The ryanodine receptor ion channel RyR1 is present in skeletal muscle and has a large cytoplasmic N-terminal domain and smaller C-terminal pore-forming domain comprising six transmembrane helices, a pore helix, and a selectivity filter. The RyR1 S6 pore-lining helix has two conserved glycines, Gly-4934 and Gly-4941, that facilitate RyR1 channel gating by providing S6 flexibility and minimizing amino acid clashes. Here, we report that substitution of Gly-4941 with Asp or Lys results in functional channels as indicated by caffeine-induced Ca\(^{2+}\) release response in HEK293 cells, whereas a low response of the corresponding Gly-4934 variants suggested loss of function. Follow-up mutagenesis and single-channel measurements. Gly-4941 replacement with Lys resulted in channels having reduced K\(^+\) conductance and reduced selectivity for Ca\(^{2+}\) compared with wildtype. RyR1-G4941K did not fully close at nanomolar cytosolic Ca\(^{2+}\) concentrations and nearly fully opened at 2 \(\mu \)M cytosolic or sarcoplasmic reticulum luminal Ca\(^{2+}\), and Ca\(^{2+}\)- and voltage-dependent regulation of RyR1-G4941K mutant channels was demonstrated. Computational methods and single-channel recordings indicated that the open G4941K variant results in the formation of a salt bridge to Asp-4945 in the pore-lining S6 helix impact RyR1 ion permeation and selectivity (13, 14). The RyR1 S6 pore-lining helix has also two conserved glycines, Gly-4934 and Gly-4941, whose replacement with uncharged residues with an increased sidechain volume altered RyR1 channel gating and ion permeation (15). High-resolution cryoelectron microscopy (cryo-EM) using open RyR1 (Fig. 1) and closed RyR1 (16) and cardiac muscle RyR2 isoforms (17) provided structural insights in the mechanisms of channel opening and closing. Binding sites for channel activators Ca\(^{2+}\), ATP, and caffeine were identified (16). It was also shown that the RyR2 pore constriction site of Ile-4868 in the closed state is shifted to Gln-4864 in the open state (17). Ion-pulling simulations that generated an open-channel conformation of RyR1 (18) from the 3.8-Å closed state of RyR1 (7) also showed that the pore constriction site was shifted from Ile-4937 in the closed channel to Gln-4933 in the open channel. The premise of this study was that Gly-4934 and Gly-4941 have critical roles in the ion permeation and Ca\(^{2+}\) regulation of RyR1. We addressed this by replacing the two glycines with two oppositely charged residues, aspartic acid and lysine. The results indicate that RyR1-G4934D, G4934K, and G4941D mutants were not gated by Ca\(^{2+}\), did not conduct Ca\(^{2+}\), and had reduced K\(^+\) conductances in single-channel measurements. In contrast to the closed wildtype (WT) channel, a K\(^{+}\)-conducting RyR1-G4941K mutant channel was recorded at nanomolar Ca\(^{2+}\) concentrations. The mutant channel was
nearly fully opened by SR luminal Ca\(^{2+}\) in a voltage-dependent manner, which suggested that luminal Ca\(^{2+}\) activated RyR1-G4941K by accessing a cytosolic Ca\(^{2+}\)-binding site. Computational methods suggested that RyR1-G4941K formed a salt bridge with Asp-4938. Single-channel measurements of RyR1-G4941K/D4938N and RyR1-G4941K/D4945N mutants validated the computational predictions.

**Results**

**Expression and functional properties of RyR1-Gly-4934 and RyR1-Gly-4941 mutants**

To assess the functional role of two conserved glycines in the S6 pore-lining helix in RyR1 gating and ion permeation, Gly-4934 and Gly-4941 were mutated to negatively charged aspartic acid and positively charged lysine. RyR1 mutant protein expression levels in HEK293 cells ranged from ~50 to 100% of WT (Table 1). In the presence of 10 mM SR luminal Ca\(^{2+}\) and -G4941D. These may have resulted from RyR1-G4934K having a reduced sensitivity to caffeine and loss of high-affinity \(^{3}H\)ryanodine binding but not Ca\(^{2+}\)-dependent channel activity is considered in single-channel measurements described below.

**Single-channel measurements with WT- RyR1 and Gly-4934 and Gly-4941 mutants**

Single channel measurements were performed with purified WT and mutant RyR1s to determine how mutagenesis of Gly-4934 and Gly-4941 affected channel gating and ion permeation. WT and mutant RyR1 channels were solubilized using the zwitterionic detergent CHAPS, purified on sucrose gradients, and reconstituted in proteoliposomes by removing the detergent by dialysis (20). In recording single channels, we took advantage of the finding that RyRs are Ca\(^{2+}\)-gated channels impermeant to Cl\(^{-}\) and conduct K\(^{+}\) more efficiently than Ca\(^{2+}\). The upper left trace in Fig. 3A shows a partially activated single WT-RyR1 channel recorded using 0.25M KCl and 2 mV trace in Fig. 3A, right panel). Reduction from 2 to 0.1 mV allowed the finding of high-affinity \(^{3}H\)ryanodine binding but not Ca\(^{2+}\)-dependent channel activity is considered in single-channel measurements described below.

