

Direct knock-on of desolvated ions governs strict ion selectivity in K⁺ channels

Wojciech Kopec¹, David A. Köpfer¹, Owen N. Vickery^{2,3}, Anna S. Bondarenko⁴, Thomas L. C. Jansen⁴, Bert L. de Groot^{1*} and Ulrich Zachariae^{1,2,3*}

The seeming contradiction that K⁺ channels conduct K⁺ ions at maximal throughput rates while not permeating slightly smaller Na⁺ ions has perplexed scientists for decades. Although numerous models have addressed selective permeation in K⁺ channels, the combination of conduction efficiency and ion selectivity has not yet been linked through a unified functional model. Here, we investigate the mechanism of ion selectivity through atomistic simulations totalling more than 400 μs in length, which include over 7,000 permeation events. Together with free-energy calculations, our simulations show that both rapid permeation of K⁺ and ion selectivity are ultimately based on a single principle: the direct knock-on of completely desolvated ions in the channels' selectivity filter. Herein, the strong interactions between multiple 'naked' ions in the four filter binding sites give rise to a natural exclusion of any competing ions. Our results are in excellent agreement with experimental selectivity data, measured ion interaction energies and recent two-dimensional infrared spectra of filter ion configurations.

Ionic currents through K⁺ channels establish the membrane voltage in all cells and terminate action potentials in electrically excitable cells. K⁺ channels facilitate the passage of K⁺ ions to near-diffusion limited rates, while reliably excluding smaller Na⁺ ions^{1,2}. How K⁺ channels achieve the combination of these seemingly incompatible features has intrigued scientists for decades^{3–7}. A clear separation between K⁺ and Na⁺ currents is necessary to ensure sharp action potentials, facilitating the rapid propagation of electric signals in excitable cells such as neurons. However, despite intense efforts, a unified explanation for how K⁺ channels maintain strict K⁺ selectivity under maximal ion conduction rates has remained elusive.

Ions pass through the K⁺ channels' selectivity filter (SF), which represents their conserved functional core, enabling rapid and selective ion permeation⁸. It contains four successive K⁺ binding sites (S₁–S₄), with two additional binding sites at its extracellular entrance (S_o) and in the water-filled central cavity (S_{cav})⁹ (Fig. 1a). Predating structural elucidation of the channels, the snug-fit model, which is still preferred by most present textbooks^{10,11}, posited that the K⁺ binding sites in a rigid SF provide an unfavourable binding environment for the smaller Na⁺ ions⁴. Upon determination of the first K⁺ channel structures by X-ray crystallography, the electron density observed in the SF K⁺ binding sites was interpreted to reflect alternate occupation with water and ions^{1,12,13}. This conclusion informed subsequent structure-based and computational models^{14–22}, which suggested that ion selectivity arises from the intrinsic flexibility of specific SF binding sites^{14,15}, the collective dynamics of ions and water within the SF^{16,17}, or energetic differences in ion binding^{13,14,16–20}. These models were based on the assumption that permeating K⁺ ions in the SF are separated by water molecules (the 'soft knock-on' mechanism). The majority of previous computational studies identified the SF K⁺ binding sites as thermodynamically K⁺ selective, due to the optimal coordination of a K⁺ ion by eight carbonyl ligands, while Na⁺ binding to the same geometry was found to be energetically less favourable^{14,18–20,23–25}. However, the complete sequence of microscopic events leading to efficient K⁺ permeation and simultaneous Na⁺

exclusion during ion permeation has never been revealed, partially due to the inability to observe high, sustained currents in previous atomistic molecular dynamics simulations²⁶.

Recently, however, a reanalysis of the crystallographic data showed that K⁺ ions form close ion pairs (that is, direct ion–ion contacts) at neighbouring SF ion-binding sites²⁷. Furthermore, accompanying molecular dynamics simulations suggested that the strong electrostatic interaction between ions at short distance plays a key role in establishing the observed high rates of K⁺ ion conduction, while water molecules are largely excluded from permeating the channel²⁷. Since the instantaneous ion occupancies of the SF under 'direct Coulomb knock-on' conditions drastically differ from those of 'soft knock-on'²⁸, the competition between K⁺ and Na⁺ ions, and therefore the mechanism of ion selectivity, is likely to be based on different principles than was previously thought. As it is now possible to directly simulate ion currents and analyse thousands of individual ion permeation events at the atomistic level, we set out to apply this methodology to achieve unfiltered insight into the determinants of ion selectivity in K⁺ channels under actual permeation conditions. Our extended simulations spanning more than 400 μs of simulated time allowed us to analyse in detail over 7,000 individual ion permeation events.

Our results present a novel mechanism of ion selectivity in K⁺ channels, providing a unifying explanation for the two major, seemingly irreconcilable, physiological characteristics of K⁺ channel conduction—highly efficient ion transfer under exquisite K⁺ selectivity. We show that full desolvation of ions in the SF—required to establish close contacts between K⁺ ions in the neighbouring sites and rapid conduction—efficiently excludes Na⁺ ions from permeating the channel. Importantly, in contrast with previous ion permeation models^{29–31}, our findings show that any significant level of water co-permeation invariably leads to drastically diminished ion selectivity and reduced ion conduction rates. Finally, we demonstrate that recent K⁺ channel data from two-dimensional (2D) infrared (2D IR) spectroscopy³⁰, which were originally interpreted to be exclusively compatible with filter occupancies under water

¹Biomolecular Dynamics Group, Max Planck Institute for Biophysical Chemistry, Göttingen, Germany. ²Computational Biology, School of Life Sciences, University of Dundee, Dundee, UK. ³Physics, School of Science and Engineering, University of Dundee, Dundee, UK. ⁴University of Groningen, Zernike Institute for Advanced Materials, Groningen, The Netherlands. *e-mail: bgroot@gwdg.de; u.zachariae@dundee.ac.uk

