

# Intracellular Calcium Release Channels Mediate Their Own Countercurrent: The Ryanodine Receptor Case Study

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**ABSTRACT** Intracellular calcium release channels like ryanodine receptors (RyRs) and inositol trisphosphate receptors (IP<sub>3</sub>Rs) mediate large Ca<sup>2+</sup> release events from Ca<sup>2+</sup> storage organelles lasting >5 ms. To have such long-lasting Ca<sup>2+</sup> efflux, a countercurrent of other ions is necessary to prevent the membrane potential from becoming the Ca<sup>2+</sup> Nernst potential in <1 ms. A recent model of ion permeation through a single, open RyR channel is used here to show that the vast majority of this countercurrent is conducted by the RyR itself. Consequently, changes in membrane potential are minimized locally and instantly, assuring maintenance of a Ca<sup>2+</sup>-driving force. This RyR autountercurrent is possible because of the poor Ca<sup>2+</sup> selectivity and high conductance for both monovalent and divalent cations of these channels. The model shows that, under physiological conditions, the autountercurrent clamps the membrane potential near 0 mV within ~150 μs. Consistent with experiments, the model shows how RyR unit Ca<sup>2+</sup> current is defined by luminal [Ca<sup>2+</sup>], permeable ion composition and concentration, and pore selectivity and conductance. This very likely is true of the highly homologous pore of the IP<sub>3</sub>R channel.

## INTRODUCTION

Intracellular Ca<sup>2+</sup> signaling is associated with many cellular phenomena. The Ca<sup>2+</sup> signals are generated either by Ca<sup>2+</sup> entry through the surface membrane or by Ca<sup>2+</sup> release from intracellular Ca<sup>2+</sup> stores like the sarcoplasmic reticulum (SR) or endoplasmic reticulum. Surface membrane Ca<sup>2+</sup> entry is mediated by the L-type calcium channel and/or its homologs (e.g., T-, N-, and P/Q-type calcium channels), while Ca<sup>2+</sup> entry from intracellular stores is generally mediated by either ryanodine receptor (RyR) or inositol trisphosphate receptors (IP<sub>3</sub>R) channels.

The RyR and IP<sub>3</sub>R channels share significant homology (1,2), but have little homology with the L-type channel (with the possible exception of their selectivity filters (3–7)). Their single-channel permeation properties reflect this dichotomy: the RyR and IP<sub>3</sub>R channels have high conductance and low (millimolar) Ca<sup>2+</sup> affinity (8–11) while the L-type channel has relatively low conductance and high (micromolar) Ca<sup>2+</sup> affinity (12,13). In this article, we show that the high conductance and low Ca<sup>2+</sup> affinity of the intracellular calcium channels make them ideal for their physiological role of conducting a large Ca<sup>2+</sup> flux over a long time (>5 ms). Our test case here is the RyR channel of striated muscles.

The RyR channel of striated muscle is found in the SR membrane and can mediate large SR Ca<sup>2+</sup> release events lasting >10 ms. Such long release events would not be possible if only Ca<sup>2+</sup> moved across the SR membrane; the rapid movement solely of Ca<sup>2+</sup> would quickly bring the SR membrane potential to the Ca<sup>2+</sup> Nernst potential ( $E_{Ca}$ , the Ca<sup>2+</sup> equilibrium potential), stopping Ca<sup>2+</sup> release. Some other ion

species like K<sup>+</sup>, Mg<sup>2+</sup>, or Cl<sup>-</sup> must provide a countercurrent to prevent the SR membrane potential from coming close to  $E_{Ca}$ .

The need for countercurrent in Ca<sup>2+</sup> release—known for many years (14–20)—can be illustrated using an equivalent circuit model of a patch of membrane that contains conduction pathways for two ion species (e.g., K<sup>+</sup> and Ca<sup>2+</sup>). These pathways could be either two separate channels or through the same channel (Fig. 1 A). At steady state (with conduction pathways open), the net ionic current is 0 and the membrane voltage is (see Eq. 4)

$$V = \frac{g_K E_K + g_{Ca} E_{Ca}}{g_K + g_{Ca}} \quad (1)$$

where  $g_j$  and  $E_j$  are the conductance and Nernst potential, respectively, of ion species  $j$ . Thus, if there is no K<sup>+</sup> current ( $g_K = 0$ ), the Ca<sup>2+</sup> current will stop because the membrane potential  $V$  becomes the Ca<sup>2+</sup> Nernst potential ( $E_{Ca} = -118$  mV, assuming normal resting physiological ion concentrations). On the other hand, if other ions (K<sup>+</sup> in this example) can cross the membrane (assuming equal K<sup>+</sup> concentrations on both sides of the membrane), then any change in the membrane potential generated by the Ca<sup>2+</sup> current will drive K<sup>+</sup> in the opposite direction. This countercurrent attenuates the Ca<sup>2+</sup>-driven change in the membrane potential and will ultimately clamp potential at some value away from  $E_{Ca}$  (Eq. 1). At this potential, there will be a constant driving force for Ca<sup>2+</sup> release. The predicted time course of the membrane potential change in each of these cases is shown in Fig. 1 B. How this affects the unit Ca<sup>2+</sup> current is shown in Fig. 1 C. Equivalent circuit modeling and interpretations like this were presented decades ago (14).

How the SR membrane potential may be affected or generated by Ca<sup>2+</sup> release has been experimentally explored. In

Submitted February 26, 2008, and accepted for publication July 1, 2008.

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Editor: Peter C. Jordan.

