The permeation of potassium ions through the lipid scrambling path of the membrane protein nhTMEM16

Xiaolu Cheng1, George Khelashvili1,2† and Harel Weinstein1,2*†

1Department of Physiology and Biophysics, Weill Cornell Medicine, New York, NY, United States, 2Institute for Computational Biomedicine, Weill Cornell Medicine, New York, NY, United States

The TMEM16 family of transmembrane proteins includes Ca2+-activated phospholipid scramblases (PLS) that can also function as non-selective ion channels. Extensive structural and functional studies have established that a membrane-exposed hydrophilic groove in TMEM16 PLS can serve as a translocation pathway for lipids. However, it is still unclear how the TMEM16 PLS conduct ions. A "protein-delimited pore" model suggests that ions are translocated through a narrow opening of the groove region, which is not sufficiently wide to allow lipid movement, whereas a "proteolipidic pore" model envisions ions and lipids translocating through an open conformation of the groove. We investigated the dynamic path of potassium ion (K+) translocation that occurs when an open groove state of nhTMEM16 is obtained from long atomistic molecular dynamics (MD) simulations, and calculated the free energy profile of the ion movement through the groove with umbrella sampling methodology. The free energy profile identifies effects of specific interactions along the K+ permeation path. The same calculations were performed to investigate ion permeation through a groove closed to lipid permeation in the nhTMEM16 L302A mutant which exhibits a stable conformation of the groove that does not permit lipid scrambling. Our results identify structural and energy parameters that enable K+ permeation, and suggest that the presence of lipids in the nhTMEM16 groove observed in the simulations during scrambling or in/out diffusion, affect the efficiency of K+ permeation to various extents.

KEYWORDS
phospholipid scramblase, molecular dynamics simulations, potential of mean force (PMF) calculations, free energy profiles, umbrella sampling method, open groove permeation

Introduction

The TMEM16 (Anoctamin) family proteins are involved in a wide range of physiological processes in eukaryotic organisms. Some family members, such as TMEM16A (Caputo et al., 2008) and TMEM16B (Stephan et al., 2009), are Ca2+-activated Cl− channels required for normal electrolyte and fluid secretion in various key physiological functions, e.g., mediating olfactory amplification (Stephan et al., 2009). Others, such as TMEM16F and TMEM16K, have dual functions as non-specific Ca2+-
activated phospholipid scramblases (i.e., they scramble phospholipids with various headgroups) and non-selective ion channels (Suzuki et al., 2010; Yang et al., 2012; Bushell et al., 2019). While not all the physiological functions of mammalian TMEM16 proteins have been elucidated thus far, mutations have been associated with various disorders (Di Zanni et al., 2018; Wanitchakool et al., 2017). Inferences from the structure of various TMEM16 family members determined with X-ray or cryo-electron microscopy (cryo-EM) (Brunner et al., 2014; Paulino et al., 2017a; Paulino et al., 2017b; Dang et al., 2017; Alvadia et al., 2019; Bushell et al., 2019; Falzone et al., 2019; Feng et al., 2019; Kalienkova et al., 2019; Khelashvili et al., 2019; Falzone et al., 2022) have reached important mechanistic insights into both ion conduction and lipid scrambling functions by these proteins. A polar groove identified in the X-ray structure of the two protomers of the fungal scramblase nhTMEM16 (Brunner et al., 2014) was first proposed to be the lipid translocation pathway (Figure 1). Subsequent structural studies of nhTMEM16 constructs showed that the groove adopts different conformations in the presence and absence of Ca$^{2+}$, including an intermediate conformation between the open and closed forms (Kalienkova et al., 2019; Khelashvili et al., 2019). The structure of TMEM16A, a mammalian Ca$^{2+}$-activated Cl$^{-}$ channel, revealed a closed groove in the presence of Ca$^{2+}$ (Paulino et al., 2017a; Paulino et al., 2017b; Dang et al., 2017). In contrast, aTMEM16 scramblase, another fungal homologue, was found to adopt an open-groove conformation in the presence of Ca$^{2+}$ even in a membrane environment in which the scrambling activity is low (Falzone et al., 2019). Of the mammalian scramblases, the structure of nTMEM16F (Alvadia et al., 2019; Feng et al., 2019) has thus far been determined only with a closed groove in the presence of Ca$^{2+}$, while TMEM16K, a scramblase resident in the endoplasmic reticulum, exhibits open and closed conformations of the groove in the presence and absence of Ca$^{2+}$, respectively (Bushell et al., 2019).