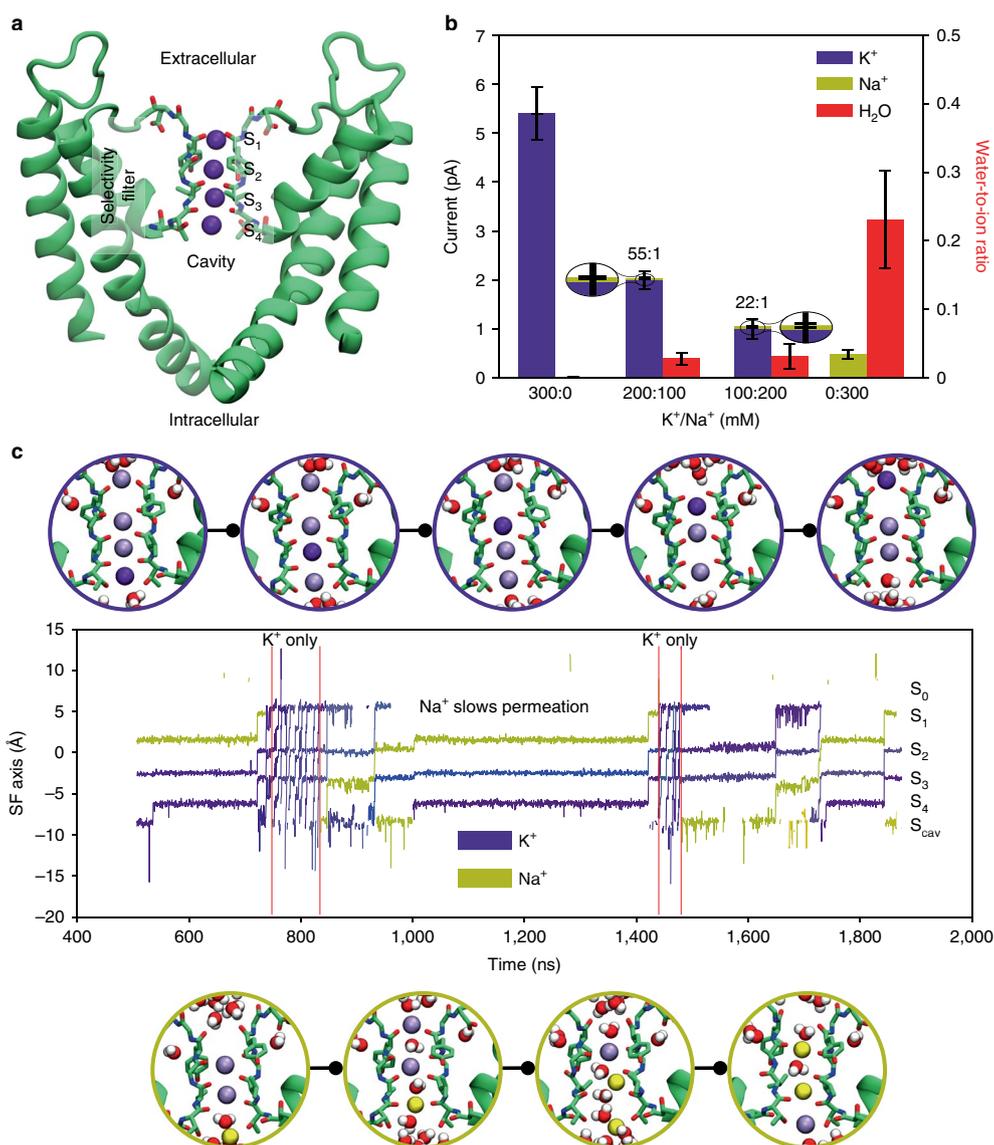


Fig. 1 | Ion selectivity of KcsA. **a**, Open-state form of KcsA with the main K⁺ binding sites of the SF (S₁–S₄) at the interface between the four channel subunits. **b**, KcsA currents and water-to-ion permeation ratios at a voltage of ~ 220 mV in solutions of 300 mM ionic strength containing both K⁺ and Na⁺ ions. At a 2:1 excess of K⁺, less than 2% of the current originates from Na⁺ flux (K⁺, purple; Na⁺, yellow), corresponding to a selectivity of 55:1 for K⁺ over Na⁺. At a 2:1 excess of Na⁺, less than 5% of all conduction events are caused by Na⁺. Only in the total absence of K⁺ ions is a more substantive Na⁺ current seen, although this is much smaller than in pure K⁺ solutions. Error bars depict s.e.m. (see also Supplementary Table 1). **c**, Representative traces of ions traversing the SF of KcsA. The positions in the SF of K⁺ ions (purple) and Na⁺ ions (yellow) are shown versus simulated time. During periods in which Na⁺ binds to the SF, the overall permeation rate is slowed down drastically, whereas in the absence of Na⁺ highly efficient K⁺ permeation takes place. Representative instantaneous SF occupancies are shown for K⁺ (purple circles) and Na⁺ permeation (yellow circles), respectively. K⁺ and Na⁺ ions are displayed as purple and yellow spheres. The permeating K⁺ ion is marked in a darker colour, and water molecules are shown as red and white spheres.

co-permeating conditions, are equally compatible with states displaying directly contacting ions in the SF.

Results

The direct Coulomb knock-on is intrinsically ion selective. We conducted atomistic simulations to directly investigate the origins of ion selectivity and the coupling of ion and water flux under K⁺ channel permeation conditions, employing both electrochemical ion gradients and applied electric fields to generate membrane voltages in the range of 220–300 mV. These acted as driving forces for the permeation of K⁺ and Na⁺ ions across a variety of channels including KcsA, MthK, Kv1.2, NaK2K and NaK2CNG-N. The data we collected comprise more than 7,000 ion permeation events in mixed and pure K⁺ and Na⁺ solutions (Supplementary Tables 1–9).