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0006-3495/08/10/3706/09 \$2.00

doi: 10.1529/biophysj.108.131987

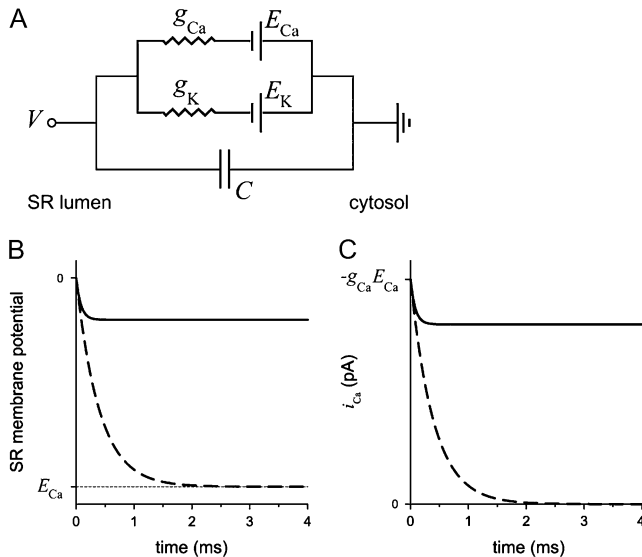


FIGURE 1 (A) Equivalent circuit of Ohmic ion currents across a capacitive membrane. (B) Time course of membrane potential with countercurrent (solid line) and without (dashed line) for the equivalent circuit in panel A. (C) Time course of  $\text{Ca}^{2+}$  current with countercurrent (solid line) and without (dashed line) for the equivalent circuit in panel A.  $g_K = 100$  pS,  $g_{Ca} = 25$  pS,  $E_K = 0$  mV, and  $E_{Ca} = -120$  mV.

1976, Chandler et al. (21) proposed a mechanical model of  $\text{Ca}^{2+}$  release control that did not involve SR potential changes. A few years later, Mathias et al. (22) proposed an alternative model where SR potential changes stimulate  $\text{Ca}^{2+}$  release. Indeed, several studies (using a variety of techniques) have reported signals that were attributed to SR membrane potential changes (23–28). However, the disagreement between these studies has made it difficult to access particular models describing SR membrane potential changes that may be caused by (or generated by)  $\text{Ca}^{2+}$  release. In a now classic work, Somlyo et al. (29) used electron-probe x-ray analysis to monitor total intra-SR ion concentrations before and after  $\text{Ca}^{2+}$  release. They provided compelling evidence against the existence of a large and/or sustained SR membrane potential change, but the possibility of small transient changes in SR membrane potential during  $\text{Ca}^{2+}$  release remains.

Many groups have also attempted to identify the required countercurrent and the channel that mediates it. The SR  $\text{K}^+$  channel has been perhaps most often considered the SR countercurrent pathway (16,26,30–34). However, block of this pathway alters SR  $\text{Ca}^{2+}$  uptake, but does not appear to eliminate SR  $\text{Ca}^{2+}$  release (19,35). Furthermore,  $\text{K}^+$  and  $\text{Cs}^+$  act almost equally well as counterions during prolonged (>10 ms)  $\text{Ca}^{2+}$  release (16,36)—even though  $\text{Cs}^+$  barely goes through the SR  $\text{K}^+$  channel (31,37). This is also true for the briefer  $\text{Ca}^{2+}$  release associated with  $\text{Ca}^{2+}$  sparks in amphibian skeletal muscle (38) and mammalian cardiac muscle (39–41). Overall, the clear implication is that the SR  $\text{K}^+$  channel may not carry the required countercurrent during  $\text{Ca}^{2+}$  release.

Recently, a new SR cation channel called TRIC has been identified (42) and suggested to mediate countercurrent during SR  $\text{Ca}^{2+}$  release. However, TRIC knockout mice have large, fast, caffeine-induced  $\text{Ca}^{2+}$  transients, implying that sufficient countercurrent exists in the absence of TRIC current (42). Countercurrent during SR  $\text{Ca}^{2+}$  release could also be mediated by the SR  $\text{Cl}^-$  channel (43). However, many studies of SR  $\text{Ca}^{2+}$  release in skeletal and cardiac muscle have been done using large impermeable anions like glutamate or aspartate instead of  $\text{Cl}^-$  (14,16,38–41). This suggests that SR  $\text{Cl}^-$  channels also do not carry a substantial countercurrent during  $\text{Ca}^{2+}$  release. Somlyo et al. (29) proposed that  $\text{H}^+$  may also play a role in SR charge compensation. Kamp et al. (44) showed that, although there was significant acidification ( $\sim 0.2$  pH units) inside the SR during  $\text{Ca}^{2+}$  release, proton fluxes could reasonably account for just 5–10% of the necessary charge compensation.

Here, we propose here that the RyR channel mediates its own countercurrent during SR  $\text{Ca}^{2+}$  release for the following reasons:

1. Only countercurrent through RyR is consistent with previous experimental results. RyR conducts  $\text{Cs}^+$  with high conductance and does not conduct anions. Thus, substituting  $\text{Cs}^+$  for  $\text{K}^+$  or use of large anions instead of  $\text{Cl}^-$  would not substantially affect  $\text{Ca}^{2+}$  release.
2. RyR is poorly  $\text{Ca}^{2+}$  selective and has millimolar  $\text{Ca}^{2+}$  affinity (8). Therefore, it is improbable that all the current through RyR is  $\text{Ca}^{2+}$  current under physiological conditions (e.g.,  $\sim 1$  mM intra-SR  $[\text{Ca}^{2+}]$  with symmetric 150 mM  $\text{K}^+$  and 1 mM  $\text{Mg}^{2+}$ ). This is not the case for the L-type calcium channel, which is highly  $\text{Ca}^{2+}$  selective and has 1000-fold greater  $\text{Ca}^{2+}$  affinity. Thus, other permeable cations ( $\text{K}^+$  and  $\text{Mg}^{2+}$ ) very likely move through open RyR channels.
3. A robust model of ion permeation through a single open RyR predicts that there is a substantial  $\text{K}^+$  current under physiological conditions (45,46). The model has reproduced all the known permeation and selectivity data wild-type and mutant RyR channels. It has predicted previously unknown permeability attributes that were later verified experimentally (45,46).

