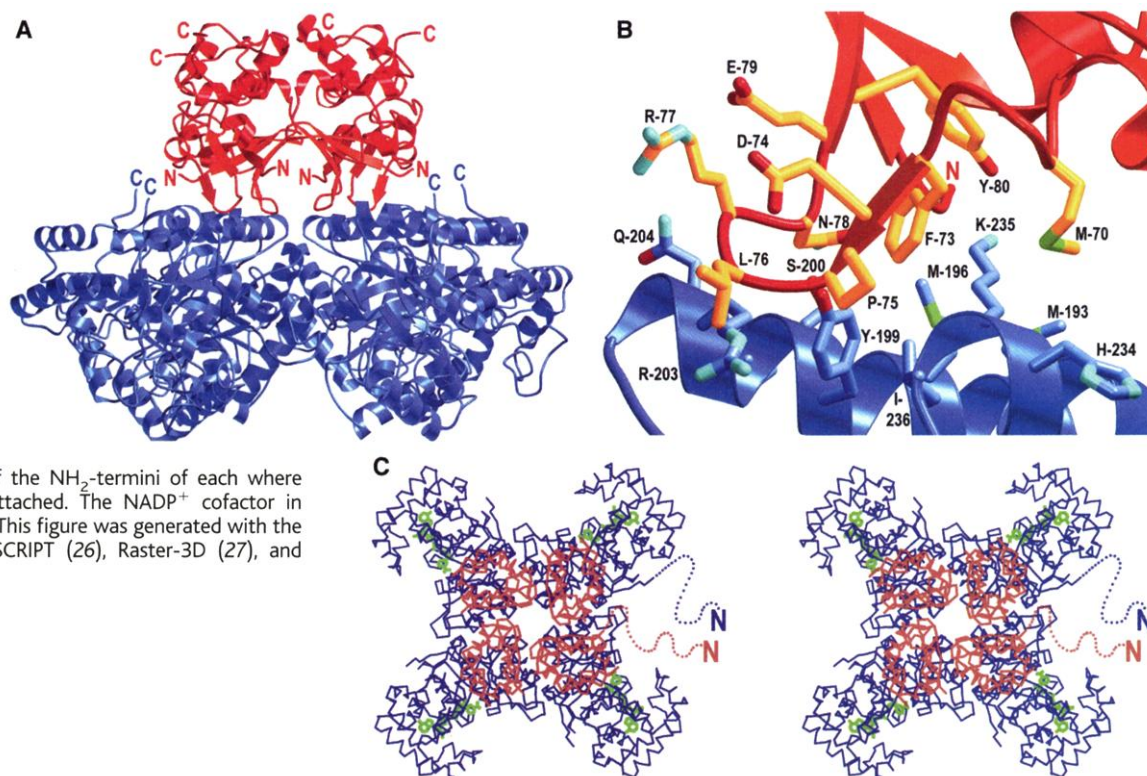
To determine which amino acids are important for $\text{T1}_4\beta_4$ complex formation in vivo, we made single-site alterations (13). On the T1 domain, only mutations involving contact loop residues abolish inactivation (Fig. 2, D and E). The effect of the point mutations is most compatible with disruption of β subunit binding to the α subunit: The same mutations do not interfere with inactivation caused by a gate on the NH_2 -terminus of the α subunit (15). On the β subunit, single-site mutations affect inactivation only when introduced onto the surface where the T1 contact loops touch the β subunit in the crystal structure (Fig. 2, D and E).

Certain mutations on the β subunit [e.g., Met¹⁹⁶ \rightarrow Trp (M196W)] reduce the extent of inactivation but do not abolish it (Fig. 2D). In such cases, the inactivating component exhibits wild-type kinetics, implying that only a fraction of channels contain a β sub-

unit because the mutant β subunit has a reduced affinity for the α subunit. The possibility of insufficient mutant β subunit expression, resulting in a mismatch between the numbers of β and α subunits, is excluded by a control experiment. Mutation M196W on the β subunit rescues the loss of inactivation caused by T1 contact loop mutation Phe²¹⁴ \rightarrow Val (F214V) (Fig. 2D). This result offers very strong support for the crystallographically defined structure in the context of a living cell. The conclusion that the T1 contact loops provide the β subunit docking surface offers insight into the specificity of α - β subunit assembly: The amino acid sequence of the T1 contact loop is highly conserved within a given family of K^+ channels but not between members of different families. Thus, $\beta 1$ and $\beta 2$ subunits associate with Kv1 but not Kv2 K^+ channels (16).