First, we focused on the most studied K⁺ channel, KcsA (Fig. 1a). The channel current in KcsA is strongly dominated by permeating K⁺ ions in all K⁺-containing solutions (Fig. 1b). In mixed solutions of a 2:1 (K⁺/Na⁺) ratio, the KcsA current shows a K⁺/Na⁺ permeability ratio of 55:1. Even under excess of Na⁺ (1:2 K⁺/Na⁺ mixture), KcsA favours K⁺ permeation with a permeability quotient of 22:1. Only in the complete absence of K⁺ is a residual Na⁺ current of $\sim 15\%$ of the maximum K⁺ flux observed.

K⁺ ions permeate the channel in a fully dehydrated state, consistent with the ‘direct Coulomb knock-on’ mechanism. In pure K⁺ solution, we therefore do not observe any co-permeation of water (Fig. 1b). In contrast, Na⁺ ions co-permeate with water. As a consequence, few water molecules are carried along during the rare event of Na⁺ traversing the SF (Fig. 1c). In both the 2:1 and 1:2 (K⁺/Na⁺)

mixed solutions, the resulting overall water-to-ion flux ratio is about 0.03 ± 0.02 (Fig. 1b). In pure Na^+ solution, the raised Na^+ current leads to increased water permeation, with a water-to-ion flux ratio of 0.22 ± 0.08 .

Experimental studies have reported K^+ channel Na^+/K^+ permeability ratios between 0.006 and 0.04 (refs 5,6,32). The simulated Na^+/K^+ ratios of 0.02–0.04 are thus in good agreement with available experimental data. Furthermore, for KcsA, we find a reduction in the outward K^+ ion current to about one-third of that in pure KCl for the 2:1 (K^+/Na^+) concentration regime, whereas it decreases to about 20% under Na^+ excess (1:2 mixture; Fig. 1b). This observation reflects the experimental observation of K^+ channel block by intracellular Na^+ , which has been widely reported in single-channel electrophysiology studies^{33,34} and is clearly visible in the ionic traces through the SF (Fig. 1c).

The SF of many K^+ channels, including KcsA, undergoes a conformational change at K^+ concentrations below ~ 20 mM on timescales of up to seconds, preventing ion flux under low- K^+ conditions^{1,35,36}. This filter gating or collapse provides an additional layer of ion selectivity to K^+ channels. However, on the shorter timescale of our simulations, the SF remains in the conductive state. Our observations made in pure Na^+ solutions are therefore compared to experiments on the constitutively conductive SF variant, KcsA^{D-Ala77}, in which filter collapse is sterically blocked. In experiments, this semisynthetic channel form conducts Na^+ when K^+ is completely absent, with currents similar to those we observe, while—as in our simulations—the addition of K^+ disrupts Na^+ conduction³⁵. The experimentally recorded ratio between maximum K^+ and residual Na^+ current in KcsA^{D-Ala77} is in remarkably good agreement with our findings³⁵.

Strict K^+ selectivity is coupled to the exclusion of water. Next, we expanded our investigation to the bacterial MthK channel and eukaryotic Kv1.2 channel (W362Y variant), as well as the engineered channels NaK2CNG-N and NaK2K (Fig. 2a). NaK2K and NaK2CNG-N are both generated by introducing mutations in the SF of the non-selective bacterial ion channel NaK^{37,38}. The SF of NaK possesses only two ion-binding sites, which are chemically equivalent to sites S_3 and S_4 of canonical K^+ channels, and displays bound water molecules within a widened vestibule^{39,40}. In NaK2CNG-N, the NaK SF is modified to yield three consecutive ion-binding sites (S_2 – S_4), while in NaK2K, the mutations reconstruct a canonical K^+ channel SF with four consecutive ion-binding sites³⁸. In experiments, MthK, Kv1.2 and NaK2K are strongly K^+ selective, while NaK2CNG-N shows similar permeability for both K^+ and Na^+ ^{41,42}.

The diminished K^+ selectivity of the NaK and NaK2CNG channels has previously been associated with the reduced number of SF ion-binding sites³⁸, increased hydration level of ions^{22,43,44}, for instance, as seen in the NaK crystal structure^{39,40}, and enhanced structural plasticity of the SF⁴⁵. This led to the conclusion that only channels with four consecutive ion-binding sites in the SF can ensure fully K^+ -selective ion permeation. However, the mechanistic basis for this phenomenon has remained unclear, in particular under permeation conditions^{38,42,46}.

In our simulations, we find strictly K^+ -selective current in KcsA, MthK, NaK2K and Kv1.2 W362Y (Fig. 2b). The total channel current shows some differences for pure K^+ solution. In mixed K^+/Na^+ solutions, the current overwhelmingly arises from K^+ permeation events, both under excess of K^+ and Na^+ . In pure Na^+ , only KcsA and NaK2K give rise to appreciable Na^+ fluxes.

In contrast, NaK2CNG-N shows greatly reduced K^+ selectivity with considerable Na^+ permeation in the K^+/Na^+ mixtures, as well as the largest level of Na^+ current in pure Na^+ among all of the investigated channels. It also displays the lowest K^+ current in pure K^+ solution. Notably, the Na^+ current is linked to a substantially raised level of water co-permeation. While the water-to-ion flux ratio is very small in all of the strictly K^+ selective channels (below 0.05 in

all regimes in which K^+ is present), the absence of the fourth binding site in NaK2CNG-N leads to a much-diminished exclusion of water, during both Na^+ and K^+ permeation, and thus allows both ion types to traverse the SF with similar probability. We observe a water-to-ion ratio of >1 under all ionic regimes in NaK2CNG-N, showing that a significant part of the ions' solvation shell is retained in the SF during permeation. These observations are in accordance with previous experiments, in which mutations reducing the number of K^+ binding sites in the SF from four to three abolished K^+ selectivity^{38,42}, and explain these experimental observations by insufficient exclusion of hydrating water around the ions. Structurally, we find that increased co-permeation of water is underpinned by an enhanced flexibility of the NaK2CNG-N SF with regard to filters with four ion-binding sites. These dynamics cause a small widening of the SF on average, and thereby slightly relax the strict geometric constraints for ion transfer, especially at the central ion-binding site. Similar findings have recently been reported for the related NaK channel⁴⁵.