In this article, we use this model (46) and the equivalent circuit approach to compute unit  $\text{Ca}^{2+}$  current,  $\text{K}^+$  and  $\text{Mg}^{2+}$  currents, and SR membrane potential (as well as their time courses) when a RyR channel opens under (approximately) physiological ionic conditions. We conclude that the large  $\text{K}^+$  and  $\text{Mg}^{2+}$  countercurrent through an open RyR is sufficient to clamp the SR membrane potential far from  $E_{Ca}$ . Therefore, RyR channels (and probably  $\text{IP}_3\text{R}$  channels as well) most likely mediate their own countercurrent during  $\text{Ca}^{2+}$  release, while the other SR ion channels likely mediate the countercurrent required for efficient SR  $\text{Ca}^{2+}$  uptake (after RyR channels have closed).

## THEORY AND METHODS

### Equivalent circuits

In our analysis, we consider two variants of the same equivalent circuit (Fig. 1 A, and see Fig. 3 A). The membrane is always modeled as a capacitor with capacitance  $C$ .

In one circuit (Fig. 1 A), two ion species cross the membrane, each with a linear current/voltage relation

$$i_j = g_j(V - E_j), \quad (2)$$

where  $i_j$ ,  $g_j$ , and  $E_j$  are the current, conductance, and Nernst potential of species  $j$ , respectively, and  $V$  is the applied voltage. The ions could be moving through two separate channels or through the same channel. This circuit was used to illustrate the need for countercurrent in the Introduction.

Such a simple equivalent circuit is not, however, a reasonable description of RyR permeability. Specifically, the RyR permeation model (see below) shows that Eq. 2 is not true for  $\text{Ca}^{2+}$  under physiological conditions; the  $\text{Ca}^{2+}$  current through RyR is very nonlinear (approximately exponential) for applied voltages between the SR membrane potential (normally  $\sim 0$  mV) and the  $\text{Ca}^{2+}$  Nernst potential ( $\sim -118$  mV). However, the net conductance through RyR (due to all the ion species) is nearly linear around the reversal potential  $V_{\text{rev}}$  under physiological conditions (Fig. 2 A). Therefore, we model (see *circuit* in Fig. 3 A) net RyR conductance (not the individual ion conductances) as a single resistor with its conductance  $g_{\text{RyR}}$  calculated using the RyR permeation model.

For both circuits, the net unitary current  $i = \sum_j i_j$  and membrane potential  $V$  at time  $t$  is the solution of the equation

$$i(t) = -C \frac{dV}{dt}, \quad (3)$$

namely

$$V(t) = \frac{\sum_j g_j E_j}{\sum_j g_j} + \frac{\sum_j g_j (V(0) - E_j)}{\sum_j g_j} e^{-t/\tau}, \quad (4)$$

$$i(t) = \left( \sum_j g_j (V(0) - E_j) \right) e^{-t/\tau}, \quad (5)$$

where the time constant of the voltage and current change is

$$\tau = \frac{C}{\sum_j g_j}. \quad (6)$$

In all calculations, we assume a  $[\text{Ca}^{2+}]_{\text{cytosol}}$  of  $0.1 \mu\text{M}$  and a membrane capacitance  $C$  of  $0.01$  pF, which approximately corresponds to a  $1 \mu\text{m}^2$  patch of membrane. This is similar to the experimental value of  $0.013$  pF/ $\mu\text{m}^2$  quoted by Baylor et al. (28).

### Model of RyR permeation

We use a model of a single, open RyR pore to compute the net current, as well as the currents carried by each permeant ion species (46). The ions are modeled as charged, hard spheres and their flux through the open pore is described by a combination of one-dimensional Poisson-Nernst-Planck theory and density functional theory of fluids (47,48) (the PNP/DFT model):

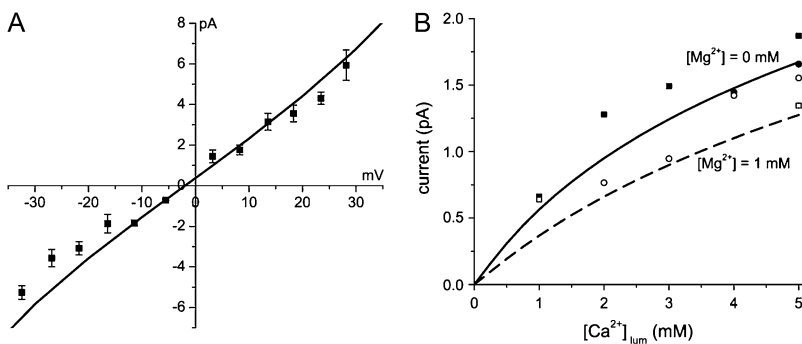
$$-J_j = \frac{1}{kT} D_j(x) A(x) \rho_j(x) \frac{d\mu_j}{dx}, \quad (7)$$

$$-\frac{\epsilon \epsilon_0}{A(x)} \frac{d}{dx} \left( A(x) \frac{d\phi}{dx} \right) = e \sum_j z_j \rho_j(x), \quad (8)$$

where  $\rho_j$  and  $\mu_j$  are the concentration and electrochemical potential, respectively, of ion species  $j$  throughout the pore and baths.  $J_j$  is the flux of ion species  $j$  and  $A(x)$  is the area of the equichemical potential surfaces that is estimated as previously described (49,50). The dielectric constant  $\epsilon$  of the system is 78.4. The value  $\epsilon_0$  is the permittivity of free space,  $k$  is the Boltzmann constant, and  $T = 298.15\text{K}$  is the temperature. The chemical potentials  $\mu_j$  are described with DFT of electrolytes (51,52).