The aqueous channel down the center of the $\text{T1}_4\beta_4$ complex is too narrow (~ 4 Å and positively charged) to allow entry of tetraethylammonium, a pore-blocking cation, or the inactivation peptide (Fig. 3). If not through the center of the $\text{T1}_4\beta_4$ complex, how does the ion pathway connect to the cytoplasm? We addressed this question by again exploiting inactivation. Inactivation occurs when an inactivation peptide binds at or near the pore (Fig. 2A). Electrostatic interactions help to mediate this process: Mutations of basic (positive charged) amino acids on the inactivation peptide influence the rate and extent of inactivation (17). Therefore, we expect to find complementary acidic (neg-

Fig. 1. Structure of the $\text{T1}_4\beta_4$ complex. (A) Ribbon representation showing four contact loops that form the primary interface between the T1 and β tetramers. The T1 tetramer is red and the β tetramer is blue. (B) Molecular detail of a T1 contact loop touching the β subunit surface (24). (C) Stereo-view of a $\text{C}\alpha$ trace of the β tetramer (blue) and T1 tetramer (red) viewed along the four-fold axis and showing the relative proximity of the NH_2 -termini of each where inactivation gates are attached. The NADP^+ cofactor in each active site is green. This figure was generated with the programs O (25), MOLSCRIPT (26), Raster-3D (27), and POVray (28).



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active charged) amino acids on the channel near the inactivation peptide-binding site. The T1-S1 linker (connecting T1 to the first membrane-spanning segment) contains several acidic amino acids. When three of these residues, located six amino acids COOH-terminal to the T1 structure (Glu²⁷³, Asp²⁷⁴, and Glu²⁷⁵ in Kv1.4), are mutated to Ala (neutral) or Lys (positive), the rates of inactivation and recovery from inactivation are affected (18) (Fig. 4, A and B), and mutations to Lys (negative to positive) have the largest influence. Mutation of Val²⁴⁷ to Lys (neutral to positive) also reduced the rate of inactivation (Fig. 4C). Val²⁴⁷ is located on the surface of the T1 domain close to where the

T1-S1 linker emerges (Fig. 5).

Mutations were also made in the inactivation peptide to reduce its positive charge. These mutations affected inactivation, but more importantly their influence depended on the charge at positions 273 to 275 in the T1-S1 linker. Thus, mutations in the inactivation peptide are coupled in mutant cycle analysis to mutations in the T1-S1 linker near the T1 domain (Fig. 4C) (19). In order for such coupling to occur, the peptide must be near the T1-S1 linker when it inactivates the pore. Because the peptide cannot fit through the center of the T1₄β₄ complex, it must reach the pore by entering lateral openings above the T1 tetramer (Fig. 5).

The structure of the T1 domain tetramer seemed to be inconsistent with several decades of K⁺ channel pharmacology: The central hole is too narrow for many of the classical organic blocking agents (4). When the T1 domain was deleted from the α subunit, the single channel conductance did not change substantially, as if the T1 tetramer does not form an extension of the pore (20). Thus, it was proposed that either the T1 domain is not a tetramer in the context of the integral membrane channel or that ions move between the cytoplasm and pore through an alternative pathway.