Fully desolvated ions in the SF are in agreement with 2D IR spectroscopy experiments. All of these observations suggested that the strict exclusion of water, and ion conduction by the direct Coulomb knock-on mechanism, underpin ion selectivity in K^+ channels under permeation conditions. We therefore re-examined a recent investigation, in which equilibrium molecular dynamics simulations were used to interpret 2D IR spectra on the isotope-labelled SF of synthetic KcsA under no voltage. In this study, it was found that only infrared spectra that were predicted from simulations of water-and-ion-occupied SFs were compatible with the experimental data, after fitting the relative population of each state and additionally taking carbonyl-flipped SF conformations into consideration³⁰.

In contrast, SF ion populations under non-equilibrium ion flux conditions were not fitted to match the infrared data, but rather taken directly from our previous simulations, which were conducted under voltage²⁷. As shown in the present work, these K^+ conducting states display direct ion–ion contacts. The corresponding spectra of these conducting states were interpreted to disagree with the infrared experiments, which led to the conclusion that water molecules are required to accompany K^+ ions during ion permeation.

To reconcile this apparent discrepancy, we used our previously reported ion configurations (together with those shown here) in the ion-conductive SF, and equilibrated these states further at 0 mV to replicate the conditions of the infrared experiment (see Supplementary Methods and Supplementary Table 10). We then followed the same protocol employed in previous studies³⁰ to calculate the final infrared spectra. Spectra calculated from a weighted (fitted) combination of these states yield good agreement with the experimental spectra, similar to a weighted sum of the water-containing states described previously and repeated independently by us (the final spectra are shown in Fig. 3 and Supplementary Fig. 4). Furthermore, when configurations with only direct ion–ion contacts in the SF are taken into consideration, a wide range of spectral parameters, such as peak positions, nodal slopes and intensity ratios, are also consistent with the experimental data (Table 1). We conclude that the analysis based on 2D IR spectra cannot differentiate between the two scenarios of SF occupancy and thus could support both mechanisms of ion permeation with and without co-permeating water.

How does ion dehydration contribute to selectivity? The number of K^+ and Na^+ ions traversing KcsA from the intracellular side during conduction shows strong divergence upon SF entry (Supplementary Fig. 6). The fraction of Na^+ ions migrating from S_4 to S_3 decreases by about 18-fold, while that of K^+ remains nearly constant. Overall, in pure KCl, 44% of all K^+ ions entering the SF at S_4 eventually permeate fully, compared with only 2.5% of Na^+ ions in pure NaCl (the presence of K^+ reduces this proportion further to less than 1%; Supplementary Fig. 7).

