Crystal Structure of a Voltage-gated K⁺ Channel Pore Module in a Closed State in Lipid Membranes

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Voltage-gated K⁺ channels underlie the electrical excitability of cells. Each subunit of the functional tetramer consists of the tandem fusion of two modules, an N-terminal voltage-sensor and a C-terminal pore. To investigate how sensor coupling to the pore generates voltage-dependent channel opening, we solved the crystal structure and characterized the function of a voltage-gated K⁺ channel pore in a lipid membrane. The structure of a functional channel in a membrane environment at 3.1 Å resolution establishes an unprecedented connection between channel structure and function. The structure is unique in delineating an ion-occupied ready to conduct selectivity filter, a confined aqueous cavity, and a closed activation gate, embodying a dynamic entity trapped in an unstable closed state.

Ion channels allow the selective and regulated diffusion of ions across membranes. Voltage-gated K⁺ channels (Kv) are pimemeric proteins organized around a central pore. Each subunit consists of two tandemly arranged structural modules: an N-terminal voltage sensor module encompassing four transmembrane segments (TMs) S1–S4 and a C-terminal pore (PM) consisting of TMs S5 and S6 (2–6). The two modules are connected through a cytoplasmic linker (S4-S5 linker) between TMs S4 and S5.

Molecular dissection of KvLm, a Kv from Listeria monocytogenes, led to the isolation of a sensorless PM that exhibits the ionic selectivity, permeation, and blocker sensitivity properties characteristic of the intact KvLm, namely a functional PM (7). Here, we present the structure of such PM obtained from diffraction quality crystals using the lipidic cubic phase approach (8, 9).

EXPERIMENTAL PROCEDURES

Protein Expression and Purification—A single amino acid mutation (A98C) was introduced in the KvLm PM (7) using the QuikChange I site-directed mutagenesis kit (Agilent), according to the manufacturer’s instructions. Mutant channel protein was expressed as previously described (7). Following centrifugation, the cells were resuspended in 50 mM Hepes, pH 7.5, 50 mM KCl, 5 mM MnCl₂, 0.1 mM CaCl₂, 10% (v/v) glycerol, 0.02% (w/v) NaN₃ (denoted as low salt buffer) supplemented with bovine serum albumin, lysozyme (Sigma-Aldrich), DNase, and protease inhibitor tablets (Roche Applied Science) and lysed with a microfluidizer. The membranes were pelleted by centrifugation at 125,000 × g, resuspended in the low salt buffer, and incubated for 1 h at 4 °C. The procedure was repeated with a high salt membrane wash buffer (same as the low salt buffer but with 0.5 M KCl). Following ultracentrifugation, the cells were resuspended in solubilization buffer (50 mM Hepes, pH 7.5, 50 mM KCl, 5 mM imidazole) prior to the addition of n-decyl-β-D-maltopyranoside (DM) (Anatrace) to 20 mM. The membranes were solubilized for 3–3.5 h at 4 °C under stirring, followed by the addition of KCl to a final concentration of 400 mM and ultracentrifugation at 125,000 × g for 1 h. The supernatant was passed through a cobalt (Co²⁺) IMAC resin (Clontech) three times before rigorous washing in three steps: first, with 20 column volumes of 50 mM Hepes, pH 7.5, 400 mM KCl, 20 mM imidazole, 10% glycerol, 15 mM DM followed by 20 column volumes with 50 mM Hepes, pH 7.5, 300 mM KCl, 20 mM imidazole, 5% (v/v) glycerol, 10 mM DM, and a final wash with 20 column volumes of 50 mM Hepes, pH 7.5, 200 mM KCl, 20 mM imidazole, 1% (v/v) glycerol, 5 mM DM. Purified KvLm was eluted in 50 mM Hepes, pH 7.5, 200 mM KCl, 500 mM imidazole, 1% glycerol, 5 mM DM. Pooled fractions were desalted in PD-10...
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desalting column (GE Healthcare) and eluted in crystallization buffer (50 mM Hepes, pH 7.5, 200 mM KCl, 3 mM DM, 0.1 mM TCEP). Protein digestion with 1-(tosylamido-2-phenyl)ethyl chloromethyl ketone modified trypsin (New England Biolabs) was carried out overnight at ~20 °C by incubation with trypsin at a trypsin to PM mass ratio of 50:1. Analysis by nano-LC/MS/MS (The Scripps Research Institute Center for Mass Spectrometry) showed that the enzymatic digestion of KvLm-PM removed the C-terminal 22 residues, including the His$_{6}$ tag, to yield a truncated PM construct at residue Arg$^{137}$ with a molecular mass of 15.6 kDa. The digested PM was isolated by passing the enzymatic reaction first through 1 ml of nickel-nitrilotriacetic acid IMAC resin (Qiagen) and second through 100 μl of benzamidine-Sepharose resin (GE Healthcare) and collecting the flow through. The sample was then concentrated to 40 ~ 45 mg ml$^{-1}$ using a 100-kDa cutoff concentrator (Sartorius). Analytical Size exclusion chromatography and SDS-PAGE were used to monitor sample homogeneity and purity, respectively.