The data presented here establish that the T1 domain is indeed a tetramer in the K⁺

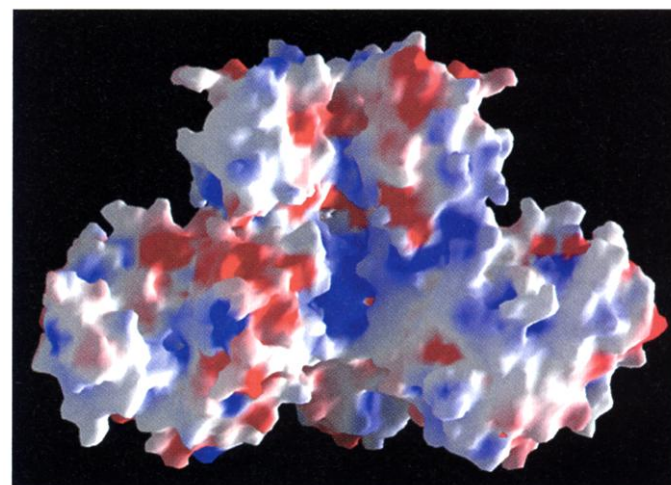
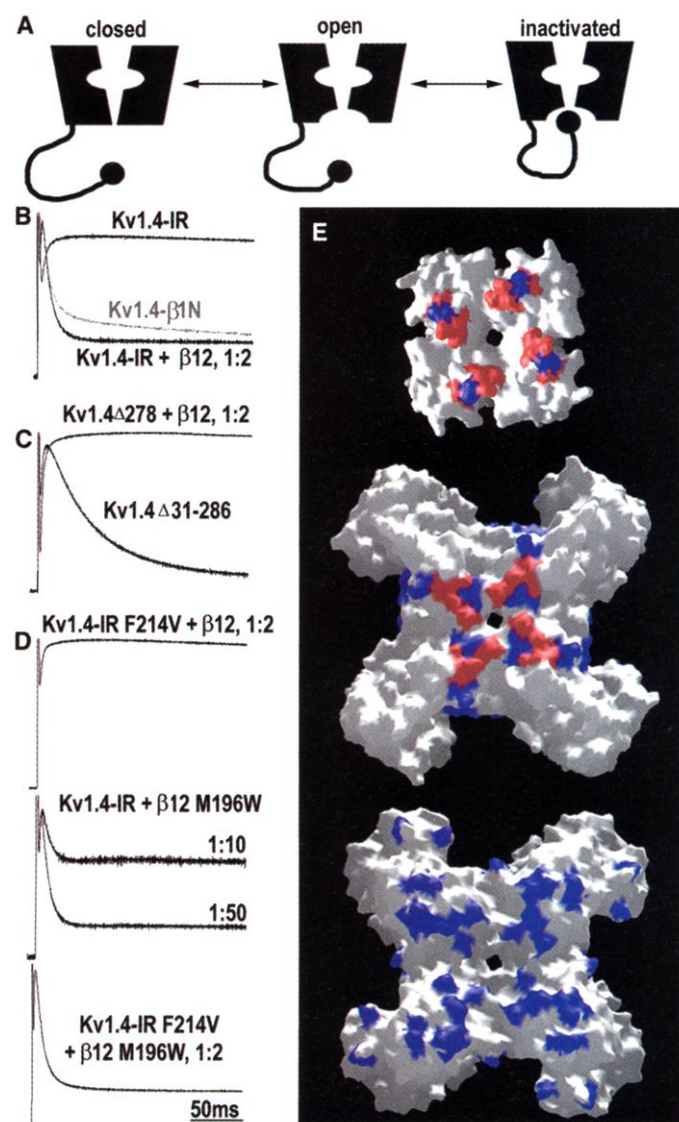


