

Single Channel Kinetics of BK Channels

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Aims

The aim of this presentation is to give a general description and interpretation of what experiments found about the gating process of BK channels.

In particular we consider what types of kinetic gating mechanism can account for the single-channel current records for the activation of BK channels by Ca^{2+} and voltage.

Introduction

Potassium is one of the most important ions involved in the cell physiology. Its concentrations inside and outside the cell membranes, together with the ones of other important ions, strongly determines the electric properties of the cell membrane, such as the resting potential.

The potassium transfer through the cell membrane is regulated by different kind of channels; among these one special mention goes to the BK channels, which are large conductance calcium-activated potassium channels, hence the name, which stands for *big potassium*.

Introduction

Remarkably, the voltage dependent activation of the BK channels creates a negative feedback system to drive the membrane potential more negative, which would then close both the open BK channels and also the voltage dependent Ca^{2+} channels which are often localized in the vicinity.

Through this feedback mechanism BK channels are involved in many physiological processes and their dysfunction can lead to diseases such autism, mental retardation, epilepsy and others.

Structure of a BK channel

We are now looking at a schematic diagram of the modular structure of a BK channel. Only two of the four subunits are shown with the front and back subunits removed.

Bk channels are comprised of four transmembrane voltage sensor domains (VSD) surrounding a central pore gate domain (PGD), and a large cytosolic domain (CTD).

Each VSD is formed by five transmembrane segments, S0 to S4 of a single subunit. The central PGD is formed by S5 and S6 from each subunit.

Structure of a BK channel

The gating ring is formed by the RCK1 and RCK2 (regulators of conductance) domains of each of the four subunits, forming a ring with a central opening. Here is located the Ca^{2+} sensibility of the channel: binding with Ca^{2+} expands the gating ring, pulling on the RCK1-S6 linkers to activate the channel.

The voltage sensibility of the channel is due to the S4 transmembrane segment, which is elevated by depolarization, pulling on the S4-S5 linkers to activate the channels.

DSM models

DSM, which stands for discrete state Markov, form the class of models mostly used to describe gating of BK channels.

They are based upon three assumptions:

- ▶ Short transition times
- ▶ Rate constants invariant for constant experimental condition
- ▶ The gating takes place by means of changing the rate constants

The main idea is that channels gate by moving among conformational states, which have exponentially distributed dwell-times.

If gating concerns multiple states the dwell-time distribution is an exponential distribution with mean dwell time which is the inverse of the sum of the rate constants for the transitions involved; this is also referred as mixed exponentials.

Gating mechanism for BK channels

The theoretical structure of a gating mechanism has to be formulated according to what is known about the microscopic structure of the considered ion channel.

The BK channel structure, as already discussed, is made of a pore gate domain and four voltage sensitive domains plus four pairs of intertwined RCK domains which are properly linked to the PGD to control its gating.

The voltage and Ca^{2+} activation system of BK channels can be incorporated into a kinetic model starting with separate models for each.

A scheme for voltage dependence

Regarding the voltage dependence, the gating mechanism is described by [SCHEME 4]; In this 10-state model, each state is comprised of four subunits, as dictated by the tetrameric structure of BK channels. The five states in the upper tier are closed and the five states in the lower tier are open, with an open state indicated by an open circle in the middle of the four subunits on the lower tier. A square subunit indicates that the voltage sensor in that subunit is relaxed (deactivated), whereas a circular subunit indicates that the voltage sensor is activated.

It is important to notice that there is not an obligatory coupling between voltage sensor movement and channel opening and closing, as channel can open with 0, 1, 2, 3 and 4 activated voltage sensors, with opening probability increasing with the number of activated voltage sensors.

A scheme for Ca^{2+} dependence

The gating mechanism for the Ca^{2+} dependent gating of BK channels is presented in [SCHEME 5]

This scheme parallels the voltage activation shown before except that the activation is by binding Ca^{2+} , indicated by a shaded subunit.

As in the previous case, there is no obligatory coupling between the Ca^{2+} binding and channel opening.

Total scheme

The combination of the voltage and the Ca^{2+} dependence is shown in [SCHEME 6]. [SCHEME 6] includes five repetitions of [SCHEME 4] with 0, 1, 2, 3, and 4 bound Ca^{2+} per channel. The scheme shows that with one Ca^{2+} sensor and one voltage sensor for each subunit the channel could potentially enter 25 closed states on the upper tier and 25 open states on the lower tier during gating, each defined by a different combination of activated voltage sensors and bound Ca^{2+} sites.