Crystallization and Data Collection—The protocol used for lipidic cubic phase crystallization has been described in detail previously (8, 9). Diffraction quality crystals were obtained in 26 ~ 30% (v/v) PEG-MME-550, 0.5 ~ 0.6 M ammonium sulfate, and 50 mM ADA, pH 7.0 (ammonium condition), and in 25% (v/v) PEG-MME-550, 0.35 M sodium nitrate, 50 mM ADA, pH 6.25, 50 mM sodium malonate, pH 7.0 (sodium condition). The ions found in the selectivity filter of both structures could be $NH_4^+$ (Fig. 1, B and C) or Na$^+$ (Fig. 1, D and E), the majority cations in each crystallization solution. However, this is unlikely, given 1) that 27 ms $K^+$ is the only constant in the two conditions and 2) the observed significant increase in the $R/\text{F}_{\text{free}}$ upon $NH_4^+$ ion substitution. For simplicity, we therefore surmise that the ions in the filter are $K^+$.

Structure Analysis—All structural analysis was performed using PyMOL (Schrodinger) and YASARA (YASARA Biosciences) on the highest resolution structure (3.1 Å) obtained in ammonium sulfate, unless otherwise indicated. A $C_α$ alignment of the two KvLm PM structures shows that they deviate by ~0.61 Å. A $C_α$ alignment of the PMs of KvLm and of the Kv1.2-Kv2.1 chimera (Protein Data Bank code 2RBR (6)) has an root mean square deviation of 1.6 Å. To calculate the electrostatic surface potential in a lipid membrane, the preoriented protein coordinates for KvLm PM were obtained from the PPM web server (21) and provided as input to the pbeq solver web server (22). The cavity profiles were obtained using the MOLE server (23).

Reconstitution in Droplet Interface Lipid Bilayers, Single-channel Recordings, and Analysis—Droplet interface lipid bilayers (24) were formed between two lipid monolayer-encased aqueous nanoliter droplets submerged in hexadecane containing 0.5 mg ml$^{-1}$ monoolein (1-oleoyl-rac-glycerol) (Avanti Polar Lipids). Liposomes were composed of 0.5 mm 1,2-diphytanoyl-sn-glycero-3-phosphocholine (Avanti Polar Lipids). Purified KvLm PM was diluted into the liposome suspension to a final concentration of 2 ~ 5 mg ml$^{-1}$. The electrode carrying the droplet with KvLm PM and desired buffer-lipid mix (0.5 ml or 27 mM KC1, 10 mM Hepes, pH 7.4) was connected to the grounded end of the amplifier head stage (Axopatch 200B). The second electrode, in a droplet containing exclusively the buffer-lipid mix, was connected to the working end of the head stage. All of the reconstitution experiments were done at room temperature. The data were acquired at 10 kHz and fil-
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Table 1
Data collection and refinement statistics