Fig. 2 (left). Functional analysis of the T1₄β₄ interface. (A) Illustration of N-type K⁺ channel inactivation. The black circle represents the inactivation peptide. (B) Potassium currents recorded from *Xenopus* oocytes under voltage clamp containing channels without an inactivation peptide (Kv1.4-IR) or with an inactivation peptide attached to the NH₂-terminus of the α subunit (Kv1.4-β1N) or β subunit (Kv1.4-IR + β12). The α and β subunits were coexpressed at an RNA ratio of 1:2 (v:v) (13). (C) The Kv1.4Δ278 channel lacks amino acids 2 to 278, which make up Kv1.4 channel's own inactivation peptide and T1 domain. The β12 chimera fails to cause inactivation (Kv1.4Δ278 + β12). The Kv1.4Δ31-286 channel (amino acids 31 to 286 deleted) lacks a T1 domain but contains its own inactivation peptide (Kv1.4Δ31-286), which differs from that of β12, accounting for the slower inactivation rate. (D) (Top) A point mutation on the contact loop of the T1 domain (Kv1.4-IR F214V) prevents β12 chimera-induced inactivation (residue Phe⁷³ in Kv1.1) (Middle) Currents carried by Kv1.4-IR channels expressed with a mutant β subunit (β12 M196W) at two RNA volume ratios, 1:10 and 1:50, undergo incomplete inactivation. (Bottom) Coexpression of Kv1.4-IR F214V mutant α subunit and β12 M196W mutant β subunit yields inactivation properties similar to the combination of wild-type α and β subunits. In all recordings, oocytes were held at -80 mV and stepped to +60 mV for 200 ms. (E) GRASP view of T1 and β2 subunits showing the effect of selected point mutations on channel inactivation (red, alteration; blue, no alteration). (Top) NH₂-terminal surface of the T1 domain. Red residues are Phe⁷³, Leu⁷⁶, and Arg⁷⁸, and the blue residue is Pro⁷⁵. (Middle) Flat surface of the Kvβ2 subunit. Red residues are Met¹⁹⁶, Tyr¹⁹⁹, Ser²⁰⁰, and Gln²⁰⁴, and blue residues are Arg²⁰³, Lys²⁶⁸, Lys¹²⁹, and Met¹⁹³. (Bottom) Concave

surface of the β subunit. Blue residues are Asp⁶⁶, Asp⁶⁷, Tyr⁹⁰, Lys¹⁰⁴, Lys¹⁰⁵, Lys¹⁰⁶, Lys¹²⁴, Glu¹⁴⁵, Gln¹⁴⁸, Glu¹⁵⁰, Asn¹⁵⁸, Arg¹⁵⁹, Arg¹⁸⁹, Pro²⁶⁰, and Pro²⁶¹. This figure was made with GRASP (29). **Fig. 3 (right).** The central pore of the T1₄β₄ complex. A molecular surface calculated over three of the four T1-β subunits reveals a central pore with unusual internal electrostatic features. The pore is coincident with the molecular four-fold axis and extends the length of the complex. At its apex, it is sufficiently wide to permit the entry of ions. The inner surface of this cavity is lined with charged amino acid side chains that are conserved in the Kv1 family. Calculation of the electrostatic potential was carried out at an ionic strength equivalent to 0.15 M KCl. Dielectric constants of 2.0 for the protein interior and 80.0 for solvent were used. Regions of the molecular surface exhibiting intense negative charge are colored red, and electropositive regions are colored blue. This figure was generated with the program GRASP (29).

Fig. 4. Evidence for pore openings above the T1 domain. (A) (Top) β 12 chimera-induced inactivation of wild-type Kv1.4-IR α subunits (WT) and α subunits with Ala (A) or Lys (K) substituted at positions 273 to 275. (Bottom) Currents from wild-type α subunits (WT) or α subunits with Ala substituted at 273 to 275 (A) coexpressed with the β 12 chimera containing a wild-type inactivation gate (+ β 12) or K13N, R15G mutant inactivation gate (+1315NG). The experimental conditions were identical to those described in Fig. 2D. (B) (Top) Rate of recovery from inactivation measured in paired pulses from a holding voltage of -80 mV and stepping the membrane to $+60$ mV. The initial conditioning pulse was 400 ms. The dashed curve corresponds to an exponential function ($\tau = 11$ s) fitted to the envelope defined by the peak current of the test pulse. (Bottom) Fraction of recovery as a function of time between the conditioning pulse and the test pulse. ■, Kv1.4-IR α subunit paired with the β 12 chimera; ●, α subunit with Ala substituted at 273 to 275 coexpressed with β 12 chimera; ○, α subunit with Lys substituted at 273 to 275 coexpressed with β 12 chimera. The solid lines are single exponential fits. Error bars are the standard error of the mean (SEM) from \geq five independent experiments. (C) Double-mutant cycle analysis of inactivation. The time constants of inactivation (τ_{inac}) and recovery (τ_{rec}) and the ratio between these two time constants are tabulated (\pm SEM, \geq five independent experiments) (18). The Ω value was calculated as described (19).