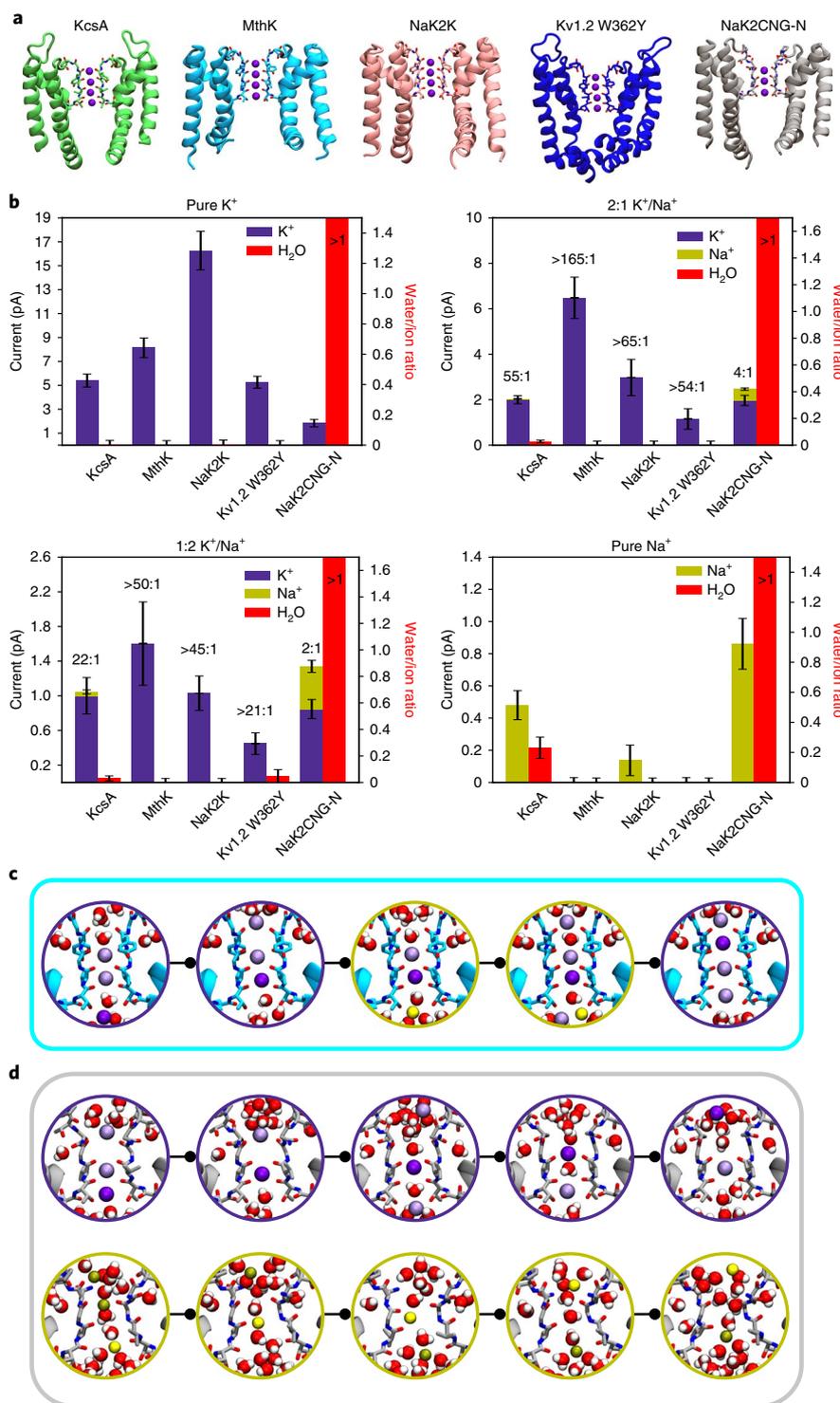


Fig. 2 | Relationship between conduction efficiency, ion selectivity and water co-permeation. **a**, The selective channels KcsA, MthK, NaK2K and Kv1.2 W362Y contain four SF ion-binding sites, whereas the non-selective NaK2CNG-N possesses three. **b**, Currents and water-to-ion flux ratios at voltages of 220–280 mV in pure and mixed K⁺/Na⁺ solutions (error bars depict s.e.m.). KcsA, MthK, NaK2K and Kv1.2 W362Y show high K⁺ currents in all solutions, lowered by the presence of Na⁺ in a concentration-dependent manner (sodium block). MthK, NaK2K and Kv1.2 W362Y strictly exclude Na⁺ permeation, with the exception of NaK2K in pure Na⁺ solution. The water-to-ion flux ratio is minimal in all selective channels. Conversely, NaK2CNG-N displays a high level of water permeation in all solutions (a water-to-ion flux ratio >1). Water molecules in the SF lead to reduced K⁺ currents. **c**, Simulation snapshots showing outward K⁺ permeation events, Na⁺ block and relief of Na⁺ block in MthK (K⁺ ions, purple spheres (with the permeating ion the darkest); Na⁺, yellow spheres; water molecules, red and white spheres). Here, Na⁺ does not reach S₄ and is instead replaced by K⁺. **d**, Simulation snapshots of K⁺ (top, purple circles) and Na⁺ permeation in NaK2CNG-N (bottom, yellow circles). Direct ion-ion contacts form transiently, but permeation in NaK2CNG-N occurs with intervening water molecules. Periods where only one ion is present in the SF lead to higher water permeation levels. Na⁺ ions cross the channel partially hydrated, allowing more water molecules to enter the SF. Control simulations show the robustness of these observations for a wide range of force-field parameters (Supplementary Figs. 1 and 2). Simulation snapshots for NaK2K and Kv1.2 W362Y are displayed in Supplementary Fig. 3.