| Structure                  | Ammonium condition | Sodium condition |
|----------------------------|--------------------|-----------------|
|                            | KvLm               |                 |
| Data collection            |                    |                 |
| Space group                | P42,2              | P2,2            |
| Cell dimensions            | a = b = 53.37 Å, c = 108.16 Å | a = 56.10 Å, b = 80.67 Å, c = 55.2 Å |
| Number of reflections measured | 21,139            | 22,028          |
| Number of unique reflections | 2.994              | 3.799           |
| Resolution (Å)             | 50–3.10 (3.21–3.10) | 50–3.35 (3.47–3.35) |
| Rmerge                     | 0.132 (0.592)      | 0.116 (0.660)   |
| Mean I/σ(I)                | 18.6 (2.45)        | 15.8 (1.10)     |
| Completeness (%)           | 94.4 (74.5)        | 95.2 (86.0)     |
| Redundancy                 | 7.1 (5.5)          | 5.8 (4.8)       |

| Refinement                 |                    |                 |
| Resolution (Å)             | 16.4–3.10          | 32.6–3.35       |
| Number of reflections (test set) | 2.674 (369)     | 3.736 (534)     |
| Rmerge/Rmerge (%)          | 28.1/30.5          | 31/34.1         |
| Number of atoms            | Protein 673        | 650 (A chain), 653 (B chain) |
|                            | K⁺ ion 3           | 4               |
|                            | Lipids 57          |                 |
|                                | Average B value (Å²) | 115          |
|                                | Protein 154        | 191 (A chain), 194 (B chain) |
|                                | K⁺ ion 167         |                 |
|                                | Lipids 103         |                 |
|                                | Root mean square deviations | Bond lengths (Å) 0.010 |
|                                |                     | Bond angles (°) 1.01 |
|                                | Ramachandran plot statistics (%) a | Ramachandran favored regions 98 |
|                                |                     | Additionally allowed regions 2 |
|                                |                     | Ramachandran outliers 0 |
|                                |                     | RAMACHANDRAN outliers 1 |

* The Ramachandran plot was calculated by using Molprobity (version 3.17).

Building on the discovery that a sensorless PM dissected from the bacterial KvLm (Fig. 1A) recapitulates the permeation properties of the full-length KvLm (7), we pursued its structural characterization. A crystal structure solved at 3.1 Å (Fig. 1, B and C) in a monooolein (1-oleoyl-rac-glycerol) membrane shows that the KvLm PM shares the funnel architecture of other K⁺ channels (5, 6, 20, 26–28). Each subunit contributes two transmembrane helices (outer helix or TM S5 (Gly15–Ile7) inner helix or TM S6 (Pro-68 to Thr-93) tilted with respect to the lipid bilayer normal, a pore helix (Tyr44–Ala55) with the C-terminus directed to the pore lumen, and the selectivity filter (Thr56–Asp62) formed at the center of the tetrameric channel (Figs. 1, A and B). The selectivity filter is occupied by three equidistant (3.1 Å) K⁺ at positions (S2–S4) (29) (“Experimental Procedures”). Despite low ion occupancy, the ion binding sites in the filter are mostly intact: the backbone carboxyls of residues Thr57, Val58, Gly59, and Tyr60 and the hydroxyl of Thr57 are facing the permeation path, positioning the filter in a ready to conduct conformation (Fig. 1, B and C). In agreement, a PM structure determined at 3.4 Å in a different condition, with two subunits in the asymmetric unit, shows a fully occupied selectivity filter (Fig. 1, D and E). A narrow central cavity appears at the center of the ion channel delimited on the extracellular side by the filter and on the intracellular boundary by the crossing of the inner helices, the activation gate (30, 31).

Mapping the electrostatic potential on the solvent-accessible surface of the activation gate at the channel bundle crossing reveals that the gate is closed and positively charged (Fig. 2B). The large positive potential at the intracellular entrance is generated by residues RCKK that cap the inner helix and presumably act as an electrostatic barrier to ion access into the channel vestibule. Residues Ile85, Thr89, and Thr93 face the pore lumen resulting in a neutral electrostatic surface. Near the entryway to the filter is a negatively charged surface (Fig. 2C), most likely arising from the focusing of the pore helices dipoles.