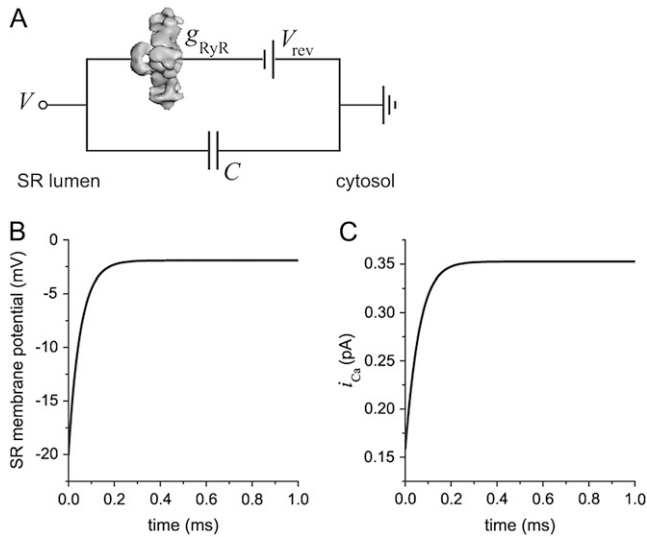
The model of the pore includes the five conserved, charged amino acids found by Gao et al. (5), Wang et al. (6), and Xu et al. (7) to significantly affect RyR selectivity and permeation. These amino acids are Asp-4899, Glu-4900, Asp-4938, Asp-4945, and Glu-4902 in the RyR1 numbering scheme (53). The current/voltage relations ( $-150$  to  $+150$  mV) in over 50 ionic solutions (45,46) (and another 50 solutions as yet unpublished by D. Gillespie, L. Xu, and G. Meissner) of wild-type and mutant RyR channels are reproduced by the model without changing any parameters. The model also predicted—before the confirming experiments were done—anomalous mole fraction effects (54) in mixtures of  $\text{Na}^+$  and  $\text{Cs}^+$  (45),  $\text{Ca}^{2+}$  and  $\text{Na}^+$ , and  $\text{Ca}^{2+}$  and  $\text{Cs}^+$  (46).

This article focuses on ionic solutions that approximate intracellular, physiological conditions at small applied voltages: symmetric  $150$  mM KCl and  $1$  mM  $\text{MgCl}_2$ ,  $1$  mM SR luminal  $\text{CaCl}_2$  and a  $\pm 10$  mV applied voltage range. To show that the model is accurate under these conditions, we compare it—without changing any parameters from those listed in Gillespie (46)—to experimental current/voltage data collected under these conditions (Fig. 2 A). These single RyR channel data were collected with standard methods described elsewhere (55). Note the close correspondence of the experimental results (*squares*) and theoretical prediction (*line*). We also compared the model to previously published experimental results reported by Kettlun et al. (55) in Fig. 2 B (*symbols*). The solid line in Fig. 2 B shows the current predicted by the model in the absence of  $\text{Mg}^{2+}$  ( $0$  mV, symmetric  $150$  mM  $\text{K}^+$ , various luminal  $\text{Ca}^{2+}$  levels). The dashed line in Fig. 2 B shows the current predicted by the model in the presence of  $1$  mM  $\text{Mg}^{2+}$  (again  $0$  mV, symmetric  $150$  mM  $\text{K}^+$ , various luminal  $\text{Ca}^{2+}$  levels). The model reproduces the experimental results very well with no adjustable parameters. These data/



**FIGURE 2** Comparison of the permeation (PNP/DFT) model predictions (*lines*) to experimental data (*symbols*) collected under physiological conditions. (A) Current/voltage curve under our approximate physiological concentrations at  $\sim V_{\text{rev}}$ . Currents are from single RyR2 channels reconstituted in planar lipid bilayers. These data were collected using the methods described by Kettlun et al. (55). The mean  $\pm$  SE are shown and  $n = 4-8$ . Between  $\pm 15$  mV, a linear least-squares fit (not shown) indicated a  $g_{\text{RyR}} = 197$  pS and  $V_{\text{rev}} = -2.3$  mV, which is almost identical to the permeation model curve ( $g_{\text{RyR}} = 192$  pS and  $V_{\text{rev}} = -1.9$  mV). (B) Comparing the model to data published in Kettlun et al. (55). (*Solid line/solid symbols*) The current at  $0$  mV applied voltage with symmetric  $[\text{K}^+] = 150$  mM as luminal  $\text{Ca}^{2+}$  is added. (*Dashed line/open symbols*) The current at  $0$  mV applied voltage with symmetric  $[\text{K}^+] = 150$  mM and  $[\text{Mg}^{2+}] = 1$  mM as luminal  $\text{Ca}^{2+}$  is added. Squares are from mammalian cardiac muscle RyR and circles are from amphibian skeletal muscle RyR.

$\text{Ca}^{2+}$  is added. (*Dashed line/open symbols*) The current at  $0$  mV applied voltage with symmetric  $[\text{K}^+] = 150$  mM and  $[\text{Mg}^{2+}] = 1$  mM as luminal  $\text{Ca}^{2+}$  is added. Squares are from mammalian cardiac muscle RyR and circles are from amphibian skeletal muscle RyR.



**FIGURE 3** (A) Equivalent circuit of net RyR current across a capacitive membrane. The representation of RyR is reproduced from Orlova et al. (72). (B) Time course of SR membrane potential for the equivalent circuit in panel A. (C) Time course of unit Ca<sup>2+</sup> current for the equivalent circuit in panel A. The initial applied potential at time  $t = 0$  is  $-20$  mV.  $g_{\text{RyR}} = 192$  pS and  $V_{\text{rev}} = -1.9$  mV.

theory comparisons and many other published ones (45,46) show that the model can accurately compute the net current, but also—and most importantly for this article—the contributions of each permeable ion species to that net current (see Figs. 5 and 6).

Details of the model have been described previously (46).

## RESULTS AND DISCUSSION

### RyR opening can change local SR membrane potential quickly

To study the time course of local membrane potential due to the opening of a single RyR, we consider a  $1 \mu\text{m}^2$  patch of SR membrane ( $C = 0.01$  pF). In the equivalent circuit shown in Fig. 3 A, the SR membrane potential is Eq. 4,

$$V(t) = V_{\text{rev}} + (V(0) - V_{\text{rev}})e^{-t/\tau}, \quad (9)$$

with time constant  $\tau = C/g_{\text{RyR}}$  where  $g_{\text{RyR}}$  is the net conductance of RyR at the reversal potential  $V_{\text{rev}}$  (Eq. 6).