Computational studies using molecular dynamics (MD) simulations based on the structural information available for the various TMEM16 family members (Bethel and Grabe, 2016; Jiang et al., 2017; Lee et al., 2018; Bushell et al., 2019; Khelashvili et al., 2019; Khelashvili et al., 2020) have aimed to complete the mechanistic picture of lipid and ion translocations by these proteins. Results from our previous μs-long all-atom MD studies of structure-function relations of nhTMEM16 (Lee et al., 2018; Khelashvili et al., 2019; Khelashvili et al., 2020) have shown that the network of polar residues on the extracellular side of the groove region (residues R432/E313/ E318 and Y439/T333, see Figure 1) functions as gates that modulate the exchange of lipids between the groove and the bulk membrane. The long simulations trajectories also revealed events of ion passage through the groove in the absence of artificially applied voltage across the membrane. This was intriguing finding, as experiments and computations have shown that ion conductance in TMEM16 scramblases is facilitated in membranes under applied voltage (Jiang et al., 2017). Here we pursue this opportunity to investigate the molecular mechanisms associated with ion permeation in these scramblases in the structural context of the conformations visited in the dynamics of the nhTMEM16 scramblase.

Early data have led to the suggestion that ion channel activity of the nhTMEM16 scramblase is mediated by a protein-delimited pore generated according to a proposed “alternating pore-cavity”

![Figure 1](image-url)
model (Kalienkova et al., 2019; Khelashvili et al., 2019). Thus, structure determination showed that a much narrower opening of the groove region in an intermediate conformation can create a protein-enclosed pore large enough to permit the permeation of ions but not lipids. Indeed, we found that the single mutation L302A of nhTMEM16 stabilizes such an intermediate conformation, and the assays of lipid scrambling and ion flux by this mutant supported the “alternating pore-cavity” model (Khelashvili et al., 2019). On the other hand, results from modeling and computational simulations of electrophysiology assays of this scramblase (Kostritskii and Machtens, 2021) support a proteolipidic-pore model, which assumes that ions and lipids are translocated through a single protein conformation. We report here the investigation of the dynamic path of potassium ion (K+) translocation through the open groove of nhTMEM16 observed in the long MD simulations, and present the free energy profile of the ion movement through the groove calculated with umbrella sampling methodology (Torrie and Valleau, 1977). Notably, as discussed further below, we did not observe any instance of chloride passing through the groove. To investigate K+ ion permeation in a model with a groove closed to lipid permeation, the same calculations are performed with the MD simulation trajectories of the nhTMEM16 L302A mutant which exhibits a stable conformation of the groove that does not permit lipid scrambling (Khelashvili et al., 2019). Our results quantify the energetic feasibility of K+ permeation through different conformations of the groove and the slowing down of the process when membrane lipids occupy the groove.

Results

The translocation pathway of K+ through WT nhTMEM16 includes specific interaction sites in the groove

We have previously reported unbiased 10 µs all-atom MD simulations of nhTMEM16 [PDBID 4WIS (Brunner et al., 2014)] in 3:1 POPE/POPG membrane system immersed in a solution containing 150 mM K+Cl− (Khelashvili et al., 2019) (see Material and methods for details). The trajectory from this MD simulation contains two separate events of ion translocating through the protein groove recorded in Supplementary Movies S1, S2. In each case, the translocated ion was a potassium ion (K+). Snapshots representing various stages of the ion translocation during the two events are shown in Figure 2.