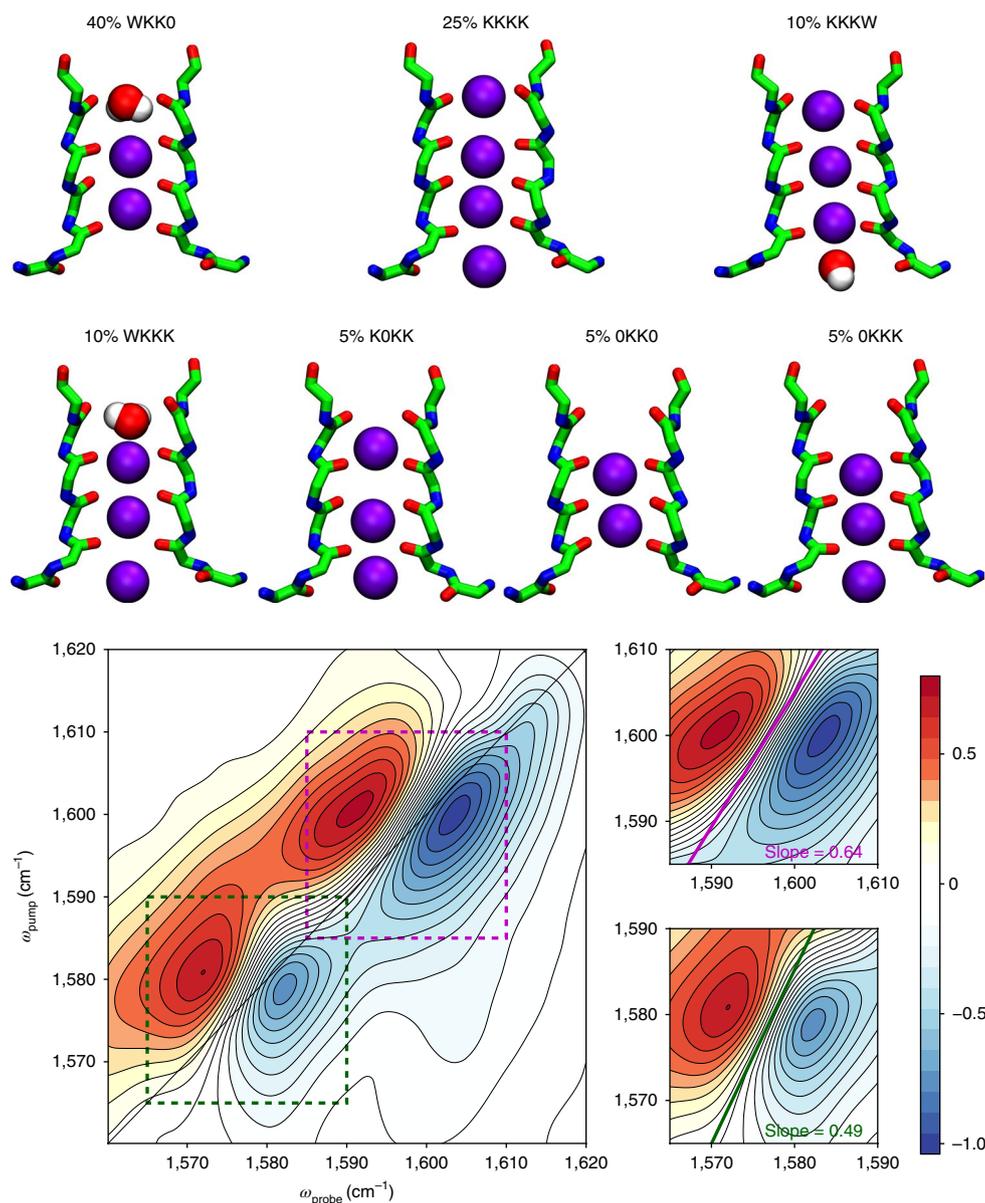


Fig. 3 | Calculated 2D IR spectrum for occupancy states characteristic of the direct Coulomb knock-on conduction mechanism. Weights used to generate the spectrum are given above the schematic representation of each occupancy. The spectrum displays two pairs of peaks at $(\omega_{\text{probe}}, \omega_{\text{pump}}) = (1,600 \text{ cm}^{-1}, 1,603 \text{ cm}^{-1})$ and $(\omega_{\text{probe}}, \omega_{\text{pump}}) = (1,579 \text{ cm}^{-1}, 1,584 \text{ cm}^{-1})$. These peaks are in good agreement with the experimentally reported values— $(\omega_{\text{probe}}, \omega_{\text{pump}}) = (1,603 \text{ cm}^{-1}, 1,610 \text{ cm}^{-1})$ and $(\omega_{\text{probe}}, \omega_{\text{pump}}) = (1,580 \text{ cm}^{-1}, 1,584 \text{ cm}^{-1})$ —when, notably, only states with direct ion-ion pairs in the SF are taken into account. The spectrum is a linear combination of the spectra of individual states that are frequently visited under permeation conditions (Supplementary Fig. 5). The weights were derived following a protocol from Kratochvil et al.³⁰.

Since Na^+ interacts more strongly with water than K^+ , the free energy required to fully dehydrate Na^+ ions exceeds that of K^+ by 74.8 kJ mol^{-1} (see Supplementary Methods and Supplementary Fig. 8). This energy difference contributes to the thermodynamic basis for ion selectivity at every SF site. Free-energy calculations on occupancy states with direct ion-ion contacts (that is, states that lead to high-efficiency ion permeation under voltage (Supplementary Fig. 8)) show that all of the internal SF K^+ binding sites in KcsA are K^+ selective when this dehydration free energy is taken into account (Supplementary Fig. 8). In particular, the SF entry sites S_4 and S_1 favour K^+ binding by up to 27.6 kJ mol^{-1} . In contrast, sites in-plane with the filter carbonyl groups between the K^+ binding positions remain slightly Na^+ selective, in agreement with our findings from ion flux simulations.

The SF is too narrow to allow ions to permeate in the presence of their full hydration shells. Most ions therefore enter and pass

through the SF either in a partially or fully dehydrated state. The vast majority of permeation events we observe follow the ‘direct Coulomb knock-on’ mechanism, where K^+ ions traverse the SF of selective channels without hydrating water.

In contrast, the classic ‘soft knock-on’ mechanism, in which one water molecule permeates along with each ion in the SF, would compromise the maximal dehydration of permeating ions. According to our observations, the thermodynamic basis for ion selectivity is thus optimized when the maximum number of solvating water molecules is removed upon entry into the SF. Simultaneously, direct contacts between fully dehydrated ions lead to maximally efficient ion permeation in our simulations. We therefore identify the complete loss of the ion hydration shell during SF entry—induced by the unique architecture of the SF—as the major contributing factor underpinning both ion selectivity and high conduction rates.

Table 1 | Properties of experimental and calculated 2D IR spectra

Spectrum	Comparison of infrared spectra			Intensity ratio
	Centre ω_{pump} (cm ⁻¹)	Centre ω_{probe} (cm ⁻¹)	Slope	
Experimental ^a	1,603	1,610	0.58 (0.01)	1.25
	1,580	1,584	0.60 (0.11)	
Soft knock-on ^a	1,603	1,607	0.51 (0.03)	1.25
	1,580	1,586	0.52 (0.09)	
Soft knock-on ^b	1,609	1,612	0.58 (0.10)	1.27
	1,570	1,575	0.60 (0.10)	
Direct knock-on	1,600	1,603	0.64 (0.10)	1.33
	1,579	1,584	0.49 (0.10)	

^aData reported in Kratochvil et al.³⁰. ^bOur data.

Thermodynamic and kinetic factors of selective ion permeation within the SF. Finally, we aimed to resolve the processes that contribute to selectivity when residual Na⁺ ions compete with K⁺ ions within the SF. To distinguish thermodynamic effects (for example, differential affinity of K⁺ and Na⁺ to filter binding sites)²⁵ from kinetic effects, which may inhibit the passage of ion types between the binding sites⁷, we recorded K⁺ and Na⁺ density profiles in the SF, based on KcsA simulation data under various ion concentration regimes (Supplementary Fig. 9). Although these density profiles do not strictly reflect equilibrium free energies, we use them here as an approximate measure to quantify differences in binding affinity for K⁺ and Na⁺ in the SF (manifested as different density minima) and kinetic barriers (peak heights).

Due to their high K⁺ affinity, the SF binding sites S₄–S₀ show strong K⁺ density in pure KCl solutions. Lower minima for K⁺ compared with Na⁺ in mixed K⁺/Na⁺ conditions demonstrate the channel's thermodynamic K⁺ selectivity, which we also observe in free-energy calculations (Supplementary Fig. 8). Considering transition kinetics, large barriers occur for Na⁺ transfer between S₃ and S₂ and near S₃; however, they do not exceed the maximum barrier to K⁺ translocation between S₄ and S₃. Only under excess of Na⁺ are slightly raised kinetic barriers for Na⁺ transfer observed.

However, in contrast with K⁺ ions, the Na⁺ density shows additional Na⁺ binding sites between the canonical K⁺ positions, which are located in-plane with the backbone carbonyl oxygen atoms. These sites can be observed even in the absence of K⁺; however, under K⁺ excess, Na⁺ occupation mainly shifts to positions between the K⁺ binding sites S₃ and S₄, and S₁ and S₂ (Supplementary Fig. 9). As a consequence, the distance between two favourable Na⁺ binding positions in the SF centre increases to ~7 Å—more than twice the distance observed for a K⁺ ion pair at S₂ and S₃, which we have found to be crucial for high-speed conduction of K⁺²⁷.