**Figure 1. Alignment of RyR pore sequences and RyR1 structures**. A, shown are the amino acid sequences of the pore region of rabbit RyR1, RyR2, and RyR3. B, open RyR1 channel structure (PDB code 5tal). C, pore structure of RyR1 (PDB code 5tal). For clarity, B and C show only two of the four RyR1 subunits.
Table 1
Single channel measurements with wildtype and mutant RyR1s

The following abbreviations are used: Ry, ryanodine; ND, not determined. Caffeine response indicates the number of HEK293 cells showing a caffeine-induced Ca\(^{2+}\) response normalized to WT. Single-channel data of mutant channels not conducting Ca\(^{2+}\) were excluded from the statistical analysis. Single-channel open probabilities (P\(_o\)) and K\(^{+}\) conductances were determined with 2 μM Ca\(^{2+}\) in the trans (SR luminal) lipid bilayer chamber. Ca\(^{2+}\) currents were determined at 0 mV.

| Intensity on immunoblots | Caffeine response | B\(_{\text{max}}\) of [\(^{3}H\)]Ry binding | P\(_o\) (2 μM calcium) | P\(_o\) (0.1 μM calcium) | γ\(_K\) | \(I_{\text{Ca}}\) (+10 mM calcium trans) | \(P_{\text{Ca}}/P_{K}\) |
|--------------------------|------------------|---------------------------------|-------------------------|--------------------------|-------|--------------------------------|---------------------|
| WT-RyR1                  | 100              | 100 % WT                        | 100 % WT                | 100 % WT                 |       |                                      |                     |
| G4934D                   | 51.4 ± 12.8(5)   | 2.0 ± 1.4(4)a,b                 | 0 ± 0.6(6)a,b,c         | 0.13 ± 0.04(9)           |       | 820 ± 3(9)                        | −2.3 ± 0.1(5)       |
| G4934K                   | 80.0 ± 24.9(5)   | 2.6 ± 1.3(4)a,b                 | 24.8 ± 6.0(9)b,c         | 0.91 ± 0.05(13)          |       | 519 ± 45(15)                      | −0.1 ± 0.1(11)      |
| G4941D                   | 106.2 ± 28.7(6)  | 72.9 ± 19.2(4)                  | 130.5 ± 14.9(15)b,c      | 0.99 ± 0.10(15)          |       | 304 ± 30(20)                      | −0.1 ± 0.1(10)      |
| G4941K                   | 80.6 ± 187.5(5)  | 63.5 ± 11.2(4)                  | 37.8 ± 9.2(10)b,c        | 0.98 ± 0.04(14)a         |       | 413 ± 79(9)                       | −0.1 ± 0.1(7)       |
| G4941M                   | 115.6 ± 11.2(5)  | 128.3 ± 14.3(8)b,c             | 0.99 ± 0.01(9)a          | 0.13 ± 0.04(15)b,c       |       | 364 ± 5(10)b,c,cd                 | −0.6 ± 0.1(7)       |
| D4938N                   | 84.1 ± 29.3(4)   | 49.9 ± 11.6(5)                  | 0.10 ± 0.01(3)b,c        | 0.13 ± 0.05(4)d          |       | 2.1 ± 0.17(17)                    | 6.0 ± 0.16(6)b,c    |
| G4941K/D4938N            | 97.5 ± 52.1(5)   | 115.3 ± 15.2(4)d,b,c           | 773 ± 3(9)b,c            | 0.10 ± 0.01(3)b,c        |       | 520 ± 6(9)                        | −0.8 ± 0.1(4)       |
| D4941N                   | 61.7 ± 33.6(4)   | 41.7 ± 3.6(5)                   | 264 ± 78(4)b,c          | 0.06 ± 0.02(11)b,c       |       | 3.3 ± 0.24(4)d,c,d,e              | <1 (4)b,c,d,e       |
| G4941K/D4945N            | 48.2 ± 12.6(5)   | 40.9 ± 10.4(4)                  | 0.06 ± 0.01(3)b,c        | 0.06 ± 0.02(11)b,c       |       | 264 ± 78(4)b,c,d,e                | <1 (4)b,c,d,e       |
|                         |                 |                                 |                         |                         |       |                                      |                     |

\(^{a}\) p < 0.05 compared with WT.
\(^{b}\) p < 0.05 compared with G4941K.
\(^{c}\) p < 0.05 as compared with D4938N.
\(^{d}\) Data are from Ref. 14.

Figure 2. Caffeine-induced Ca\(^{2+}\) release in HEK293 cells expressing WT and mutant RyR1s. Membrane potential was −40 mV. Arrowhead indicates the position of caffeine addition to WT and mutant cells. Number of mutant cells showing a caffeine response were summarized in Table 1. Changes in KCL concentration indicated maximal \(P_{\text{Ca}}/P_{K}\).
Luminal RyR1 Ca\textsuperscript{2+} gating

A. WT-RyR1

B. G4934D

C. G4934K

D. G4941D

E. G4941K

F. G4941M

G. D4938N

H. G4941K/D4938N

I. D4945N

J. G4941K/D4945N
conductances for WT-RyR1 and RyR1-G4941K near 600 mosm K⁺ (Fig. 4). Maximal conductances and apparent $K_D$ values were 453 pS and 38 mM for G4941K versus 943 pS and 28 mM for WT. In the presence of 10 mM luminal Ca²⁺, the Ca²⁺ current (−0.6 pA versus −2.3 pA for WT at 0 mV) and permeability ratio of Ca²⁺ over K⁺ ($P_{Ca}/P_K = 2.1$ versus 6.6 for WT) were greatly reduced (Table 1).

Channel open probability of RyR1-G4941K was elevated at 2 μM cytosolic Ca²⁺ (0.89 versus 0.13 for WT) and 0.1 μM Ca²⁺ (0.56 versus 0.01 for WT) (Fig. 3E, Table 1). At 2 μM Ca²⁺ in the bath solution, the RyR1-G4941K mutation increased the duration of open events (34.6 ms versus 0.47 ms for WT) and decreased the duration of the closed events (0.32 ms versus 6.5 ms for WT) (Table 2). The results suggest that RyR1-G4941K exhibited altered Ca²⁺-dependent gating and reduced ion conductances compared with WT.

As a control, RyR1-G4941M was analyzed with methionine having a side-chain volume comparable with lysine. RyR1-G4941M exhibited caffeine-induced Ca²⁺ release in HEK293 cells (Fig. 2) and bound [³H]ryanodine in an in vitro ligand-binding assay, albeit at reduced levels compared with WT (Table 1). Single-channel measurements showed that replacement of a positively charged residue in G4941K with methionine with a comparable side-chain volume but uncharged side chain resulted in channel open probability, $K^+$ conductance, and permeability ratios of Ca²⁺ over K⁺ close to WT (Fig. 3F and Table 1).