The K+ ion permeation occurring at 5.8 µs time-point in the MD trajectory shown in Supplementary Movie S1 is from the extracellular (EC) to the intracellular (IC) end. In the other recorded translocation event – at 7 µs in the trajectory (Supplementary Movie S2), a K+ traverses the groove in the IC-to-EC direction. The duration of the permeation events was 11 ns for the first and 63 ns for the second one (see panels A and B in Figure 3). As can be seen from these snapshots and the Supplementary Movies, the translocated K+ ion forms interactions at two main sites in the groove during both events. These are 1) the EC gate (Lee et al., 2018) formed by the E313-R432 residue pair, and 2) a POPG lipid situated in the middle of the groove and coordinated by several residues (Supplementary Figure S1). Specifically, during the EC-to-IC permeation event, the EC gate is broken (Figure 3G) and the incoming K+ ion partitions at E313 (Figure 3C), which is disengaged from R432. After spending 3 ns at E313 (Figure 3C), the K+ ion continues down the groove and engages in electrostatic interactions with the headgroup of a POPG lipid (Figure 3E) that was seen to penetrate the groove from the IC side at earlier time-point in the trajectory (Supplementary Figure S2). From this interaction site the ion continues to travel down the groove and diffuses away into the solution.

As observed also for the scrambling events recorded in our MD simulations (Lee et al., 2018), the K+ ion translocation pathway in the IC-to-EC direction mirrors the one described above, but in reverse order. Specifically, the K+ ion engages first with the same POPG lipid in the groove as described above for the EC-to-IC event (Figure 3F), and then proceeds towards the EC gate which opens transiently as K+ remains engaged with E313 (Figures 3D,H). After interacting with E313 for 58 ns (Figure 3D), the ion diffuses into the EC milieu completing the translocation (Figure 3B). During this time, a network of interactions was observed among four sites: the potassium ion, E313, R432 and another POPG in the EC region (Figure 2B).

Together, the simulation results suggest a mechanistic role for the electrostatic interactions between K+ ions and the two sites along the groove (the E313 residue, and the groove-bound POPG lipid), akin to the ion-binding sites in ion channels that are considered to serve as “stepping stones” for the ions to pass through the conducting pore (Doyle et al., 1998; Bethel and Grabe, 2016). The involvement of the groove-bound POPG lipid in the permeation sequence is particularly intriguing, since it has been debated whether lipids and ions share the same translocation pathway in the TMEM16 scrambles and thus can be translocated concomitantly, or the lipid scrambling and ion permeation events use different pathways and thus are mechanistically decoupled (Kalienkova et al., 2019; Khelashvili et al., 2019). We find that the POPG lipid that associates with the translocated K+ ion was not scrambled itself (Supplementary Figure S2), but its strategic location in the middle of the groove region and its involvement in the ion translocation in both directions, suggest that the negative charge of its headgroup may lower the free energy barrier for the K+ ion permeation process. To test this hypothesis, we used umbrella sampling simulations (Torrie and Valleau, 1977) to calculate the free energy of K+ ion translocation in the presence or absence of the groove-bound POPG lipid, as well as under conditions when the negatively charged PG headgroup of the bound lipid was substituted by a zwitterionic PC headgroup. The
umbrella sampling simulation of wild type nhTMEM16 were done under four different conditions (see Methods): System 1—the original system in which the two K⁺ permeation events were observed, consisting of the scramblase with the POPG lipid in the groove, embedded in a 3:1 POPE/POPG membrane; System 2—the same a System 1 but without the in-groove POPG; System 3—a system consisting of the scramblase with a POPC in the groove (instead of POPG), embedded in a POPC membrane; System 4—the same a System 3 but without the in-groove POPC. The headgroup of the in-groove POPC in System 3 is at the same Z position along the membrane axis as the head group of the in-groove POPG of System 1 (see Supplementary Figure S3). System 2 and System 4 are without lipid in the groove, but they differ in the membrane composition as detailed above.

The potential of mean force profile for the translocation of K⁺ in the open groove of nhTMEM16

The PMF in each of the four Systems was calculated from the umbrella sampling simulations along the reaction coordinate taken as the distance in the Z direction (perpendicular to the membrane) between the K⁺ ion and the Cα atom of residue T381...
FIGURE 3
Dynamic changes during the K⁺ ion translocation events in the 10 μs unbiased MD simulation of nhTMEM16. In all panels, red arrows indicate the beginning and end of the translocation process. (A,B) Changes in the Z position of the translocated K⁺ ion during the EC-to-IC (A) and IC-to-EC (B) permeation events. The beginning of the EC-to-IC process (t = 0) was defined by the time-point in the trajectory when the ion partitioned from the bulk into the groove (established interaction with E313), while the end of the IC-to-EC process was defined by the time-point when the ion disengaged from E313 and diffused away to the EC milieu (see Supplementary Movies). Z = 0 Å represents the position of the Cα atom of T381. (C,D) Distance between the translocated K⁺ ion and E313. (E,F) Distance between the translocated K⁺ ion and the closest headgroup P atom of the in-
The results of our analyses suggest that the free energy barrier for the permeation of the K+ ion is dependent on the presence of lipids within the groove.