The simultaneous presence of K⁺ and Na⁺ in the SF therefore has a major effect on the SF occupancy pattern, as under excess of K⁺, binding of Na⁺ between the K⁺ positions leads to a disruption of the optimal ion–ion distance for efficient conduction. Accordingly, Na⁺ ions cannot permeate the channels at high kinetic rates to generate sizable Na⁺ currents, and instead, they block K⁺ conduction, as observed in experiments^{33,34} and also suggested by calculations of potentials of mean force in a previous computational study²³.

Since the typical ion concentration of K⁺ is 140 mM and that of Na⁺ is near 10 mM in the cytoplasm⁴⁷, only few Na⁺ ions would typically compete with an excess of K⁺ ions during their physiological outward permeation. As there is a higher concentration of Na⁺ on the extracellular side of the membrane, we also investigated the selectivity for extracellular Na⁺ upon outward (and also inward) K⁺ flux in additional sets of simulations (Supplementary Figs. 10 and 11), confirming strict selectivity also under these conditions.

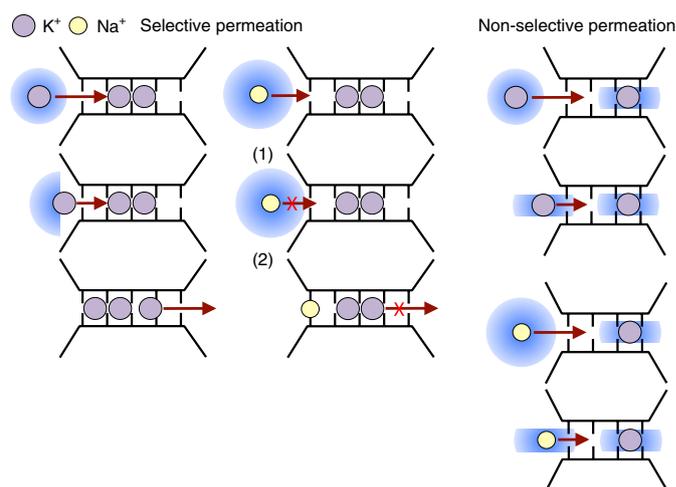


Fig. 4 | Schematic of the mechanisms of K⁺ ion selective and non-selective channel permeation. In selective K⁺ channels containing four SF binding sites, ions are observed to permeate largely without their solvation shell (blue). In contrast, we find that non-selective permeation in channels with three SF ion-binding sites is based on partial retention of the hydration shell around the ions. The desolvation penalty associated with dehydration of Na⁺ ions prevents most Na⁺ ions from SF entry (centre, (1)) in the K⁺ selective channels. Na⁺ ions that surmount this barrier in K⁺ selective channels tend to adopt positions between the K⁺ binding sites, adding to their ion selectivity (centre, (2)) by impeding efficiently permeating (direct knock-on) conditions.

Discussion

The results we present demonstrate an intimate relationship between ion selectivity, rapid ion conduction and the dehydration of permeating ions in K⁺ channels (Fig. 4). They also highlight the additional role of complex multi-ion interactions between the preferred K⁺ ions and competing Na⁺ ions within the SF for enhanced selectivity.

Across a range of different channel types, we show that the level of water co-permeation in the SF is closely linked to the ion selectivity and conduction efficiency of the channels. An increased amount of water in the SF invariably decreases both conduction rates and ion selectivity (Fig. 2 and Supplementary Figs. 12–14). Channels with only three K⁺ binding sites in their SF do not dehydrate permeating ions to the same degree as the canonical K⁺ channels, which possess four stacked ion-binding positions in their filter. As a consequence, they lack the exquisite ion selectivity of the canonical K⁺ channels (Fig. 4). The selectivity for K⁺ is optimized when the ions are maximally dehydrated, as observed, for example, for the KcsA, MthK and NaK2K channels. Notably, Na⁺ channels possess a much wider SF (~8 Å) than K⁺ channels (~3 Å)⁴⁸, and consequently, Na⁺ ions permeate these channels in a mostly hydrated form⁴⁹.

In all of the K⁺ channel types we investigated, Na⁺ ions typically cross the SF together with some remaining hydrating water molecules. However, in the SF of the selective channel types, competing K⁺ ions displace residual Na⁺ ions to positions located between the crystallographically determined K⁺ binding sites, where they cannot fulfil the criterion for highly efficient ion conduction; that is, ion pair formation in the K⁺ binding sites (Fig. 4, centre). Free-energy calculations also identify these intermediate sites as preferred positions for Na⁺ binding (Supplementary Fig. 8). These results, together with the data we obtained from analysing ~7,000 individual permeation events, therefore challenge the snug-fit model, which posits that the smaller Na⁺ is coordinated less optimally than K⁺ by the carbonyl groups of the canonical SF K⁺ binding sites. This model is, to date, cited most often as the basis for K⁺ channel selectivity in current textbooks^{10,11} despite having been disproven many times^{14,18,19,23,50}. Our results now combine aspects of the previously proposed alternative mechanisms into a unified model, which simultaneously explains rapid ion permeation and selectivity by multi-ion, direct Coulomb knock-on.

This mechanism is in agreement with a wide array of available experimental data, including crystallographic, electrophysiological and spectroscopic investigations. By calculating 2D IR spectra for a large number of possible occupancy states of the SF, and using the same averaging scheme as was used in previous work³⁰, we demonstrate that states that exclusively display direct ion–ion contacts in the SF are fully compatible with recent experimental 2D IR spectra. Notably, the carbonyl-flipped states that are necessary to explain the infrared data on the basis of water-separated ion configurations in the SF correspond to non-conductive filter states in our simulations in all K⁺ channels except the non-selective NaK2CNG-N. Moreover, our observations explain the results from previous experiments studying the interactions of a range of cations with K⁺ channels (see Supplementary Text)⁵¹. We therefore obtain a convergent picture of high-conductance K⁺ channel permeation and exquisite ion selectivity based on maximally dehydrated ions, which occupy neighbouring positions in a linear array of at least four SF ion-binding sites. In this unified model, efficient throughput and strict selectivity are no longer contradictory, but arise as two necessary consequences of a single mechanism.