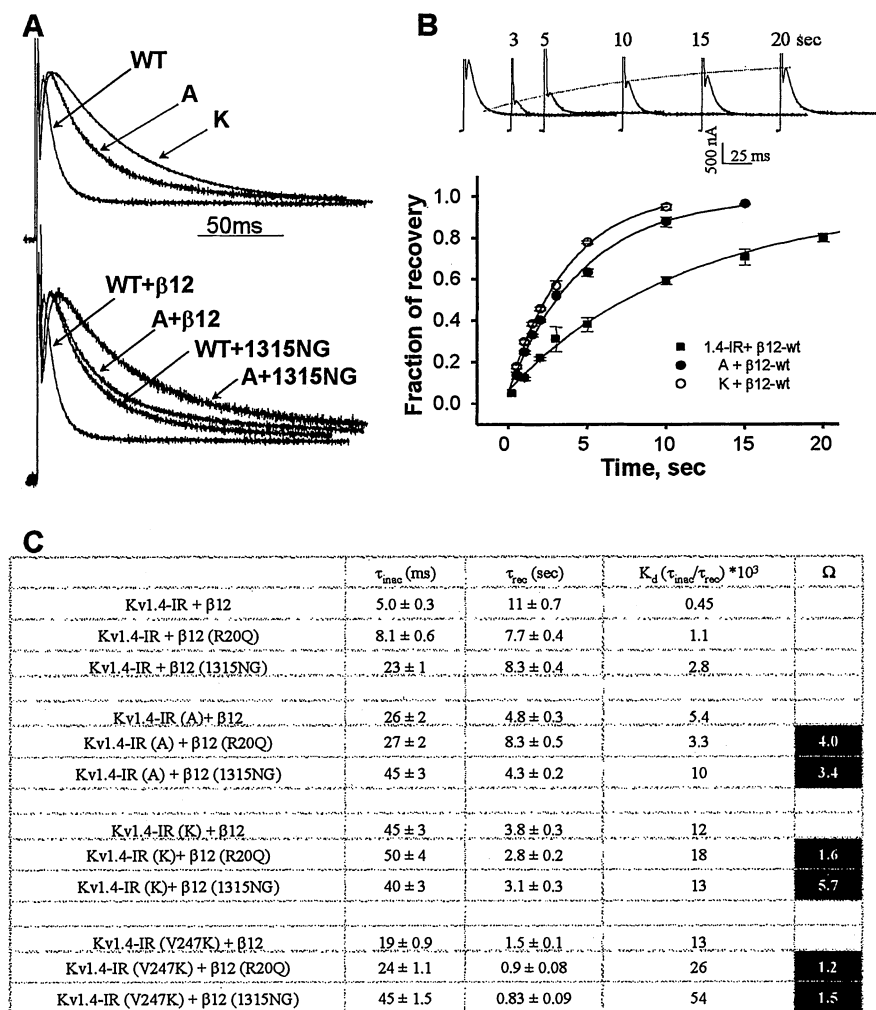
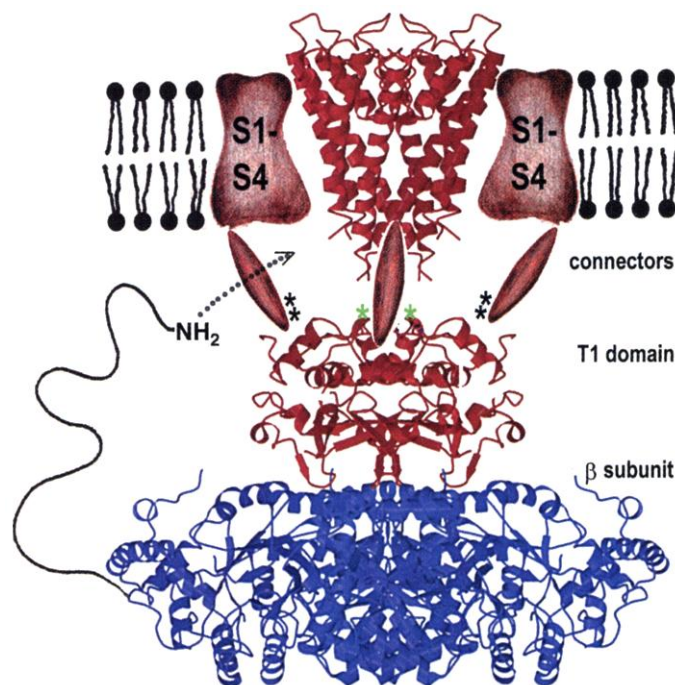