Effective number of visited states

The figure [figure 6] presents the single channel recordings obtained from a single BK channel at three different Ca^{2+} concentrations. The Ca^{2+} induced increase in channel activity is readily apparent. Measuring the open and closed interval durations from long records of stable data and plotting them as frequency histograms on log-log coordinates we get the open and closed dwell-time distribution for a fixed Ca^{2+} .

Effective number of visited states

The open distributions are well described by the sum of three significant exponential components and the closed distribution by the sum of five significant exponential components.

This is a rather general result, as open distributions are typically described by 3-4 significant exponential components and closed distribution are typically described by 4-6, suggesting gating in a minimum of 3-4 open states and 4-6 closed states.

Then why isn't the BK gating over 25 open and 25 closed states? One reason is that the 50 states are potential states based on the modular structure of the BK channels. For any given Ca^{2+} and voltage, gating would be effectively limited to only a small subset of 50 states, so only one subset would generate exponentials with sufficiently large areas to detect.

Effective number of visited states

Furthermore, calculations on large multistate models show that many of the expected time constants for 50-state models can be nearly identical or too close to detect.

Activation of BK channels by Ca^{2+}

In [FIGURE 6a] we see the activation of a BK channel at the single-channel level by Ca^{2+} , measured at a fixed +30 mV potential. The dependence of the opening probability on the Ca^{2+} concentration is steep.

Increasing the Ca^{2+} from 0.1 to 1000 μM increased the mean open interval duration 10 times and decreased mean closed interval duration about 1000 times. Hence the major action of Ca^{2+} is an increase in the effective rate constant for opening (leaving closed states), indicating destabilization of the closed states with a much smaller decrease of the closing rate constant (leaving open states), which gives a much smaller stabilization of open states.

Activation of BK channels by voltage

The activation of a single BK channel by voltage is shown in [FIGURE 9a] for voltages ranging from -70 to +100 mV with fixed 95 μM Ca^{2+} .

Changing the voltage from -100 to +100 mV decreased the mean observed closed time 4500 times, while increasing the observed mean closed time of 12 times. Hence the predominant action of depolarization is to destabilize the closed states, with a much smaller action on stabilizing the open states.

Synergistic activation of BK channels by voltage and Ca^{2+}

The joint activation of a BK channel by voltage and Ca^{2+} is shown in [FIGURE 11].

Jointly increasing of Ca^{2+} and depolarization led to synergistic increases in the opening probability.

Increasing Ca^{2+} gives approximately parallel shifts in the opening probability vs. voltage curves, suggesting independent activation by Ca^{2+} and voltage, consistently with the modular model discussed before.

Does the model describe the data?

The consistency of the 50-state scheme with the synergistic activation by voltage and Ca^{2+} was inferred after testing it on a model with limited number states, obtaining good to excellent description of the data.

The key assumption in this procedure is that if the reduced scheme can account for the single-channel kinetics, then the full 50-state scheme is also able to account for the kinetics of the system because it contains the reduced model.

The reduced model lowers the number of constant in such a way that they can be determined through simultaneous fitting of data obtained over a range of Ca^{2+} and voltage.

On the action of the gating ring

To explore the action of the gating ring and the RCK1-S6 linkers, experiments on mutated BK channels have been made. In particular the mutations involved the absence of the gating ring and the length of the RCK1-S6 linkers.

It has been shown that the linkers passively pull on the PGD to bias the channel towards opening in absence of Ca^{2+} , while they apply an active opening force in presence of Ca^{2+} .

Conclusion

The Ca^{2+} dependent ring and the voltage dependent ring can each be approximated by a two-tiered 10-state models with five closed states on the upper tier and five open states in the lower tier.

The joint activation by Ca^{2+} and voltage is obtained with five repeats of the 10-state activation mechanism, where each repeat has 0 to 4 bound Ca^{2+} , or alternatively with five repeats of the 10-states Ca^{2+} activation mechanism, where each repeat has 0 to 4 activated VSD.

Ca^{2+} and depolarization activate BK channels by predominantly increasing the rate of channel opening, or destabilizing the closed states.

To a first approximation, voltage and Ca^{2+} can act independently to activate the channel.

The possibility of interaction among the various sensors arises considering that both VSDs and Ca^{2+} sensors are attached to the PGD, then they could interact allosterically through it.

Conclusion

A preliminary study has shown that a highly constrained 50-state model can approximate the single-channel data. A viable model would describe both the single channel kinetics and the kinetics of the macro current data.

Future studies need to expand the BK models to take into account the two high affinity Ca^{2+} binding sites rather than the usual assumption of one high affinity site. This would shift the 50-state model to a 250-state model, with increased computational complexity.

Altro?