**TABLE 1** Illustration of frame-counting results for the 3Å proximity to K⁺ in an umbrella sampling window (the center of the window $Z_0$ is −1.5 Å away in Z direction from the Cα atom of residue T381) in System 1. The total number of frames in the PMF windows is 4500 and an interaction with the potassium ion is considered “critical” if it is present in >2250 frames (i.e., >50%). Note, however that the total number of interactions is >4500 because in a single frame, the K⁺ ion can interact with several residues. Thus, the sum of percentage in the second column is not 100%.

| Residue | Number of “interacting” frames |
|---------|-------------------------------|
| POPG 909 | 3732 (82.9%) |
| THR 381 | 953 (21.2%) |
| VAL 337 | 135 (3.0%) |
| SER 382 | 53 (1.2%) |
| TYR 513 | 37 (0.8%) |
| THR 340 | 18 (0.4%) |
| LEU 336 | 9 (0.2%) |
| PHE 440 | 2 (0.04%) |

*Note that the total number of interactions is >4500 because in a single frame, the K⁺ ion can interact with several residues. Thus, the sum of percentage in the second column is not 100%.*
indeed the presence of POPC in the groove results in the free energy barrier that is ~5 kcal/mol higher than in the other three systems.

Interestingly, the interval of the K+/POPC interactions in System 3 is longer than for the K+/POPG interactions (from Z = −3.25 Å to Z = 1.25 Å) in System 1. This difference could be due to the smaller size of the PG head group compared to PC. Notably, this short K+/POPG interacting range does not include the local minimum at Z = 2.7 Å in the PMF profile for System 1. Rather, this minimum appears to be due to the combined longer range electrostatic interaction as well as a favorable environment generated by the polar residues T381 and S382 (see Supplementary Figure S4) located just above the POPG lipid binding site (Supplementary Figure S8).

A narrowing of the groove leads to higher free energy barriers for K⁺ ion translocation in the L302A mutant of nhTMEM16

We recently described the L302A mutant of nhTMEM16 in which the groove is significantly narrower, which abrogates lipid scrambling while retaining ion channel activity (Khelashvili et al., 2019). The much narrower opening of the groove region is seen to generate a protein-enclosed pore that can sustain the translocation of certain ions but not lipids. To determine the effect of this structural change on the energetics of K⁺ ion permeation we used the same umbrella sampling protocol to obtain the PMF profile for a K⁺ ion permeation through the nhTMEM16 L302A mutant in a 3:1 POPE/POPG membrane system. The groove of the mutated scramblase remains in its original conformation during the umbrella sampling simulation (Supplementary Figure S9) and there is no lipid in the groove.

Figure 5 shows the PMF profile of K⁺ permeation in the L302A system with the ranges of “critical interactions” marked by the double arrows. The free energy barrier of the K⁺ ion permeation for this construct is ~8.0 kcal/mol, similar to the value calculated for System 3 in which a POPC lipid resided in the open groove of the wild type nhTMEM16. The maximum is at Z = 3.5 Å, where the structure shows the pore to be narrowest, and the translocated K⁺ ion encounters the hydrophobic residue L336, which dynamically occludes the pathway (Supplementary Figure S10). The EC gate in the simulation of this construct is mostly closed as E313 interacts with R432 (Supplementary Figure S11). Nonetheless, E313 still acts as an attracting site for the K⁺ ion, as indicated by the location of the minimum in the PMF curve near the range of the critical interaction with (Figure 5). We note that a POPG at the EC gate interacts with R432 and the K⁺ ion (Supplementary Figure S12), as observed also in the unbiased simulation of wild type.