Figure 3. Single-channel measurements of WT and mutant RyR1 channels. Left panels show representative single-channel currents at −35 mV (upper and middle traces) or 0 mV (bottom traces) as downward deflections from the closed states (c−). Channels were recorded in symmetrical 250 mM KCl, 20 mM HEPES, pH 7.4, 2 μM Ca²⁺ bath solution (upper traces) and following the subsequent addition of EGTA to yield free cytosolic Ca²⁺ of 0.1 μM (middle traces) or the addition of 10 mM Ca²⁺ to the luminal chamber (bottom traces). Right panels show representative current voltage relationships in 250 mM symmetrical KCl (●) and after the addition of 10 mM luminal Ca²⁺ (○). Averaged $P_o$ values and ion permeation properties are summarized in Table 1.

Figure 4. Conductances as a function of K⁺ activity. A, representative current voltage relations of G4941K at indicated symmetrical KCl concentrations. B, relationships between WT (●) and G4941K (○) conductances and K⁺ activities give $K_o = 28$ mM and $\gamma_{max} = 943$ pS for WT and $K_o = 38$ mM and $\gamma_{max} = 453$ pS for G4941K, using equation $\gamma = \gamma_{max} (1 + K_o/K)$ ⁻¹.

Activation of RyR1-G4941K by 2 μM cytosolic or 2 μM luminal Ca²⁺

Dependence of WT-RyR1 and RyR1-G4941K single-channel activities by 2 μM cytosolic or 2 μM luminal Ca²⁺ was determined at −35 mV membrane potential, which promoted the movement of luminal Ca²⁺ to the cis cytosolic side of the bilayer (Fig. 5). In the presence of 0.01 μM cytosolic and luminal Ca²⁺, WT $P_o$ was essentially 0, and G4941K $P_o$ was 0.13. An increase in luminal or cytosolic Ca²⁺ to 2 μM Ca²⁺ nearly fully activated RyR1-G4941K ($P_o = 0.86$ and 0.98, respectively), whereas WT was incompletely activated by 2 μM cytosolic Ca²⁺ ($P_o = 0.16$). The results indicate that in contrast to WT, (i) RyR1-G4941K had a $P_o > 0.1$ at low cytosolic and luminal Ca²⁺, which indicated Ca²⁺-independent open channel state(s), and (ii) was nearly fully opened ($P_o$ close to 1.0) by 2 μM luminal or 2 μM cytosolic Ca²⁺.

Altered regulation of RyR1-G4941K by cytosolic Ca²⁺ in the presence of 2 μM luminal Ca²⁺

In agreement with rapid ⁴⁰Ca²⁺ flux measurements of SR vesicles (21). WT-RyR1 channels were activated by micromolar cytosolic Ca²⁺ and inhibited by millimolar cytosolic Ca²⁺ at −35 mV (Fig. 6). Ca²⁺-dependent regulation of G4941K differed in several respects from WT. G4941K had elevated open probability at 0.01 μM cytoplasmic Ca²⁺ in the presence of 2 μM luminal Ca²⁺. G4941K was also nearly fully activated at 2 −100
Luminal RyR1 Ca\(^{2+}\) gating

Table 2

Time analysis of wildtype and mutant RyR1s

Single channels were recorded at 2 \(\mu\)M cytosolic and 2 \(\mu\)M luminal Ca\(^{2+}\) were analyzed.

|         | \(P_0\)  | No. of events/min | \(T_o\)  | \(T_c\)  |
|---------|---------|-------------------|---------|---------|
| WT-RyR1 | 0.12 \(\pm\) 0.04(6) | 13,242 \(\pm\) 2776(6) | 0.47 \(\pm\) 0.06(6) | 6.50 \(\pm\) 2.60(6) |
| G4941K  | 0.96 \(\pm\) 0.01(10)\(^{a,b}\) | 7651 \(\pm\) 1828(10) | 34.6 \(\pm\) 17.9(10) | 0.32 \(\pm\) 0.03(10)\(^{a}\) |
| G4941M  | 0.15 \(\pm\) 0.06(7)\(^{c}\) | 8036 \(\pm\) 1676(7) | 0.93 \(\pm\) 0.29(7) | 8.90 \(\pm\) 2.37(7) |
| D4938N  | 0.13 \(\pm\) 0.04(14)\(^{d}\) | 16,829 \(\pm\) 2445(14) | 0.39 \(\pm\) 0.05(14) | 6.80 \(\pm\) 3.30(14)\(^{a}\) |
| G4941K/D4938N | 0.13 \(\pm\) 0.06(4)\(^{e}\) | 4291 \(\pm\) 1193(4) | 1.29 \(\pm\) 0.19(4) | 28.8 \(\pm\) 15.3(4) |
| D4945N  | 0.05 \(\pm\) 0.03(8)\(^{f}\) | 8237 \(\pm\) 4044(8) | 0.31 \(\pm\) 0.02(8) | 71.4 \(\pm\) 34.6(8)\(^{e}\) |
| G4941K/D4945N | 0.99 \(\pm\) 0.01(5)\(^{g,h,i,d}\) | 125 \(\pm\) 76(5)\(^{d}\) | 2091 \(\pm\) 1115(5)\(^{h,i,d}\) | 0.40 \(\pm\) 0.11(5)\(^{h}\) |

\(^{a}\) \(p < 0.05\) compared with WT.
\(^{b}\) \(p < 0.05\) compared with D4945N by analysis of variance.
\(^{c}\) \(p < 0.05\) compared with D4938N.
\(^{d}\) \(p < 0.05\) compared with G4941K.

\(\mu\)M cytosolic Ca\(^{2+}\) and displayed elevated activity at 10 mM cytosolic Ca\(^{2+}\) compared with WT.