Fig. 5. Composite model of a voltage-dependent K^+ channel. The α subunit is shown in red, and the β subunit is in blue. The model of the pore region is based on the KcsA K^+ channel (30). The structures of the voltage-sensing region and connectors are unknown (depicted schematically). An NH_2 -terminal inactivation peptide is shown entering a lateral opening to gain access to the pore. Proposed locations of amino acids 273 to 275 on the T1-S1 linker are shown (black asterisks) with the location of Val²⁴⁷ on the T1 domain surface (green asterisks).



channel and that the T1 domain tetramer forms the docking platform for the β subunit. The contact loops extending from the NH_2 -terminal face of the T1 domain are obvious structures for interfacing with intracellular proteins—in this case the β subunit. We suggest that the major role of the T1 domain is to form a docking site for proteins whose activities are coupled to K^+ channel function.

The apparent paradox of a too narrow pore in the T1 domain has a simple explanation. Our data suggest the presence of lateral openings between the T1 domain and the integral membrane part of the α subunit (Fig. 5). This explanation accounts for the interaction between the inactivation peptide and amino acids on the top (COOH-terminal face) of T1 and in the T1-S1 linker close to T1. Because the peptide must access the pore through lateral openings, K^+ ions must also pass through the openings. Lateral openings in the cytoplasm, with negative charged amino acids, are also a structural feature of nicotinic acetylcholine receptor ion channels (21).

A single inactivation gate binds to its receptor near the intracellular pore entryway in an exclusive fashion when inactivation occurs (22,

23). Our findings are entirely consistent with this conclusion. Although we expect there to be four openings above the T1 domain, there is presumably only one site near the pore entryway where a single peptide can induce inactivation. We propose that the inactivation peptide is poorly structured and snakes its way into a lateral opening to reach the central pore and this is why mutations in the T1-S1 linker influence the process. In some cases, four peptides may even be bound above T1, poised and ready, before inactivation.

The central question as to why K^+ channels contain an oxido-reductase enzyme subunit remains unanswered. In this report, we have shown how the β subunit attaches to a voltage-dependent K^+ channel through its interaction with the T1 domain. We previously suggested that interactions between α and β subunits may allow cellular redox regulation of the channel or channel regulation of enzyme activity (1). The structure of the $T1_4\beta_4$ complex takes us one step closer to understanding how such energetic coupling might occur.