The solvent-accessible ion-conducting pathway displays a narrow infundibulum with several constrictions (Fig. 2D). At the selectivity filter, four constrictions are created by rings of the carbonyl oxygens of the residues forming the filter (Thr57, Val58, Gly59, and Tyr60). The central cavity reveals three prominent constrictions generated by rings of Ile, Thr, and Thr side chains at positions 85, 89, and 93, respectively. The constric-
tions range from 1.9 Å diameter at the narrowest (Thr<sup>93</sup>, C<sub>y</sub>–C<sub>y</sub>) to 3.3 Å at its widest (Ile<sup>85</sup>, C<sub>y</sub>–C<sub>y</sub>). The funnel-like assembly (Fig. 2, D and E) measures 38 Å in length, with a wide entry way into the pore at the cytoplasmic side (8.8 Å at Arg<sup>97</sup>), a narrow channel vestibule (4.2 Å at Ile<sup>85</sup> C<sub>y</sub>–C<sub>y</sub>), and a wide exit (7.6 Å at Gly<sup>61</sup> O). A comparative analysis of the cavity profile highlights that the KvLm PM diverges from the stand-alone 2-TM channels KcsA and the inward rectifier Kir2.2, in that the central water accessible cavity is narrower (4.2 Å versus 9.1 Å in KcsA, and 8.9 Å in the inward rectifier) (Fig. 2E). Surprisingly, a similarly tight central cavity occurs in the structure of the cyclic nucleotide gated channel (MlotiK1), which, akin to KvLm, is assembled from four 6-TM subunits. The constricted central cavity in KvLm and MlotiK1 may account for the absence of K<sup>+</sup> within the cavity of the closed structure: the cavity is too narrow to harbor a K<sup>+</sup> ion octa-coordinated to water as observed in KcsA (Protein Data Bank code 1K4C (20)).

![Figure 1: Structure of the KvLm PM](image)

A sequence alignment of KvLm-PM with KcsA (Protein Data Bank code 1K4C) (20) and Kv1.2 PM (Protein Data Bank code 2R9R) (6) highlighting conserved residues (black). The alignment is annotated to display secondary structure and lipid-binding sites (cyan). The mutation (A98C) is highlighted in magenta. Residue numbering is according to Protein Data Bank; for KvLm it is indicated above the sequence. B and D, stereo profiles of two diametrically facing subunits displaying the electron density map (2F<sub>o</sub>−F<sub>c</sub>) contoured at 1 σ used to fit activation gate closed for crystals obtained in 27 mM K<sup>+</sup> and 0.4 M ammonium sulfate (B) or 0.3 M sodium nitrate (D). Yellow spheres depict K<sup>+</sup> coordinated by the residues 57TVG. C and E, view of the selectivity filter showing the experimental electron density map (2F<sub>o</sub>−F<sub>c</sub>) contoured at 1 σ used to fit the position of the bound ions and the filter in a conductive conformation with the carbonyl oxygens for residues Thr<sup>57</sup>, Val<sup>58</sup>, Gly<sup>59</sup>, and Tyr<sup>60</sup> and the hydroxyl of residue Thr<sup>57</sup> facing the ions in two different crystal conditions (“Experimental Procedures”).

**FIGURE 1. Structure of the KvLm PM.** A. sequence alignment of KvLm-PM with KcsA (Protein Data Bank code 1K4C) (20) and Kv1.2 PM (Protein Data Bank code 2R9R) (6) highlighting conserved residues (black). The alignment is annotated to display secondary structure and lipid-binding sites (cyan). The mutation (A98C) is highlighted in magenta. Residue numbering is according to Protein Data Bank; for KvLm it is indicated above the sequence. B and D, stereo profiles of two diametrically facing subunits displaying the electron density map (2F<sub>o</sub>−F<sub>c</sub>) contoured at 1 σ used to fit activation gate closed for crystals obtained in 27 mM K<sup>+</sup> and 0.4 M ammonium sulfate (B) or 0.3 M sodium nitrate (D). Yellow spheres depict K<sup>+</sup> coordinated by the residues 57TVG. C and E, view of the selectivity filter showing the experimental electron density map (2F<sub>o</sub>−F<sub>c</sub>) contoured at 1 σ used to fit the position of the bound ions and the filter in a conductive conformation with the carbonyl oxygens for residues Thr<sup>57</sup>, Val<sup>58</sup>, Gly<sup>59</sup>, and Tyr<sup>60</sup> and the hydroxyl of residue Thr<sup>57</sup> facing the ions in two different crystal conditions (“Experimental Procedures”).
Thus, the closed structure of KvLm is unique in that it excludes ions from the cavity and couples a wide electrostatic gate at the permeation entrance with a sterically occluded access to the central cavity that prevents ion conduction.