RyR1-G4941K regulation by luminal Ca\(^{2+}\)

RyR1 has been reported to be regulated by luminal Ca\(^{2+}\) by binding to luminal channel sites (22) or cytosolic sites following passage of luminal Ca\(^{2+}\) through the open channel (23). To distinguish these mechanisms, dependence of channel open probability on the luminal Ca\(^{2+}\) concentration was probed at −35 and +35 mV (Fig. 7). Cytosolic Ca\(^{2+}\) was 0.01 or 2 \(\mu\)M. In WT at 0.01 cytosolic Ca\(^{2+}\), an increase from 0.01 to 10 mM luminal Ca\(^{2+}\) did not increase the low \(P_o\) at −35 or +35 mV (Fig. 7A). At 2 \(\mu\)M cytosolic Ca\(^{2+}\), an increase of luminal Ca\(^{2+}\) from 0.01 to 10 mM significantly decreased \(P_o\) at −35 mV but not at +35 mV. Regulation of RyR1-G4941K by luminal Ca\(^{2+}\) differed greatly from WT (Fig. 7B). At 2 \(\mu\)M cytosolic Ca\(^{2+}\), RyR1-G4941K was fully activated at both membrane potentials at 0.01 \(\mu\)M to 10 mM luminal Ca\(^{2+}\). At 0.01 \(\mu\)M cytosolic Ca\(^{2+}\), which kept the mutant channel open, luminal Ca\(^{2+}\) was more effective in increasing \(P_o\) at negative than positive membrane potentials. This suggested that luminal Ca\(^{2+}\) flowing through the channel accessed cytosolic Ca\(^{2+}\) activation sites in RyR1-G4941K. This was tested further by recording voltage dependence of RyR1-WT and -G4941K channel activities at 0.01 \(\mu\)M cytosolic Ca\(^{2+}\) and 0.01 or 2 \(\mu\)M luminal Ca\(^{2+}\) (Fig. 8). At 0.01 and 2 \(\mu\)M luminal Ca\(^{2+}\), WT channels were closed, displaying no voltage-dependent activity. At 0.01 \(\mu\)M luminal Ca\(^{2+}\), RyR1-G4941K displayed a weak voltage-dependent \(P_o\) at membrane potentials ranging from −80 to +80 mV. At 2 \(\mu\)M luminal Ca\(^{2+}\), \(P_o\) was higher at negative than positive membrane potentials. Thus, a larger number of Ca\(^{2+}\) ions appeared to flow through the channel at negative than positive membrane potentials to activate RyR1-G4941K to a greater extent at negative potentials.

RyR1-WT and -G4941K activities in the presence of FKBP12 and in membrane isolates

We considered the possibility that elevated RyR1-G4941K channel open probability may result from the absence of FKBP12, a closely associated subunit of RyR1. Absence of FKBP12 in the WT-RyR1 macromolecular complex activates Ca\(^{2+}\) release from the SR and increases channel activity in lipid bilayers (24, 25). Proteoliposomes containing purified WT and RyR1-G4941K channels were therefore treated for 30 min with 5 \(\mu\)M FKBP12 before addition to the cis bilayer chamber. Treatment of WT and RyR1-G4941K with FKBP12 did not significantly affect \(P_o\) at 0.1 and 2 \(\mu\)M Ca\(^{2+}\) compared with channels recorded in the absence of FKBP12, respectively (Fig. 9, A and B). To increase...
the probability of preserving normal function, experiments were also performed with channels that were not subjected to detergent treatment. Fig. 9, A and B, shows that RyR1-G4941K channels obtained using membrane preparations isolated from HEK293 cells maintained an elevated $P_o$ at 0.1 and 2 $\mu$M cytosolic Ca$^{2+}$ compared with WT, respectively. The results indicate that preincubation with FKBP12 and elimination of detergent treatment did not decrease the single-channel activities of RyR1-G4941K to those of WT.

RyR1-G4941K interaction with RyR1-Asp-4938 and Asp-4945

Modeling revealed an interaction of RyR1-G4941K with Asp-4938 (Fig. 10, see under “Discussion”). The computational predictions were tested by performing single-channel measurements with RyR1-G4941K/D4938 and G4941K/D4945 double mutants. Their properties were compared with WT, RyR1-G4941K, and two previously studied single-site mutants (RyR1-D4938N and -D4945N) (Tables 1 and 2) (14). The two single-site (D4938N and D4945N) and double-site (G4941K/D4938 and G4941K/D4945) mutants had cellular caffeine-induced Ca$^{2+}$ release and bound $[^3H]$ryanodine that indicated function was retained (Fig. 2 and Table 1). In single-channel measurements, substitution of Asp-4938 with Asn in RyR1-D4938N and RyR1-G4941K/D4938 and Asp-4945 with Asn in RyR1-G4941K/D4945 reduced K$^+$ conductances and Ca$^{2+}$ currents at 0 mV (Fig. 3 and Table 1). In contrast, RyR1-D4945N maintained K$^+$ conductance and Ca$^{2+}$ current close to WT–RyR1.

RyR1-D4938N, -D4945N, and -G4941K/D4938N had channel open probabilities comparable with WT–RyR1 at 2 $\mu$M cytosolic Ca$^{2+}$ (Fig. 3 and Table 1). In contrast, RyR1-G4941K/D4945N maintained high $P_o$ comparable with RyR1-G4941K ($P_o \sim 1.0$ versus 0.89). Significant differences were also obtained for the open and closed times (Table 2). Compared with the D4938N mutant, the mean open time of RyR1-G4941K/D4938N increased 4-fold ($p < 0.05$, Student’s $t$ test), and compared with G4941K, it decreased $\sim 25$-fold. Mean closed time of RyR1-G4941K/D4938N increased $\sim 4$-fold compared with WT–RyR1 and RyR1-D4938N ($p < 0.05$, Student’s $t$ test). The mean open time of RyR1-G4941K/D4945N increased $\sim 4000$-, $\sim 60$-, and 6,500-fold, respectively, compared with WT, RyR1-
Luminal RyR1 Ca\(^{2+}\) gating

![Graph](image)

Figure 9. Effects of FKBP12 on purified WT and G4941K mutant channel open probabilities and fusing membrane fractions with lipid bilayers. Single open probabilities of WT and G4941K in 250 mM symmetrical KCl were determined at −35 mV. SR luminal Ca\(^{2+}\) was 2 μM. Single-channel open probabilities were at 0.1 μM (A) and 2 μM (B) cytosolic Ca\(^{2+}\) of WT and G4941K, membrane isolates (M), purified WT, and G4941K mutant channels not incubated (P), and incubated with 5 μM FKBP12 for 30 min at 24 °C (P + FKBP). Bars show means of data.