Our experiments show (Fig. 2 A) that  $g_{\text{RyR}} = 197$  pS gives a time constant  $\tau$  of  $51 \mu\text{s}$ . With this time constant, the SR membrane potential and unit Ca<sup>2+</sup> current reach 95% of their eventual steady-state values in  $\sim 150 \mu\text{s}$ . This is illustrated in Fig. 3, B and C. This time constant is likely an upper limit because a smaller patch of SR membrane or a  $1 \mu\text{m}^2$  patch with more RyR channels would decrease  $\tau$  by decreasing  $C$  or increasing  $g_{\text{RyR}}$ , respectively. This implies that that RyR opening (by itself) will clamp the SR membrane potential to a steady-state value in much less than a millisecond ( $\sim 150 \mu\text{s}$ ). If this steady-state potential is well away from  $E_{\text{Ca}}$ , then an open RyR will continue to conduct Ca<sup>2+</sup> out of the SR. Experimentally, the minimum detectable RyR open event

duration in bilayers is probably near  $358 \mu\text{s}$  (i.e.,  $2 \times 0.179/\text{Fc}$ ; recording system dead time assuming  $\text{Fc} = 1$  kHz (56)). Thus, RyR autocurrent probably clamps the SR membrane potential in less time than the briefest single RyR open events detected in typical planar lipid bilayer studies.

At steady state ( $t > 0.5$  ms), the SR membrane potential becomes the single-channel reversal potential  $V_{\text{rev}}$  (Eq. 9). Here, we compute  $V_{\text{rev}}$  with the model (46), but experimentally measured  $V_{\text{rev}}$  could be used as well.  $V_{\text{rev}}$ —and consequently SR membrane potential—depends on the ionic composition of the cytosol and SR lumen. In the cytosol, the normal physiological divalent concentrations are believed to be  $1$  mM Mg<sup>2+</sup> and  $0.1 \mu\text{M}$  Ca<sup>2+</sup>. Inside the SR (luminal), there is  $1$  mM Mg<sup>2+</sup> and  $1$  mM Ca<sup>2+</sup>. Luminal Ca<sup>2+</sup> concentration  $[\text{Ca}^{2+}]_{\text{lum}}$  (i.e., SR Ca<sup>2+</sup> load) is an important and variable physiological parameter in cardiac muscle and some nonmuscle cells. Also, Mg<sup>2+</sup> concentrations can and have been experimentally manipulated in studies of RyR-mediated SR Ca<sup>2+</sup> release. Therefore, we have evaluated how different  $[\text{Mg}^{2+}]$  and  $[\text{Ca}^{2+}]_{\text{lum}}$  affect  $V_{\text{rev}}$  (and, later, RyR autocurrent).

Fig. 4 A shows how  $V_{\text{rev}}$ , achieved after a RyR channel opens, varies as SR load ( $[\text{Ca}^{2+}]_{\text{lum}}$ ) is increased with symmetric  $1$  mM Mg<sup>2+</sup> present. Fig. 4 B shows  $V_{\text{rev}}$  as a function of symmetric  $[\text{Mg}^{2+}]$  at a constant  $[\text{Ca}^{2+}]_{\text{lum}}$  ( $1$  mM). Note that the  $V_{\text{rev}}$  values here are all between  $0$  and  $-7$  mV and far from  $E_{\text{Ca}}$ . For example,  $V_{\text{rev}}$  is  $\sim -2$  mV and  $E_{\text{Ca}}$  is  $-118$  mV under physiological conditions (at  $1$  mM Ca<sup>2+</sup> in Fig. 4 A). Such small  $V_{\text{rev}}$  values have been measured in single RyR channel studies, even under Ca<sup>2+</sup> and Mg<sup>2+</sup> gradients of  $>5$  mM (46). A reversal potential so far from the Ca<sup>2+</sup> Nernst potential is consistent with the relatively poor Ca<sup>2+</sup> selectivity of the RyR channel and implies that there must be a large countercurrent of other permeable cations through the open channel.

There are, albeit indirect, measurements of the SR membrane potential (e.g., (14,28)). These studies generally have applied a model to convert an optical signal into a predicted SR potential change. Vergara et al. (14) and Baylor et al. (28) used an equivalent circuit model similar to the one we apply here. Both computed that a stationary SR membrane potential is achieved when there is zero net current flowing across the membrane—which is our conclusion as well (Eq. 5). However, they predicted that there is a large SR potential and we predict a small one ( $-2$  mV). The difference in these predictions is generated by the assumptions used to describe ion permeation and selectivity. For example, Vergara et al. (14) described—very reasonable at the time—that SR Ca<sup>2+</sup> currents were Ohmic (i.e., linear). As described above, the RyR-mediated Ca<sup>2+</sup> current is highly nonlinear between  $0$  mV and  $E_{\text{Ca}}$ , and the linear assumption resulted in Ca<sup>2+</sup> current being severely overestimated. Baylor et al. (28), on the other hand, used a Goldman-Hodgkin-Katz model to model individual ion currents. Because their study was performed before RyR permeation properties were characterized, they reasonably

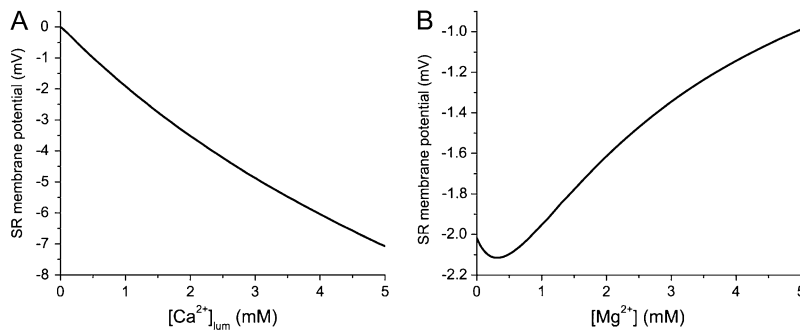


FIGURE 4 Changes in stationary SR membrane potential as divalent concentrations vary, as computed by the PNP/DFT model. (A)  $[Ca^{2+}]_{lum}$  is increased with symmetric  $[K^+] = 150$  mM and  $[Mg^{2+}] = 1$  mM. (B)  $[Mg^{2+}]$  is increased symmetrically as  $[K^+] = 150$  mM and  $[Ca^{2+}]_{lum} = 1$  mM. The minimum in the potential is probably due to an anomalous mole fraction effect similar to the one previously discovered by the model from Gillespie (46).

assumed that the  $Ca^{2+}$  current was conducted by a highly  $Ca^{2+}$  selective channel; the smallest  $Ca^{2+}/K^+$  permeability ratio they considered was 100. We now know that this value is near 7, and if that were used in their formulation, then they would have predicted an SR potential close to  $-2$  mV.