A comparison of the KvLm PM structure with that of the chimera Kv1.2-Kv2.1 (6) (Fig. 3A), a tandem fusion of two eukaryotic Kv modules (a sensor and a pore), is informative because it underscores significant differences and similarities attributable in part to the removal of the sensor from the KvLm PM. First, the crystallized KvLm-PM contained the S4-S5 linker, yet no density was observed for any of its residues suggesting that the linker was not immobilized at the position found in the eukaryotic chimera (6) (Fig. 3B). For the chimera, the inner helices are more distant and disclose an open activation gate that leads to an unobstructed pore lumen (Fig. 3, B–D); by contrast, in the KvLm PM a narrower permeation path appears occluded by a closed activation gate. Therefore, we surmise that the structure of the activation gate is sensor-dependent.

At variance, the near identical conformation of the selectivity filters in a conductive conformation (Fig. 3E), and the conservation of the lipid immobilization surface at the inter-
subunit crevice (Fig. 3F) suggests that the presence of a voltage sensor module is not required for conservation of these PM features. Importantly, the conservation of the lipid-binding sites (Fig. 3F) implies that both full length and PM are permissive to modulation by membrane lipid composition. Hence, the KvLm PM structures underscore the essential role of the voltage sensor in stabilizing the conductive state of the channel.

There are conspicuous differences between the PM of KvLm and that of the chimera that are not attributable to the missing sensors. Salient among them is that, for KvLm, the extracellular facing surface is flat, whereas in Kv1.2–Kv2.1, the extra-long linker that connects the outer helix and the pore helix protrudes ~15 Å out toward the extracellular milieu and contributes both sterically and electrostatically to the docking of other interacting proteins, including toxins (33–35). In contrast, access to the selectivity filter in KvLm is unobstructed, implying that the structural element connecting outer helix and pore helix may be of modular nature, a site adaptable to respond to different gating modifying signals.

To establish a rigorous connection between structure and function, the KvLm PM was reconstituted in lipid bilayers composed of the lipid used for crystallization (monoolein) and bathed in symmetric 27 mM KCl, thereby mimicking the crystallization condition (“Experimental Procedures”). The single-channel conductance (\( \gamma = 4 \pm 2 \) pS) of all event current amplitude histograms of a 6-min-long recording in symmetric 27 mM KCl, 10 mM Hepes, pH 7.4, at 150 mV. The total number of opening events (N) analyzed was 34,225; \( \gamma = 4 \pm 2 \) pS. \( \gamma \) was calculated from Gaussian fits (black lines) to current amplitude histograms (gray lines); c and o denote closed and open states. D, currents at 100 mV in 0.5 M KCl. E, the section marked by the asterisk in D is displayed at a 1000-fold higher time resolution; \( \gamma = 68 \pm 9 \) pS. F, all event current amplitude histogram of a 5-min-long recording in symmetric 0.5 M KCl, 10 mM Hepes, pH 7.4, at 100 mV. N = 20,200 and \( \gamma = 68 \pm 9 \) pS. G, currents at 100 mV in 0.5 M KCl before addition of blocker. After 2 min at 0 mV, 100 μM tetrabutylammonium (TBA) was injected onto one side of the bilayer (arrow) followed by a 2-min waiting time at 0 mV. H, the voltage was stepped to 100 mV evoking the single-channel currents. I, the segment immediately succeeding the application of the second 100-mV step indicated by the asterisk in H, displayed at 100-fold higher time resolution, demonstrates that the channel is already blocked at the onset of the voltage step. \( \Delta t \) signifies the time interval at 0 mV.