G4941K, and D4945N. The mean closed time of RyR1-G4941K/D4945N decreased 14-fold compared with WT, was comparable with RyR1-G4941K, and decreased ~160-fold compared with RyR1-D4945N. The results suggest that attenuation of the G4941K interaction with Asp-4938 favors the closed state, whereas attenuation of the G4941K-D4945 interaction favors the open state.

Discussion

The principal finding of this study is that RyR1-G4941K exhibits a marked increase in the sensitivity of SR luminal Ca\(^{2+}\) regulation in single-channel recordings. The RyR1-G4941K mutant exhibited a caffeine-induced Ca\(^{2+}\) response in HEK293 cells and bound \(^{3}H\)ryanodine, but it had reduced Ca\(^{2+}\) over K\(^{+}\) permeability ratio compared with WT. This was in contrast to the finding that RyR1-G4934D and -G4934K mutants lost caffeine-induced Ca\(^{2+}\) response in HEK293 cells, had a pronounced decrease in \(^{3}H\)ryanodine binding, and Ca\(^{2+}\) currents decreased to background levels. These findings suggest loss-of-function channels in HEK293 cells, membrane isolates, and single-channel measurements. By comparison, the G4941D mutation resulted in a more complex behavior. RyR1-G4941D responded to caffeine, which suggested the formation of a functional Ca\(^{2+}\)-conducting channel in HEK293 cells. However, \(^{3}H\)ryanodine binding was low, and Ca\(^{2+}\) currents decreased to background levels, which suggested loss of function on removal from HEK293 cells. The mutagenesis results support the modeling data described below that RyR1-G4941K forms a salt bridge to Asp-4938 and luminal Ca\(^{2+}\) accesses a recently identified Ca\(^{2+}\)-binding site in the open-channel conformation.

Our results suggest that the RyR1-G4941K mutant undergoes a structural change during extraction from HEK293 cells. The caffeine-induced Ca\(^{2+}\) release suggested an intracellular Ca\(^{2+}\) store in HEK293 cells in the absence of caffeine. In contrast, single-channel measurements suggested that RyR1-G4941K rapidly released Ca\(^{2+}\) from the intracellular membrane compartment. Neither the potential loss of the FKBP12 subunit from the RyR1 channel complex during purification nor exposure to detergent was found to be responsible for the elevated channel activity in single-channel measurements. The results may be interpreted that the mutant channel underwent a structural change upon removal from the cellular environment.

Previous single-channel studies showed that mutagenesis of the two conserved glycines in the pore-lining S6 pore-lining helix altered RyR function. Substitution of RyR2-Gly-4864 (corresponding to RyR1-Gly-4934) with alanine resulted in no significant changes of RyR2 function, whereas replacement with valine and proline profoundly altered both channel gating and ion permeation (26). Single-channel studies were done using purified RyR2 channel preparations. We reported that replacement of RyR1-Gly-4934 and Gly-4941 with Ala altered RyR1 channel function in single-channel measurements using purified RyR1 preparations (15). A mutation further increasing the side-chain volume at Gly-4934 (G4934V) resulted in loss of function. In contrast, function was maintained when membrane fractions isolated from HEK293 cells were fused with lipid bilayers (18). In this study, loss of function by the G4934D and G4941D mutant channels was seen when proteoliposomes containing the purified mutants or membrane fractions were fused with lipid bilayers.

Aspartic acid has a smaller residue volume than valine (115 Å\(^{3}\) versus 138 Å\(^{3}\) (27)). This suggested that the introduction of additional negative charges near Asp-4938 and Glu-4942 rendered RyR1-G4943D structurally unstable in HEK293 cells and RyR1-G4941D when removed from the cellular environment.

Modeling the RyR1-G4941K mutation in the open-channel conformation revealed the preferential formation of an inter-subunit salt bridge with Asp-4938 (Fig. 10) compared with Asp-4938 versus Asp-4945. Among the four chains the interaction persists for 34% of the simulation, meaning that on average between one and two salt bridges with Asp-4938 are present at any point in the simulation. Single-channel recordings suggested that the formation of a salt bridge stabilizes the open channel state. In the G4941K/D4945N double mutant, we observed an increase in the G4941K and Asp-4938 interaction in the open-channel conformation and in single-channel measurements a prolonged duration of channel openings. As expected, mutating Asp-4938 to Asn attenuated salt-bridge formation with D4938N. G4941K now showed a preference for intrasubunit salt bridge formation with Asp-4945, and single-
channel measurements indicated decreased stability of the open-channel in favor of a closed-channel conformation.

Time analysis of the single-channel data is in agreement with these conclusions. The RyR1-G4941K/D4945N double mutant showed a substantial increase in $T_o$ compared with RyR1-G4941K and D4945N, which indicated increased stability of the open-state or a barrier to return to the closed state. In contrast, the G4941K/D4938N double mutation resulted in an elevated $T_c$ compared with G4941K and D4938N, which implies that the closed state was more stable or that there was an additional barrier to opening. As a control, we modeled the open channel conformation of the G4941M mutation and monitored the distances between G4941M and Asp-4938 or Asp-4945 (Fig. 11). We observed closer contacts of G4941M with Asp-4945; however, neither showed significant contacts below the 3.5 Å range required for salt-bridge formation.