Methods

Computational electrophysiology simulations. We studied spontaneous ion permeation through potassium channels, in pure and mixed K⁺/Na⁺ solutions, driven by the transmembrane voltage induced by charge imbalances as implemented in the CompEL (computational electrophysiology) method in GROMACS^{52,53} or by applied external electric fields⁵⁴. For open-state KcsA, we used the X-ray structure (Protein Data Bank (PDB) ID: 3f5w⁵⁵), with the SF in the conductive configuration (PDB ID: 1k4c²), embedded in a patch of a 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine (POPC) membrane, as reported in our previous simulations²⁷.

For open-state MthK, we used the high-resolution structure (PDB ID: 3ldc (ref. 56)). For open-state Kv1.2, we used the pore domain from the ‘paddle-chimera’ Kv1.2-Kv2.1 structure (PDB ID: 2r9r), which remains open in simulations at positive voltages⁵⁶. For the open-state NaK2K and NaK2CNG-N, we used high-resolution structures (PDB IDs: 3ouf and 3k06, respectively). All simulations were carried out with GROMACS 5.0 or 5.1 (refs 57–60), using the dual membrane setup typical of the CompEL scheme, with an ionic imbalance between the compartments of 2e⁻, yielding a transmembrane voltage of ~220 mV, or by applying an external electric field⁵⁴, yielding a transmembrane voltage of ~280 mV. The ionic currents were estimated by counting the number of ions permeating the SF per time unit. A list of all simulations is shown in Supplementary Tables 1–9. All simulation details and data on other channels are provided in the Supplementary Information. Kv1.2 W362Y was used instead of wild-type Kv1.2 due to SF instabilities in molecular dynamics simulations (see Supplementary Fig. 14). An equivalent mutation in Kv1.6 yields a channel with almost the same conduction and selectivity properties as wild-type Kv1.6 (ref. 61) (see also Supplementary Information). In NaK2K and NaK2CNG-N, the F92A mutation was used to increase the ionic current, as in previous experiments⁴².

2D IR spectral calculations. In our 2D IR calculations, we used a single KcsA channel embedded in a POPC membrane, similar to the system reported in Kratochvil et al.³⁰. We used the SF occupancy states observed in our computational electrophysiology simulations and compared them with the experimental

spectrum. To do so, a number of specific occupancy states of the SF, corresponding to ion-conductive configurations, were selected (see Supplementary Table 3), and further equilibrated at 0 mV for 10 ns per state, to mimic the conditions of the infrared experiments. Subsequently, a number of snapshots (at least 9 per single occupancy state) were randomly selected from the last 2 ns of equilibration. Each snapshot was then used as a starting point for a simulation of 1 ns length, during which the positions of atoms were saved every 20 fs. These trajectories were then used for spectral calculations. The spectral calculations were performed by first extracting the amide I Hamiltonian, transition dipoles and site frequencies from the 1 ns trajectories. Next, the amide units corresponding to those labelled in the experiment³⁰ were selected and their frequencies were shifted by -66 cm^{-1} to account for the isotope label. One-dimensional and 2D IR spectra were calculated with the numerical integration of the Schrödinger equation method^{62,63}. The linear one-dimensional spectrum was then obtained by a Fourier transform of the linear response function, while the 2D spectrum was obtained with a 2D Fourier transform with respect to the coherence times. The full computational procedure is described in the Supplementary Information.

Free-energy calculations. We performed free-energy calculations for individual binding sites in the SF of KcsA to assess their thermodynamic selectivity between K⁺ and Na⁺ ions for the occupancy states most frequently visited during ion permeation. Consequently, we focused on KK0K and 0KKK occupancies (Supplementary Fig. 8). Three snapshots per occupancy pattern were selected from the computational electrophysiology simulations, transformed back to a single membrane setup and further equilibrated for 40 ns at 0 mV. The final snapshots from these simulations were then used for K⁺ to Na⁺ alchemical free-energy calculations to obtain $\Delta G_{\text{K,Na}}^{\text{site}}$ for each occupied site. To assess the existence and selectivity of potential Na⁺ binding sites in the SF, as previously suggested^{13,25}, we introduced a single Na⁺ ion instead of a K⁺ ion at each site for both starting occupancies, ultimately resulting in the following occupancies: KNaK, NaKK and KKNa (Fig. 3b). After 40 ns of equilibration, the introduced Na⁺ ions bound to their preferred binding sites, in-plane with four SF carbonyl oxygen atoms. The final snapshots from these simulations were then used for Na⁺ to K⁺ alchemical free-energy calculations. To calculate the free-energy differences between K⁺ and Na⁺, we employed the free-energy code in GROMACS. The free-energy differences were calculated using the multistate Bennett acceptance ratio method⁶⁴, as implemented in the ‘alchemical analysis’ utility⁶⁵. The full computational procedure is described in the Supplementary Information.

Code availability. Custom computer code used for simulation analysis is available from the authors upon request.

Data availability. All relevant data are available from the authors upon request.

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Author contributions

U.Z. and B.L.d.G. conceived and supervised the project. W.K. performed and analysed the ion channel simulations under ion gradients and voltage, free-energy simulations, and infrared spectral calculations. D.A.K. performed and analysed the KcsA and initial MthK simulations. O.N.V. performed and analysed additional ion channel simulations under voltage. A.S.B. assisted with the infrared spectral calculations. T.L.C.J. designed and supervised the spectral infrared calculations. W.K., D.A.K., O.N.V. and U.Z. prepared the figures. U.Z., B.L.d.G. and W.K. wrote the manuscript with comments from all authors.

Competing interests

The authors declare no competing interests.

Additional information

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Correspondence and requests for materials should be addressed to B.L.d.G. or U.Z.

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