### RyR mediates a large countercurrent

The equivalent circuit model of the capacitive membrane (Fig. 3 A) indicates that the stationary net current through the membrane will be 0 (Eq. 5). If the RyR channel is the only channel present, then the membrane potential is  $V_{rev}$  (Fig. 4) and the net current through the RyR channel will be 0. Since there is a large  $Ca^{2+}$  driving force ( $E_{Ca} - V_{rev} = -118 - (-2) = -116$  mV), there will be a substantial  $Ca^{2+}$  efflux. To have zero net current, there must be an equal countercurrent of other permeable cations,  $K^+$  and  $Mg^{2+}$ . This is illustrated in Fig. 5 where the individual ion currents around the predicted  $V_{rev}$  (Fig. 4) under physiological conditions are shown. Note that the unit  $Ca^{2+}$  current at 0 mV here (0.38 pA) is quite close to the extrapolated unit  $Ca^{2+}$  (0.48 pA) experimentally defined by Kettlun et al. (55) in such salt solutions.

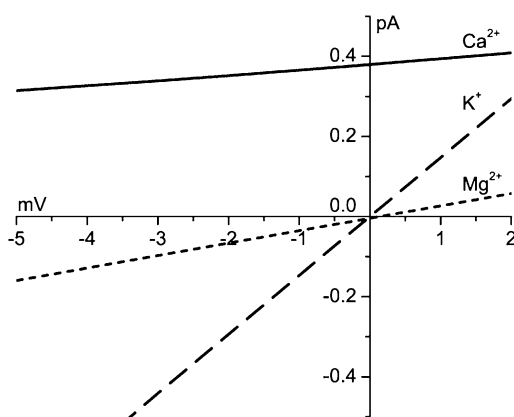


FIGURE 5 Individual  $Ca^{2+}$  (solid line),  $K^+$  (dashed line), and  $Mg^{2+}$  (dotted line) currents at small applied voltages, as computed by the PNP/DFT model. The conductances are 14 pS for  $Ca^{2+}$ , 31 pS for  $Mg^{2+}$ , and 147 pS for  $K^+$ , which were obtained by linear least-squares fits between  $\pm 3$  mV.  $[K^+] = 150$  mM and  $[Mg^{2+}] = 1$  mM symmetric and  $[Ca^{2+}]_{lum} = 1$  mM. The model results for the net current under these conditions are compared to experiments in Fig. 2 A.

Since the ionic concentrations of  $Ca^{2+}$  and  $Mg^{2+}$  in cells may vary (e.g., during exercise or SR overload), we calculated unit currents (at the  $V_{rev}$  of Fig. 4) over a range of  $[Ca^{2+}]_{lum}$  (SR loads) and symmetric  $[Mg^{2+}]$ . Fig. 6 A shows how unit currents vary as SR load ( $[Ca^{2+}]_{lum}$ ) is increased with symmetric 1 mM  $Mg^{2+}$  present. As SR  $Ca^{2+}$  load is increased from 0 to 5 mM,  $K^+$  consistently provides the vast majority (82%) of the countercurrent while  $Mg^{2+}$  provides the rest (18%). Larger  $Ca^{2+}$  loads result in larger  $Ca^{2+}$  currents, but the unit  $Ca^{2+}$  current changes sublinearly with intra-SR  $Ca^{2+}$  levels: doubling intra-SR  $[Ca^{2+}]$  (from 1 to 2 mM) increases unit  $Ca^{2+}$  current by  $\sim 70\%$ ; halving intra-SR  $[Ca^{2+}]$  (to 0.5 mM) reduces the current  $\sim 43\%$ . Some modeling of SR release and sparks (e.g., (57)) assume a linear  $i_{Ca}$ -versus- $[Ca^{2+}]_{lum}$  relationship. The nonlinearity illustrated here could alter the interpretation of those studies.

Fig. 6 B shows how unit  $Ca^{2+}$  currents vary as symmetric  $Mg^{2+}$  is increased at a constant SR load (1 mM). Increasing  $[Mg^{2+}]$  has the opposite effect of increasing SR load: stationary  $Ca^{2+}$  current decreases as  $[Mg^{2+}]$  increases. Also, as  $[Mg^{2+}]$  increases from 0 to 5 mM,  $Mg^{2+}$  provides a larger percentage of the countercurrent, from 0% to 52%. Both of these results are because  $Mg^{2+}$  competes very effectively with  $Ca^{2+}$  for the RyR pore; high  $[Mg^{2+}]$  displaces  $Ca^{2+}$  from the pore, decreasing  $Ca^{2+}$  current and increasing the  $Mg^{2+}$  current (46,54,58–62).

The effect of elevated cytosolic  $Mg^{2+}$  on RyR-mediated SR  $Ca^{2+}$  release has been studied experimentally and our predictions are consistent with those results. For example, elevated cytosolic  $Mg^{2+}$  reduces both the amplitude and spatial width of local  $Ca^{2+}$  release (63). Our calculations suggest that one contributing factor is that the extra  $Mg^{2+}$  reduces the unitary  $Ca^{2+}$  current. Likewise, the marked increase in cytosolic  $Mg^{2+}$  associated with skeletal muscle fatigue (64) will also reduce RyR unitary  $Ca^{2+}$  current, contributing to tension decline.