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12. DNA encoding the NH_2 -terminal cytoplasmic T1 domain (residues 1 to 135) of the rat Kv1.1 α subunit was subcloned into a pAcG2T (Pharming) baculovirus transfer vector as a glutathione S-transferase (GST) fusion construct with Bam HI and Eco RI restriction sites. DNA encoding residues 36 to 367 of rat Kv β 2 was subcloned into pFastBac1 (Gibco) between Bam HI and Eco RI restriction sites. For coexpression, cells were infected with two viruses. At 72 hours, cells were harvested by centrifugation and resuspended in a cold buffer (I) comprising 20 mM tris-Cl (pH 8.0), 150 mM KCl, 5 mM $MgCl_2$, and 5 mM dithiothreitol. Phenylmethylsulfonyl fluoride was added to 1 mM, and the cells were promptly lysed with a French pressure cell. After adding deoxyribonuclease, the lysate was incubated for 20 min on ice and then centrifuged to pellet cellular debris. The filtered lysate was incubated with glutathione sepharose 4B beads (Pharmacia) for 2 hours at 4°C. Proteolytic (thrombin) removal of the GST tag was preceded by extensive washing of the matrix-bound material with a buffer (II) containing 20 mM tris-Cl (pH 8.0), 150 mM KCl, and 2 mM $CaCl_2$. The cleaved protein was eluted, concentrated, and loaded onto a size-exclusion (Superdex-200 10/30; Pharmacia) column equilibrated in buffer (I). Peak fractions corresponding to a heterooctamer ($T1_4\beta_4$) were pooled and cleaved with trypsin, yielding a $T1(27-129)_4\beta_4$ octamer. Size exclusion was used as a final purification step. Column fractions corresponding to the heterooctameric complex were collected, pooled, and concentrated to 15 mg/ml. Tetragonal crystals containing a single $T1\beta$ heterodimer ($P42_2$; $a = 100.7$ Å, $c = 110.9$ Å) grew in hanging drops; 1 μ l of the protein solution was mixed with 1 μ l of a reservoir solution containing 50 mM glycine (pH 8.6) and 6% PEG400 and allowed to equilibrate by vapor diffusion. Crystals were cryoprotected by transfer into a similar solution containing 25% glycerol, mounted in nylon loops, and frozen in propane. Diffraction data to 2.1 Å were collected from single plates at beamlines A1 (CHESS) and X25 (Brookhaven NSLS) with charge-coupled device detectors. Images were processed with DENZO and intensities were merged with SCALEPACK [Z. Otwinowski, in *Proceedings of the CCP4 Study Weekend* (SERC Daresbury Laboratory, Warrington, UK, 1993), pp. 56–62; Z. Otwinowski and W. Minor, *Methods Enzymol.* **276**, 307 (1997)]. A set of 400,177 measured reflections (33,907 unique) in the resolution range 40.0 to 2.1 Å gave a 98.8% complete data set (92.9% in the outer shell; 2.15 to 2.10 Å). The overall agreement factor R_{merge} (on intensity) is 8.0% (30.0% in the outer shell). The average intensity/standard deviation is 17.8 (4.3 in the outer shell). The $T1_4\beta_4$ structure was solved by molecular replacement with AMORE [J. Navaza, *Acta Crystallogr. Sect. A* **50**, 157 (1994)]. A refined model of the conserved core of rat Kv β 2 (119; residues 36 to 360) was used to initiate rotation and translation searches in the resolution range 8.0 to 4.0 Å. A peak 8.8 σ above the mean was the highest in the rotation function. Translation searches yielded a peak at 18.0 σ . Cycles of rigid body refinement of the heterodimer with data between 8.0 and 4.0 Å reduced the R factor to 0.393 and gave a correlation coefficient of 57.7. Before refinement, all temperature factors were fixed at 30 Å², and several residues in mobile regions together with several side chains were deleted from the model. The β subunit T1 complex model was then refined by minimization and simulated annealing procedures with the maximum likelihood target implemented in CNS [A. T. Brunger et al., *Crystallogr. D. Biol. Crystallogr.