Electrostatic properties of the nhTMEM16 groove play mechanistic role in regulating energetics of the K⁺ ion permeation process

The space sampled by the K⁺ ion as it progresses along the permeation path is summarized in Figure 6 where the purple spheres indicate positions visited by the potassium ion in all the frames from the equilibrated umbrella sampling simulations. For System 1, the groove volume sampled by the potassium ion does not enclose the whole POPG head group, as the ion passes the in-groove lipid head group “on the side” (see details in the “zoomed-in” insert), without being blocked by it. However, for System 3, the head group of POPC is immersed in the volume of sampling ions, which supports the inference from the corresponding PMF profile that this lipid hinders the potassium ion permeation. Overall, the space sampling is very similar in all wild type nhTMEM16 constructs with open-groove conformations, but different at the narrow “neck” of the groove in the L302A where the side chains of Y439 and L336 form a constriction (see Supplementary Figure S10).

The “average trajectory” paths for the K⁺ ion translocation in the grooves of the various nhTMEM16 constructs (shown in Figures 7A,B) were obtained by averaging the XY coordinates of the positions sampled by the K⁺ ion (purple spheres in Figure 6) at a particular Z. These trajectories are seen to be closely aligned with the groove axes obtained with the HOLE software (Smart et al., 1996) (http://www.holeprogram.org/) which considers only geometric measures of the molecular components (i.e., atom radii). The paths of the translocated ion shown in Figures 7A,B
are seen to follow the groove axis enabling a comparison of the steric environment in the WT and L302A systems using HOLE (Smart et al., 1996) (see Figures 7C,D) and of the electrostatic potential (EP) along the "averaged trajectories" (Figure 8). The values of EP along the trajectories in the four WT constructs and the L302A mutant of nhTMEM16 (Figure 8) were calculated by solving the linearized Poisson-Boltzmann equation (Im et al., 1998) in CHARMM (Brooks et al., 2009) (see Material and methods for details) for only one static frame of the protein (the starting configuration). The results showing that the EP in the nhTMEM16 groove region can be positive or negative in various regions of the different constructs are consistent with the experimental findings that their ion channel activity is not selective for cations or anions (Lee et al., 2016). The generally slightly negative value of the electrostatic potential in Systems 2 and 4, is enhanced by the presence of the POPG lipid in the groove (System 1), consonant with the relatively low free energy barriers calculated for the potassium ion permeation. In contrast, the presence of a POPC lipid in the groove of the wild type nhTMEM16 (System 3), and the narrowing of the groove in the L302A mutant, result in an overall positive EP, which is further consistent with our finding relatively high free energy barriers for the potassium ion translocation in these systems (~7–8 kcal/mol).
Discussion

The permeation events observed from our conventional MD simulations, with no voltage applied, show that potassium ions can pass through the hydrophilic groove of the nhTMEM16 scramblase, consistent with the reports that it can function as a non-selective ion channel (Lee et al., 2016). But the translocation efficiency is dependent on the presence of lipids in the groove, as suggested by the potential of mean force calculations for the permeation paths under various conditions. When the protein is in the Ca\textsuperscript{2+}-activated state, the presence of lipids that are being scrambled or are diffusing in and out of the groove as observed in simulations, can reduce potassium ion permeation to various extents. To interpret the mechanistic reasons for the calculated free energy profile for the translocation, we identified the effects of specific residues involved in K\textsuperscript{+} permeation along the ion path. These include multiple polar residues, such as E313 (in the EC gate region (Lee et al., 2018)), T333, T381, and S382 (see Figure 1B), as well as the in-groove POPG lipid—all of which generate favorable conditions for the ions to pass through the open groove conformation of nhTMEM16. In contrast, the presence of electrostatically neutral POPC lipid in the groove creates a free energy barrier to ion movement, which is ~5 kcal/mol higher than the in-groove POPG.

To consider the role of the groove 3D topography we carried out parallel calculations for the nhTMEM16 mutant L302A, in which groove occlusions prevent lipid scrambling even in the presence of Ca\textsuperscript{2+} (Khelashvili et al., 2019). In the L302A mutant, the free energy barrier for the K\textsuperscript{+} translocation is found to be 6 kcal/mol higher than in the WT construct, and a significantly lower K\textsuperscript{+} permeation under the same condition (Figure 4). While we found that the EP calculated in the grooves of the four WT systems simulated contains both positive and negative patches, indicating that the ion channel is not charge selective, the profile of the electrostatic potential in the permeation path of the L302A mutant is different and suggests that anion permeation may be favored.