### Substituting $K^+$ for other monovalent cations

One prediction of the autocountercurrent idea is that any cation conducted by the open RyR channel can provide countercurrent. Fig. 7 (bars) shows that the permeation model predicts that  $Li^+$ ,  $Na^+$ ,  $Rb^+$ , and  $Cs^+$  can all provide

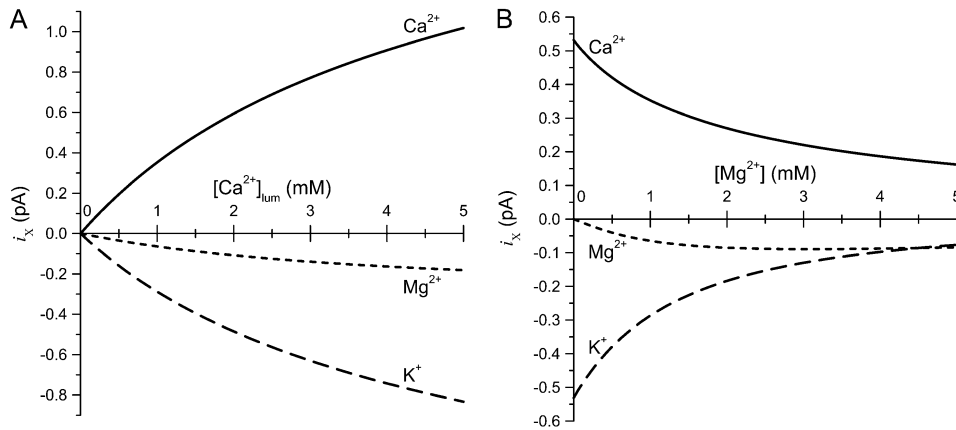


FIGURE 6 (A) Individual  $\text{Ca}^{2+}$  (solid line),  $\text{K}^+$  (dashed line), and  $\text{Mg}^{2+}$  (dotted line) currents as a function of  $[\text{Ca}^{2+}]_{\text{lum}}$  (0 to 5 mM), as computed by the PNP/DFT model.  $[\text{K}^+] = 150$  mM and  $[\text{Mg}^{2+}] = 1$  mM, symmetric. (B) Currents of  $\text{Ca}^{2+}$  (solid line),  $\text{K}^+$  (dashed line), and  $\text{Mg}^{2+}$  (dotted line) as symmetric  $[\text{Mg}^{2+}]$  is increased from 0 to 5 mM, as computed by the PNP/DFT model.  $[\text{K}^+] = 150$  mM, symmetric, and  $[\text{Ca}^{2+}]_{\text{lum}} = 1$  mM. For each  $x$ -axis concentration, the reversal potential is calculated and each ionic species current at that potential is shown. The reversal potential was used because equivalent circuit analysis indicates that this is the

stationary SR membrane potential (Eq. 4). The ionic concentrations and applied potential are very close to the conditions of Fig. 2 B, where the permeation model reproduced experimental results.  $[\text{Ca}^{2+}]_{\text{lum}}$  and  $[\text{Mg}^{2+}]_{\text{lum}}$  are also changed in Gillespie (46), and the model reproduced those results (shown in that article).

substantial countercurrent (relative to that provided by  $\text{K}^+$ ). The predictions shown in Fig. 7 use salt solutions used by Abramcheck and Best (16) in early  $\text{Ca}^{2+}$  release (i.e., initial rate of rise) studies in skinned fiber with different monovalent cations. The Abramcheck and Best (16) results are shown as bold horizontal lines in Fig. 7. Our predictions match very well with the Abramcheck and Best (16) results—except for the  $\text{Li}^+$  case. They reported that  $\text{Ca}^{2+}$  release rate in  $\text{Li}^+$  was reduced to 55%, not 85% as our permeation model predicts.

The skinned fiber studies assume open probability ( $P_o$ ) of RyR is the same in each monovalent cation tested. However, ryanodine binding studies suggest this may not be true. Ryanodine binding is often used to assess RyR channel  $P_o$  since binding is nearly always proportional to  $P_o$  (65). Ryanodine binding (and presumably  $P_o$ ) is substantially suppressed when  $\text{Li}^+$  is present compared to when  $\text{K}^+$ ,  $\text{Na}^+$ , or  $\text{Cs}^+$  are present (66,67). Thus, the reduced  $\text{Ca}^{2+}$  release in skinned fibers observed by Abramcheck and Best (16) likely reflects a  $\text{Li}^+$ -dependent  $P_o$  reduction rather than a reduction of countercurrent effectiveness.

Another prediction of the autocurrent idea is that  $\text{Ca}^{2+}$  release should be severely diminished if permeant counterions are replaced by an impermeant cation. Consistent with this, Abramcheck and Best (16) reported a substantial reduction in  $\text{Ca}^{2+}$  release when  $\text{K}^+$  was replaced by choline. However, they reported that 50% and 100% exchange of  $\text{K}^+$  for choline were equally effective. This is likely explained by slow choline permeation through open RyR channels (68) and occlusion of the pore. Our own pilot measurements of unit  $\text{Ca}^{2+}$  current in the presence and absence of choline are consistent with this notion (data not shown).