* **54** (part 5), 905 (1998)]. $3F_o - 2F_c$ (where F_o and F_c are the observed and calculated structure factors, respectively) maps were calculated on the basis of an average of 10 refined structures and were of sufficiently high quality to enable almost complete tracing of the T1 domain. A number of cycles of building and refinement were required to place the remainder of the structure. All data falling between resolution limits of 40.0 and 2.1 Å, with the exception of a random 5% used for calculation of the R_{free} , were used in the refinement. Bulk solvent and anisotropic temperature factor corrections (final anisotropic tensor elements: B11, 0.26 Å²; B22, 0.26 Å²; and B33, -0.52 Å²) were applied to the reflection data. The crystallographic R_{free} was monitored to reduce model bias. Individual temperature factors were refined for all nonhydrogen atoms. The refined model has good geometry with one Ramachandran outlier near NADP and contains 418 residues and 1 nicotinamide adenine dinucleotide phosphate (NADP⁺) and 267 water molecules and has an R_{free} of 0.229 and an R_w of 0.211. Six residues at the COOH-terminus of the β subunit, eight at the NH_2 -terminus of T1, and three at the COOH-terminus of T1 were omitted from the model because of weak electron density in the region. Side chains were not modeled beyond the β carbon for the following amino acids: Lys¹²⁴, Lys²⁶⁸, and Ser³⁶¹.
13. The Kv1.4-IR (residues 2 to 110 deleted), Kv1.4 Δ 278 (residues 2 to 278 deleted), and Kv1.4 Δ 31-286 (residues 31 to 286 deleted) were generated by the overlapping polymerase chain reaction method. The β 12 chimera and the Kv1.4- β 1N were made by splicing DNA encoding residues 1 to 70 of β 1 onto DNA encoding residues 36 to 366 of β 2 or DNA encoding residues 137 to 655 of Kv1.4, respectively. All constructs were cloned into pBluescript KS⁺. Point mutations were made by the QuickChange method (Stratagene) and confirmed by sequencing the entire insertion. RNA was prepared by T7 polymerase transcription and injected into *Xenopus laevis* oocytes [Z. Lu and R. MacKinnon, *J. Gen. Physiol.* **104**, 477 (1994)]. RNA ratios refer to volumes mixed before injection. K^+ currents were recorded under two-electrode voltage clamp (OC725B; Warner Instrument, Hamden, CT). Electrodes were drawn from borosilicate glass capillaries (World Precision Instrument, Sarasota, FL) to a resistance of ~0.5 megaohms (3 M KCl). Bath solution contained 96 mM NaCl, 2 mM KCl, 0.3 mM $CaCl_2$, 1 mM $MgCl_2$, and 5 mM Hepes (pH 7.6) with NaOH. Analog data from the amplifier were filtered (2 kHz, -3 dB) with an eight-pole Bessel filter (Frequency Devices, Haverhill, MA), digitized at 10 kHz, and transferred to a computer hard disk with an Axon 1200 digitization board and supporting software (Clampex version 8; Axon Instrument, Foster City, CA).
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18. Inactivation time constants were obtained by fitting the current decay with exponential functions with Clampfit (Axon) software. Expression of Kv1.4-IR with β 12 chimera yielded two exponentials: a predominant (>90%) fast component (time $\tau < 60$ ms) and a minor (<10%) component ($\tau \sim 500$ ms). The slower time constant was relatively invariable and coincided with the COOH-type inactivation of Kv1.4-IR [S. H. Heinemann, J. Rettig, H. R. Graack, O. Pongs, *J. Physiol.* **493**, 625 (1996)]. In double-mutant cycle analysis, only the predominant component was used. Inactivation recovery was studied with paired pulses (22). The peak of the second pulse in the pair was graphed as a function of time after the first pulse, defining an envelope of recovery that was fit to an exponential function with Origin (Microcol Software, Northampton, MA). Recovery from inactivation for Kv1.4-IR paired with the β 12 chimera was well described by single exponential function.
19. The double-mutant cycle parameter $\Omega = \frac{K_d^{wt,wt} \times K_d^{mut,mut}}{K_d^{wt,mut} \times K_d^{wt,wt}}$ was used to quantify the degree of coupling between two mutations [P. Hidalgo and R. MacKinnon, *Science* **268**, 307 (1995)]. Here, K_d refers to a dimensionless parameter defined by the ratio of inactivation and recovery time constants, $\tau_{\text{inac}}/\tau_{\text{rec}}$. Because $\tau_{\text{rec}} \gg \tau_{\text{inac}}$, K_d is very close to the equilibrium ratio of open to inactivated channels. A value of Ω greater than unity indicates that the effects of two mutations are coupled.
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24. Single-letter abbreviations for the amino acid residues are as follows: A, Ala; C, Cys; D, Asp; E, Glu; F, Phe; G, Gly; H, His; I, Ile; K, Lys; L, Leu; M, Met; N, Asn; P, Pro; Q, Gln; R, Arg; S, Ser; T, Thr; V, Val; W, Trp; and Y, Tyr.
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