In our unbiased 10 µs long MD simulation of the nhTMEM16 wild type in 3:1 POPE/POPG membrane system, we did not observe any instance of chloride passing through the groove in the absence of an applied external voltage across the membrane that is known to modulate ion conductance in TMEM16 scramblases.

**FIGURE 7**
Comparison of the groove axis obtained from the Hole calculations to the “average K+ trajectory” obtained by averaging the XY positions of K+ at each Z value in the umbrella sampling calculations. Panels (A) and (C) are from System 2 calculations of the wild type nhTMEM16 in a 3:1 POPE/POPG membrane. Panels (B) and (D) are from parallel simulations of the nhTMEM16-L302A mutant. The groove axis calculated with the HOLE software is in orange, and the K+ trajectory in purple. For simplicity, only the protein region composed of the TM3–TM7 helices is shown (rendered in teal cartoon), and Ca\textsuperscript{2+} ions are shown as red spheres. Panels (C) and (D) show the radius of the pore along the pathway calculated with the HOLE software as described above.
(Jiang et al., 2017). Notably, the number of Cl\(^-\) ions in the simulation box is much smaller than the number of potassium ions (18 Cl\(^-\) versus 223 K\(^+\)), because the membrane is negatively charged, reducing the probability of a successful Cl\(^-\)/protein encounter. Indeed, our unbiased 15 \(\mu\)s long MD simulation of nhTMEM16 in POPC membrane system (Khelashvili et al., 2019), where the number of Cl\(^-\) is similar to that of K\(^+\) (124 Cl\(^-\) and 130 K\(^+\)), we did observe spontaneous Cl\(^-\) ion penetration into the groove. However, coming from the IC end and moving up only to the middle of the groove, the ion did not complete the translocation in this run, precluding the initiation of a comparative umbrella sampling simulations for Cl\(^-\) similar to those for K\(^+\). In future simulations of much longer times this permeation is expected to be completed and thereby enable full comparative evaluations.

Material and methods

We have previously described in detail the unbiased MD simulation protocols used here for the nhTMEM16 wild type and the L302A mutant (Lee et al., 2018; Khelashvili et al., 2019). Here, the 3:1 POPE/POPG membrane containing the wild type nhTMEM16 was constructed of 851 lipids and a solution box containing 86,011 waters and 150 mM K\(^+\)Cl\(^-\), resulting in a total of 388,447 atoms. The POPC membrane containing the wild type scramble had 679 lipids in a solution box of 90,189 water molecules and 150 mM K\(^+\)Cl\(^-\), for a total of 385,207 atoms. The 3:1 POPE/POPG membrane containing the L302A mutant had 1,505 lipids and 146,737 waters in addition to the 150 mM K\(^+\)Cl\(^-\), resulting in a total of 652,878 atoms. For all simulations we used the latest CHARMM36m force-field for proteins and lipids (Huang et al., 2017), as well as the recently revised CHARMM36 force-field for ions which includes non-bonded fix (NBFIX) parameters (Venable et al., 2013). Umbrella sampling simulations (see below) were performed using NAMD software version 2.13 (Lee et al., 2016), with a 2 fs integration time-step, under the NPT ensemble (at \(T = 310\) K).

Umbrella sampling

System preparation

Before umbrella sampling, Systems 2 and 4 (WT nhTMEM16 with no lipids in the groove) were equilibrated for 11.8 and 6.2 ns after the in-groove lipid was removed from Systems 1 and 3 respectively. Because ion permeation is generally much faster than lipid translocation (104 lipids per second (Suzuki et al., 2010)), 1D umbrella sampling calculations of ion permeation were carried out for all systems with and without a lipid in the groove. The head group of the POPC in the groove of System 3 is at about the same height (Z position along the membrane axis) as the head group of the in-groove POPG of System 1, shown in Supplementary Figure S3. This was achieved by identifying a frame from our earlier unbiased MD simulations of nhTMEM16 in the POPC membrane system, in which there was a POPC lipid at about the same height in the groove,
and the EC gate was broken. This POPC lipid was not scrambled either but diffused towards the middle of the groove from IC, and then moved back to IC. It is worth mentioning that there are always lipids in the IC region of the groove; here a lipid is called “in-groove” if it is situated in the middle of the groove, near or above V447, which is positioned in the middle of TM6 and faces the groove (see Supplementary Figure S8); when we removed the “in-groove” lipid, some lipids in the IC region can move up along the groove, but on the relatively short timescales of the umbrella sampling simulations (2.4 ns), they don’t move high up the groove to be considered as the “in-groove” lipids in the umbrella sampling simulations.