Single-channel measurements showed that replacement of glycine with positively charged lysine reduced $K^+$ conductance and Ca$^{2+}$ selectivity of RyR1-G4941K, G4941K/D4938N, and G4941K/D4945N mutants to a greater extent than G4941M, D4938N, and D4945N mutants. Pore profiles for the open-channel RyR1-G4941K, G4941M, G4941K/D4938N, and G4941K/D4945N mutants are compared with WT in Fig. 12A.

Figure 10. Salt bridges in open mutant channels. Upper panels show representative images of salt bridges in G4941K, G4941K/D4945N, and G4941K/D4938N mutant channel structures. Lysine, aspartate, and asparagine are colored blue, red, and green, respectively. Salt-bridge contacts are shown with dotted lines. In the G4941K mutant, G4941K preferentially forms intersubunit salt bridges with Asp-4938 (observed between two of the four chains). An increase in the G4941K-D4938 interaction in the G4941K/D4945N mutant is observed in three of the four chains. In the G4941K/D4938N double mutant, we observe attenuation of the G4941K-D4938N interaction. A small preference of G4941K to form an intrasubunit salt bridge with Asp-4945 is observed in one of four subunits. The lower panels show distance distributions for the three mutants. Distances were monitored between the terminal amine groups and the oxygen or nitrogen atoms in the carboxyl or amide side chains. Distances were computed among all four chains for 11 structures sampled from MD simulations of each of the three open-channel RyR1 structures (PDB codes 5tv, Stal, and Sta3). Distances less than 3.5 Å are considered as potential salt bridges. For G4941K, we observe a salt bridge preference between G4941K and Asp-4938. When Asp-4945 is mutated to Asn, we observe an increased preference for the G4941K-Asp-4938 salt bridge. As expected, mutating Asp-4938 to Asn attenuates the salt-bridge formation, and G4941K now shows a preference for Asp-4945.

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Figure 11. G4941M contacts in the open channel. A, as a control the conformations of G4941M are shown with respect to Asp-4938, Asp-4945, and the pore. G4941M is observed to move into an intersubunit pocket that is out of the pore and below Asp-4938 explaining contacts between Asp-4938 and G4941M and the increased pore radius of the G4941M mutation compared with G4941K. The two right panels monitor distances between G4941M and Asp-4938 (B) or Asp-4945 (C). We observe more close contacts between G4941M and Asp-4938 than between G4941M and Asp-4945; however, neither show significant contacts below 3.5 Å (the range expected for salt-bridge formation).
and mutants. Changes in the electrostatics of Ca$^{2+}$ lysine in Gly-4941. The G4941K/D4938N double mutation resulted in a small decrease in pore radius compared with G4941K, which was likely due to the decreased formation of intersubunit salt bridges. The decreased formation of intersubunit salt bridges caused G4941K to either protrude into the pore or form an intrasubunit salt bridge with Asp-4945, which caused the β, γ, or δ carbons of lysine to protrude into the pore. RyR1-G4941M had a pore radius of ~4.5 Å. A move out of the pore into an intersubunit pocket may explain the larger pore radius for methionine compared with G4941K.

The data of Fig. 12B and C shows the interaction energies of the pore residues with Ca$^{2+}$ and K$^+$ ions as the two ions move through the pore. The results show that the electrostatic interaction energies of WT varied between −5 and −2 kcal/mol for Ca$^{2+}$ and −2.5 and −1 kcal/mol for K$^+$. A rightward shift from the RyR1-G4941K position was observed for the energy maxima. The negative interaction energies of Ca$^{2+}$ and K$^+$ are likely primarily due to a favorable interaction with negatively charged amino acid residues (RyR1-Asp-4899 and -Glu-4900) immediately after the GGGIG selectivity motif in the luminal vestibule and negatively charged residues in the cytoplasmic vestibule (Fig. 14). Substitution of glycine in RyR1-G4941K with four positively charged lysine increased interaction energies for Ca$^{2+}$ carrying a +2 charge than on K$^+$ carrying a +1 charge. To estimate the electrostatics of the ions within the pore, we calculated the electrostatic Ca$^{2+}$- and K$^+$-protein interaction energies using APBSmem software tool (28, 29). Fig. 12, B and C, illustrates these interaction energies using APBSmem software tool (28, 29).

Recent cryo-EM studies identified a binding site for Ca$^{2+}$ at the interface of the RyR1 CSol and C-terminal domains (16). Earlier studies indicated that mutations at other sites affected Ca$^{2+}$-gated channel activity. Substitution of RyR1-Glu-4032 by alanine and the corresponding amino acid residues in the RyR3 isoform reduced Ca$^{2+}$-gated channel activity (30, 31). RyR1-Glu-4032 is not part of the Ca$^{2+}$-binding site but was suggested to stabilize the interaction between two RyR1 regions by H$^+$ bonding to amide nitrogens (16). Mutations that give rise to central core disease in skeletal muscle exhibit loss of Ca$^{2+}$-dependent channel activity and Ca$^{2+}$ conductance (32–34). Many
of the central core disease mutations are located in the pore-forming region of RyR1. To our knowledge, G4941K mutant has not been associated with a human pathology.

Other sites involved in the regulation of RyR1 by Ca\textsuperscript{2+} include residues in S2 (35), the S4-S5 linker (36, 37), and S6 of the cardiac RyR2 isomorph corresponding to RyR1 amino acid residues Gln-4946 to Met-4954 of RyR1 (38). This study extends these findings and shows that sensitivity of RyR1-G4941K to cytosolic and luminal Ca\textsuperscript{2+} increased compared with WT.