**TABLE 2**

| Experimental condition | 0.5 M KCl | 27 mM KCl |
|------------------------|----------|----------|
| Conductance (pS)       | 68 ± 9   | 4 ± 2    |
| Open occupancy          | 0.1 ± 0.1| 0.3 ± 0.2|
| Closed occupancy        | 0.9 ± 0.1| 0.7 ± 0.2|
| \( \tau_{\text{open}} \) (ms) | 1.3 ± 0.3 | 12 ± 5 |
| \( \tau_{\text{close}} \) (ms) | 1.8 ± 0.5 (0.62) | 13 ± 5 (0.66) |
| \( \tau_{1\text{ close}} \) (ms) | 350 ± 40 (0.38) | 113 ± 50 (0.54) |
| Burst length (ms)       | 14 ± 3   | 70 ± 10  |
| Total openings analyzed | 20,200   | 85,200   |

by the asterisk in Fig. 4I at higher time resolution in Fig. 4E: \( \gamma \) is 68 ± 9 pS (Fig. 4F and Table 2). To explore the gating dynamics, we assessed whether at 0 mV the pore lumen was accessible to the open channel blocker tetrabutylammonium. Fig. 4 (G–I) demonstrates that tetrabutylammonium enters the permeation pathway and blocks the current at 0 mV. This observation establishes that the structure of the KvLm PM in monoolein membranes is that of a pore in transit, quickly sampling multiple conformations before being captured in one of them.

**DISCUSSION**

Functional assays showing PM channel block at 0 mV by the open channel blockers tetrabutylammonium (36) (Fig. 4, D–F)
and 4-aminopyridine (37) (data not shown) demonstrate that despite the crystal structure displaying an unequivocally closed activation gate, the reconstituted PM in lipid bilayers gates open at 0 mV long enough for the blocker to enter the cavity and abort permeation. This implies that the PM structure (a composite of 32 individual crystals) depicts a pore with a mobile activation gate able to open and close yet captured, on average, in a closed conformation. As such, the structure of KvLm PM represents the first structure of a Kv channel pore trapped with the activation gate closed. Comparison with other Kv channel structures that contain both the voltage sensor module and PM and exhibit an open PM implies that the voltage sensor module determines the stability of the open conformation of the PM (Fig. 3).

What is the significance of these data? Three main points emerge from our analysis: 1) The two new structures are the first structures of a channel obtained by crystallization of the protein in its native environment: a lipid bilayer. Specifically for Kvs, it represents the first module dissected out from a whole channel protein subunit that recapitulates function and structure in its native and identical environment: the epitome of a folded functional unit in a membrane. 2) The new structures of KvLm show for the first time a Kv channel pore in a closed conformation. The structural similarity of the KvLm pore to other reported K+ channel pores is indeed striking: they share the same fold and display a conformation of the two gates that has been depicted in another K+ channel pore that of KcsA. However, KcsA is not a Kv channel, nor is it a structural pore module, because both gating sensor and pore are present in the structure. Moreover, the closed structure of KcsA, determined at pH 7.5, has been demonstrated to be closed and stable in such a conformation (38). In contrast, the pore of KvLm is indeed a module given that, in the absence of its gating sensor, it resides in an unstable closed conformation. This may appear at first sight as structurally similar to KcsA, yet it is, in essence, energetically distinct. A comparison with the two available structures of the eukaryotic Kv1.2 pore (Protein Data Bank codes 2R9R and 2A79) (5, 6) reveals another crucial difference: in KvLm, the pore gates are closed, whereas in the other two existing structures the pores are open. 3) The fundamental tenet behind the modular design of Kvs prompts the question of how do the sensor and pore modules interact? One way to answer this question about how the machine operates is to determine whether the structure of the pore alone, in the absence of an activating signal, is closed or open. The demonstration that, in the absence of sensor, the KvLm pore in a lipid bilayer resides in an unstable and functional closed state indicates that the closed and open structures are in close energetic equilibrium. The implication is that the work that the sensor module must do on its attached pore module to elicit the transition is limited. By the same token, it also demonstrates that to keep the channel stable open or closed, the sensor must play an important role. Therefore the new information explains how the two independent modules in a Kv channel interact to generate the observable signal.

Our findings provide compelling evidence that the structure of a K+ channel in lipidic cubic phase can be determined close to atomic resolution. The new findings give further support to nature’s conservation of the pillar principles of protein science in the context of the unique environment conferred by the lipid bilayer for protein folding.

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