### Possible role of other SR channels

If RyR is not the only channel providing countercurrent, then the stationary SR membrane potential is (by Eq. 4)

$$V = \frac{g_{\text{RyR}}V_{\text{rev}} + g_{\text{cc}}V_{\text{cc}}}{g_{\text{RyR}} + g_{\text{cc}}}, \quad (10)$$

where  $g_{\text{cc}}$  and  $V_{\text{cc}}$  are the other countercurrent channel's conductance and reversal potential, respectively. For both  $\text{K}^+$  and  $\text{Cl}^-$ ,  $V_{\text{cc}}$  is 0. The SR membrane potential  $V$  with another countercurrent channel present will be a fraction of  $V_{\text{rev}}$ . Because  $V_{\text{rev}}$  is already close to zero ( $\sim -2$  mV; Fig. 4) with only the RyR present, the contribution of another countercurrent channel is limited to 2 mV. For example, if  $g_{\text{cc}} = g_{\text{RyR}} = \sim 200$  pS, then the presence of  $g_{\text{cc}}$  would change SR membrane potential from  $-1.9$  mV to  $\sim -1$  mV. Even with a large countercurrent carried by another channel, there would still be significant and sufficient RyR auto-

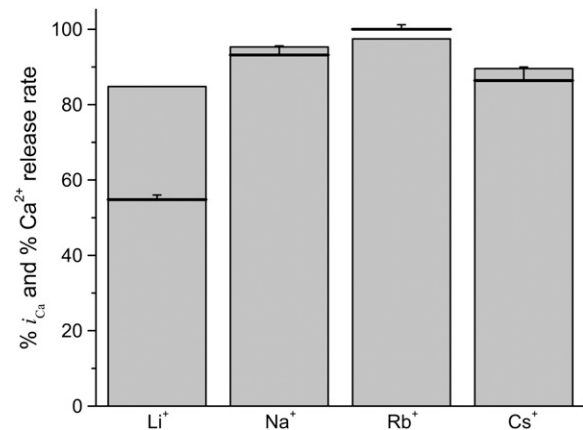


FIGURE 7 Substituting  $\text{K}^+$  for other monovalent cations  $\text{X}^+ = \text{Li}^+$ ,  $\text{Na}^+$ ,  $\text{Rb}^+$ , and  $\text{Cs}^+$ . The shaded bars show the percent  $i_{\text{Ca}}$  compared to when  $\text{K}^+$  is present, as computed by the PNP/DFT model at  $V_{\text{rev}}$ . The solutions contain symmetric 103 mM  $\text{X}^+$ , 7 mM  $\text{Na}^+$ , and 0.1 mM  $\text{Mg}^{2+}$ ;  $[\text{Ca}^{2+}]_{\text{cytosol}} = 0.1 \mu\text{M}$  and  $[\text{Ca}^{2+}]_{\text{lum}} = 1$  mM. These are the same solutions used by Abramcheck and Best (16) for their 100%  $\text{K}^+$  substitution experiments. The Abramcheck and Best (16) measurements (percent of initial rate of rise of  $\text{Ca}^{2+}$  release) are shown as bold horizontal lines with error bars.

countercurrent. That is, there is little gained by having another channel providing countercurrent. For every millivolt of SR membrane potential controlled by another high conductance channel, the  $\text{Ca}^{2+}$  current changes only  $\sim 4\%$  because of calcium's small conductance (see Fig. 5).

If RyR carries its own countercurrent, then what do the SR  $\text{K}^+$  and  $\text{Cl}^-$  channels do? A growing body of work suggests that these channels likely provide the counterion flux necessary for efficient SR  $\text{Ca}^{2+}$  uptake (69) and perhaps carry the countercurrent necessary to return SR potential to resting conditions after the RyR channels close. The SR of striated muscle is divided into two functionally/structurally distinct regions (70,71). The longitudinal SR contains few RyR channels but many SR  $\text{Ca}^{2+}$  pumps (i.e., SERCA). It could be that the countercurrent function of SR  $\text{K}^+$  and  $\text{Cl}^-$  channels is important in the longitudinal SR during  $\text{Ca}^{2+}$  uptake, which is slower and more dispersed than  $\text{Ca}^{2+}$  release. On the other hand, the fast, localized SR  $\text{Ca}^{2+}$  release requires fast, focal countercurrent—exactly what RyR autocountercurrent provides.

## CONCLUSION

We propose that open RyR channels carry their own counterion flux. This follows directly from the equivalent circuit analysis which demonstrates that the stationary SR membrane potential must be between the RyR reversal potential and zero (Eq. 10 for  $V_{\text{cc}} = 0$ ). Since the experimentally defined  $V_{\text{rev}}$  is already very close to zero (46) (see also Figs. 2 A and 4), significant autocountercurrent is inevitable given the poor selectivity of the RyR channel. This general result is independent of the PNP/DFT permeation model used here. Indeed, in many ways the model used here simply plays a pedagogical role by enabling the decomposition of the current into species components (Figs. 5 and 6).

We showed that SR membrane potential will change to  $E_{\text{Ca}}$  in  $< 1$  ms in the absence of countercurrent. Thus, any SR  $\text{Ca}^{2+}$  release event lasting  $> 1$  ms requires a counterion flux be present. This includes SR  $\text{Ca}^{2+}$  release observed in cells during prolonged voltage-clamp depolarizations which last  $> 10$  ms. It very likely includes the release underlying spontaneous  $\text{Ca}^{2+}$  sparks which have rise times lasting  $> 5$  ms. It may even include the very fast and brief  $\text{Ca}^{2+}$  release events stimulated by an action potential in mammalian skeletal muscle (rise times  $\sim 1$  ms). In fact, autocountercurrent may be essential to nearly any RyR-mediated SR  $\text{Ca}^{2+}$  release observed in a cell.

Since  $\text{IP}_3\text{R}$  and RyR channels have homologous pores and permeation characteristics, we predict this may also be true for  $\text{IP}_3\text{R}$ -mediated  $\text{Ca}^{2+}$  release as well. We would also predict that mutations in or near the RyR (and  $\text{IP}_3\text{R}$ ) selectivity filter could be pathogenic if they alter the autocountercurrent process. In wild-type channels, we predict that the early rate of rise of  $\text{Ca}^{2+}$  transients and sparks should depend nonlinearly on ion composition and concentrations.

Lastly, it is possible that TRIC,  $\text{K}^+$ , or  $\text{Cl}^-$  channels may contribute countercurrent during SR  $\text{Ca}^{2+}$  release, but their contribution is relatively small and perhaps not necessary. However, these other ion channels likely carry the required counterion flux during the SR  $\text{Ca}^{2+}$  uptake process.

This work was supported by National Institutes of Health grant No. AR054098.

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