Regulation of the RyRs by luminal Ca\textsuperscript{2+} has been attributed to luminal Ca\textsuperscript{2+}-sensing channel sites (22, 39–41), the binding to cytosolic sites following the passage of Ca\textsuperscript{2+} through the open channel (23, 42, 43), or the involvement of Ca\textsuperscript{2+}-sensing sites on both the luminal and cytosolic channel sites (44). To distinguish between luminal and cytosolic Ca\textsuperscript{2+}-regulatory sites, we previously determined the voltage- and Ca\textsuperscript{2+}-dependent regulation of WT-RyR1 (23). As shown in this study, closed WT-RyR1 channels are not activated by luminal Ca\textsuperscript{2+}, and single channels were partially opened by cytosolic ATP in the presence of a low cytosolic Ca\textsuperscript{2+} concentration. Channels were activated by 50 \(\mu\text{M}\) luminal Ca\textsuperscript{2+} to a greater extent at negative membrane potentials that favored Ca\textsuperscript{2+} flow through the RyR1 pore than at positive membrane potentials that disfavored passage of Ca\textsuperscript{2+} through RyR1. This suggested that luminal Ca\textsuperscript{2+} bound to cytosolic channel sites following their passage through the open channel. A similar result was obtained when the partially open G4941K mutant channel was recorded at a low cytosolic Ca\textsuperscript{2+} concentration. G4941K was activated to a greater extent by 2 \(\mu\text{M}\) luminal Ca\textsuperscript{2+} at negative than positive membrane potentials. These results are best rationalized by luminal Ca\textsuperscript{2+} accessing cytosolic channel sites.

A model consistent with the results of this study suggests that cytosolic Ca\textsuperscript{2+} ions access a recently identified Ca\textsuperscript{2+} activation site located in the cytosolic foot structure of RyR1 (16). The model suggests that cytosolic Ca\textsuperscript{2+} can access the Ca\textsuperscript{2+} activation site when channels are closed or open (Fig. 13). In contrast, luminal Ca\textsuperscript{2+} can access the site in WT only when channels are open. RyR2 channel opening shifts the pore constriction site of Ile-4968 in the closed state to Gln-4864 in the open state (corresponding to Ile-4937 and Gln-4933 in RyR1, respectively) (17). According to the model, luminal Ca\textsuperscript{2+} activates RyR1 when gaining access to the Ca\textsuperscript{2+} activation site following passage through the open-channel RyR1-Gln-4933 restriction site. RyR1-G4941K differs in two respects from WT in its regulation by Ca\textsuperscript{2+}. First, the RyR1-G4941K mutant channel remains open at low cytosolic Ca\textsuperscript{2+} concentrations; second, it exhibits an increased sensitivity to activation by cytosolic and luminal Ca\textsuperscript{2+} compared with WT. Our results suggest that luminal Ca\textsuperscript{2+} accesses Ca\textsuperscript{2+} activation sites as they pass through the pore rather than traveling to openings that lie outside the pore.

**Experimental procedures**

**Materials**

[\textsuperscript{3}H]Ryanodine was obtained from PerkinElmer Life Sciences; protease and phosphatase inhibitors were from Sigma, and phospholipids were from Avanti Polar Lipids.

**Preparation of wildtype and mutant channels**

RyR1 mutations G4934D, G4934K, G4941D, and G4941K were introduced by Pfu polymerase-based chain reaction using mutagenic oligonucleotides and the QuikChange II site-directed mutagenesis kit (Stratagene, La Jolla, CA) (45). RyR1-G4941M, G4941K/D4938N, and G4941K/D4945N were prepared using a gene synthesis method and a proprietary protocol (Genewiz, Inc., South Plainfield, NJ). WT-RyR1 and mutants were transiently expressed in HEK293 cells using FuGENE 6 (Roche Applied Sciences) or jetPRIME (Polyplus, New York) according to the manufacturers’ instructions. Cells were maintained in DMEM/F-12 (1:1) medium containing 10% fetal bovine serum at 37 °C and 5% CO\textsubscript{2} and plated the day before transfection. Following transfection, cells were maintained at 35 or 37 °C and harvested ~70 h after transfection. Crude membrane isolates (45) and proteoliposomes containing purified WT or mutant RyR1 channels were prepared as described (20).

**Cellular Ca\textsuperscript{2+} release**

Stored Ca\textsuperscript{2+} release was determined as described (46). Briefly, HEK293 cells were grown on glass coverslips, and Ca\textsuperscript{2+} transients were monitored with the fluorescence Ca\textsuperscript{2+} indicator Fluo-4. The addition of ~8 mM caffeine induced cellular Ca\textsuperscript{2+} release. Individ-

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**Figure 13. Schematic for cytosolic and luminal Ca\textsuperscript{2+} access to Ca\textsuperscript{2+}-binding site in closed and open RyR1 channels.** Two of the four subunits are shown (PDB code Stal). Closed- and open-channel RyR1 structures are shown in grey and white, respectively. The position of the Ca\textsuperscript{2+}-binding site is shown in magenta. Shown in blue is a cutaway of the structure highlighting a cavity through which Ca\textsuperscript{2+} can access the Ca\textsuperscript{2+}-binding site. In the closed-channel RyR1, Ca\textsuperscript{2+} can only access the Ca\textsuperscript{2+}-binding site from the cytosolic side of the channel. However, in the case of an open-channel RyR1 Ca\textsuperscript{2+} can also reach the Ca\textsuperscript{2+}-activation site by approaching from the luminal side of the channel.
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Rulli cells were monitored using the EasyRatioPro algorithm (Photon Technology International, Lawrenceville, NJ).

**SDS-PAGE and immunoblot analyses**

Proteins in crude membrane fractions (20 μg of protein/lane) were separated using 3–12% gradient SDS-PAGE, transferred overnight to nitrocellulose membranes (37), and probed with primary rabbit anti-RyR1 polyclonal antibody 6425 prepared by ProSci against RyR1 sequence FIKGLDSFSGKPRGSG. Immunoblots were developed using enhanced chemiluminescence and quantified using the Bio-Rad ChemiDoc MP Imaging System and ImageQuantTL analysis software.

[^3]H]Ryanodine binding

Ryanodine binds with high specificity to RyR1 and is widely used to probe RyR activity and content (19).[^3]H]Ryanodine binding was measured by incubating crude membrane isolates for 4–5 h at 24 °C with a near-saturating concentration of 20 nM[^3]H]ryanodine in 20 mM imidazole, pH 7.0, 0.6 mM KCl, protease inhibitors, 1 mM oxidized glutathione (GSSG), and 0.1 mM Ca\textsuperscript{2+}. Non-specific binding was determined using a 100-fold excess of unlabeled ryanodine. Aliquots of samples were diluted 9-fold with ice-cold water and placed on Whatman GF/B filters saturated with 2% polyethyleneimine. Filters were washed with ice-cold 0.1 M KCl, 1 mM imidazole, pH 7.0. Radioactivity remaining on the filters was determined by liquid scintillation counting to obtain bound[^3]H]ryanodine.