Potential of Mean Force calculations: The Z position of the potassium ion relative to the Z position of the Ca atom of T381 was chosen as the reaction coordinate for the calculation. The residue T381 in TM5 was chosen as it is positioned in the middle of the transmembrane region of the protein (Supplementary Figure S8), and its dynamic fluctuations are relatively small during simulations. We defined 84 umbrella sampling windows covering \( Z = -10.25 \) Å to \( Z = 10.5 \) Å range (\( Z = 0 \) Å representing the position of the Ca atom of T381), with a spacing of 0.25 Å between the centers of neighboring constraints. The bias is a harmonic potential function with a force constant of 20 kcal/mol/Å² to improve the sampling for overlaps of umbrella sampling histograms. For each window, we performed 5 independent runs of 2.4 ns, including 0.6 ns equilibration that was discarded for further analysis. The initial structural configurations for umbrella sampling windows are taken from the unbiased simulation frames where we observed a K⁺ ion in the groove. The potential of mean force was calculated using the WHAM program by setting up a correlation time of 20 ps.

For the mutant nhTMEM16 L302A system, the initial structural configuration is from a frame at the end of a 2 µs unbiased MD simulation of the system. To obtain starting configurations for all the umbrella sampling windows for this system, a steered MD simulation was performed with force constant of 5 kcal/mol/Å² and movement velocity of 10 Å/ns. The setup of the umbrella sampling for the L302A system was the same as the one used for the wild type nhTMEM16, except that we used 96 umbrella sampling windows covering \( Z = -12.25 \) Å to \( Z = 11.5 \) Å range.

Electrostatic potential calculations

The electrostatic potential was calculated by solving the linearized Poisson-Boltzmann equation (Im et al., 1998) in CHARMM (Brooks et al., 2009). The protein was assigned a dielectric constant of \( \varepsilon_p = 2 \). Its TM region was embedded in a membrane slab, which includes a 26-Å-thick hydrophobic core of the bilayer (with \( \varepsilon_{\text{es}} = 2 \) dielectric constant), surrounded by two 8-Å-thick regions, representing the lipid headgroups (\( \varepsilon_{\text{head}} = 30 \)). The membrane slab contained a cylindrical hole (with the dielectric of \( \varepsilon_w = 80 \)) with a radius of 18 Å centered around the groove of the protomer. The system was surrounded by solvent (\( \varepsilon_w = 80 \)) containing 150 mM of monovalent mobile ions. The cylindrical hole was accessible to the ions, but ions were not permitted into the membrane region.

Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

Author contributions

HW and GK supervised the project; XC performed the MD simulations and energy calculations, XC, GK, and HW wrote the manuscript.

Funding

This work was supported by National Institutes of Health (NIH) Grant R01GM106717.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Publisher’s note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

Acknowledgments

We gratefully acknowledge stimulating discussions with Prof. Alessio Accardi and member of his lab. GK and HW acknowledged support from the 1923 Fund. The following computational resources are gratefully acknowledged: the
Anton 2 supercomputer at the Pittsburgh Supercomputing Center (PSC) provided through Grant R01GM116961 from the National Institutes of Health (The Anton 2 machine at PSC was generously made available by D.E. Shaw Research); resources of the Oak Ridge Leadership Computing Facility (INCITE allocation BIP109) at the Oak Ridge National Laboratory, which is supported by the Office of Science of the U.S. Department of Energy under contract no. DE-AC05-00OR22725; and the computational resources of the David A. Coﬁn Center for Biomedical Information in the HRH Prince Alwaleed Bin Talal Bin Abdulaziz Alsaud Institute for Computational Biomedicine at Weill Cornell Medical College.

Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmolb.2022.903972/full#supplementary-material