**Single-channel recordings**

Single-channel measurements were performed using planar lipid bilayers (14). Unless otherwise indicated, proteoliposomes containing purified recombinant RyR1s were added to the cis (cytosolic) chamber of a bilayer apparatus containing 0.25 M KCl in the cis chamber and 20 mM KCl in the trans (SR luminal) chamber in 20 mM KHEPES, pH 7.4, and 2 μM Ca\textsuperscript{2+}. After the appearance of channel activity, the trans KCl concentration was increased to 0.25 M KCl. A strong dependence of WT channel activity on cis Ca\textsuperscript{2+} concentration indicated that the cytosolic region faced the cis chamber of the bilayer. The trans side of the bilayer was defined as ground. Electrical signals were filtered at 2 kHz (0.5 kHz for Ca\textsuperscript{2+} currents at 0 mV), digitized at 10 kHz, and analyzed at 50% threshold setting (14). Data acquisition and analysis of 2-min recordings were performed using a commercially available software package (pClamp, Axon Instruments). Single-channel activities were also recorded in symmetrical 0.25 M KCl solution with 10 mM Ca\textsuperscript{2+} on the trans side, and the reversal potential (\(E_{\text{rev}}\)) was measured to determine permeability ratios. The permeability ratio of Ca\textsuperscript{2+} versus K\textsuperscript{+} (\(P_{\text{Ca}}/P_{\text{K}}\)) was calculated using a modified form of the Goldman-Hodgkin-Katz Equation 1,

\[
E_{\text{rev}} = -\frac{RT}{F} \ln \left( \frac{[K]^2}{[C]} \right) + \frac{1}{2} \left( \frac{P_{\text{Ca}}}{P_{\text{K}}} \right) \left( \frac{[C]}{[K]} \right)^{1/2}
\]

(Eq. 1)

**Computational methods**

To model the effects of the mutations, in silico mutations were performed for each of the three open-channel cryo-EM structures (PDB codes 5tal, 5t9v, and 5ta3). Amino acid substitutions were performed using Eris (47, 48) followed by side-chain optimization in GROMACS 4.6.1 (49). Starting structures from Eris were obtained from 40 independent Monte-Carlo simulations with 20 steps each. The lowest energy structure among the 40 structures was selected as the starting point for GROMACS simulations. Each system was solvated using the TIP3P water model. Simulations were performed using the CHARMM36 force field (50, 51). The systems were equilibrated with 10,000 steps of steepest descent energy minimization followed by 200 ps of constant volume MD simulation performed at 310 K. Simulations were performed using a time step of 2 fs with all bonds constrained using the LINCS algorithm (52). Particle mesh Ewald was used for long-range electrostatic interactions. A 10-Å cutoff was used for non-bonded interactions. For production runs, 10 ns of MD simulations were performed at constant temperature and pressure of 310 K and 1 atm followed by 4 ns of annealing. For annealing simulations, pressure was held constant at 1 atm, whereas temperature was linearly decreased from 310 to 200 K. A 10,000 kJ/(mol·nm) potential was applied to fix the backbone atoms throughout the simulations and prevent large structural rearrangements in the absence of the membrane. For analysis, 10 structures sampled evenly over the last 5 ns of the simulation were extracted. Pore profiles were calculated using the HOLE program (53). Data are the mean and standard error among 30 pore profiles computed from the sampled structures.

The Adaptive Poisson-Boltzmann Solver (APBS) as applied in APBSmem (28) was used to examine the electrostatics of Ca\textsuperscript{2+} and K\textsuperscript{+} ion passage through the pore. APBSmem estimates the protein-ion electrostatic interaction energy by subtracting the energies of the membrane-embedded protein without the ion and the energy of the ion in bulk water from the total energy of the protein-ion assembly embedded in the lipid bilayer. For analysis, 10 WT and mutant structures sampled evenly over the last 5 ns of the respective simulations were extracted. Two focusing layers with a grid length of 90 Å and a total of 97 grid points in the x, y, and z dimensions were employed. Ion concentrations were set at 0.1 M, both for Ca\textsuperscript{2+} and K\textsuperscript{+}; the temperature was 298 K, and non-linear Poisson-Boltzmann method was engaged in Adaptive Poisson-Boltzmann Solver (29). To omit the membrane from the pore region, the upper and lower exclusions were set at 16 Å. Based on previously reported calculations of Ca\textsuperscript{2+} solvation energy profile of the TRPV1 receptor (54), membrane thickness was 42.5 Å, and headgroup thickness was 7 Å. During the calculation, the dielectric constant for protein and membrane was set at 2.0, and the dielectric constant for headgroups and solvent was set at 80.0. The ion step size was 1.0 Å along the channel axis and the ionic radius used for Ca\textsuperscript{2+} and K\textsuperscript{+} were 1.03 and 1.41 Å, respectively.

**Biochemical assays and data analysis**

Free Ca\textsuperscript{2+} concentrations were determined by adding varying amounts of Ca\textsuperscript{2+} to 1 mM EGTA solutions. Free Ca\textsuperscript{2+} concentrations were measured using a Ca\textsuperscript{2+}-selective electrode. Differences between samples were analyzed using Student’s t
test or analysis of variance followed by Tukey’s test, where $p < 0.05$ was considered significant.

**Author contributions**—L. X., D. D. M., N. V. D., and G. M. designed the experiments; L. X., D. D. M., V. R. C., Y. W., and D. A. P. performed the experiments, and D. D. M. and G. M. wrote the paper. All authors reviewed the results and approved the final version of the manuscript.

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