The Cytoplasmic Region of Inner Helix S6 Is an Important Determinant of Cardiac Ryanodine Receptor Channel Gating*

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The ryanodine receptor (RyR) channel pore is formed by four S6 inner helices, with its intracellular gate located at the S6 helix bundle crossing region. The cytoplasmic region of the extended S6 helix is held by the U motif of the central domain and is thought to control the opening and closing of the S6 helix bundle. However, the functional significance of the S6 cytoplasmic region in channel gating is unknown. Here we assessed the role of the S6 cytoplasmic region in the function of cardiac RyR (RyR2) via structure-guided site-directed mutagenesis. We mutated each residue in the S6 cytoplasmic region of the mouse RyR2 (K4876QQEQVKEDM4884) and characterized their functional impact. We found that mutations Q4876A, V4880A, K4881A, and M4884A, located mainly on one side of the S6 helix that faces the U motif, enhanced basal channel activity and the sensitivity to Ca2+ or caffeine activation, whereas mutations Q4877A, E4878A, Q4879A, and D4883A, located largely on the opposite side of S6, suppressed channel activity. Furthermore, V4880A, a cardiac arrhythmia-associated mutation, markedly enhanced the frequency of spontaneous openings and the sensitivity to cytosolic and luminal Ca2+ activation of single RyR2 channels. V4880A also increased the propensity and reduced the threshold for arrhythmogenic spontaneous Ca2+ release in HEK293 cells. Collectively, our data suggest that interactions between the cytoplasmic region of S6 and the U motif of RyR2 are important for stabilizing the closed state of the channel. Mutations in the S6/U motif domain interface likely destabilize the closed state of RyR2, resulting in enhanced basal channel activity and sensitivity to activation and increased propensity for spontaneous Ca2+ release and cardiac arrhythmias.

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The ryanodine receptor type 2 (RyR2)5 is an intracellular Ca2+ channel located in the sarcoplasmic reticulum (SR) membrane of cardiac muscle cells and mediates the release of SR Ca2+, which triggers muscle contraction (1–3). RyR2-mediated SR Ca2+ release can also influence membrane excitability and cardiac rhythm. Naturally occurring mutations in RyR2 have been linked to various forms of cardiac arrhythmias and cardiomyopathies. Most of the disease-associated RyR2 mutations characterized enhance RyR2 activity by sensitizing the channel to activation by luminal and/or cytosolic Ca2+ (4, 5). However, the exact molecular mechanisms underlying the action of most disease-causing RyR2 mutations are unknown. The crystal structures of the N-terminal and other domains of RyR have provided precise and detailed insights into the structural basis of action of disease-causing mutations located in these domains (6–13). For instance, most of the N-terminal disease mutations are located in domain-domain interfaces. It is believed that domain-domain interactions in the N-terminal region are important for stabilizing the closed state of the RyR channel. Mutations in these domain interfaces may weaken the domain-domain interactions and thus destabilize the closed state of the channel, rendering the channel to open spontaneously and more sensitive to activation (6, 8, 12). Hence, identifying domain-domain interactions that are important for stabilizing the closed state of the channel is important for understanding the molecular mechanisms of action of disease-associated RyR mutations. Unfortunately, atomic resolution structures are not available for most part of the RyR channel. Our understanding of the molecular mechanisms of disease-associated mutations located in regions without detailed structural information has been challenging.

Recently, a near-atomic resolution (3.8 Å) 3D structure of the skeletal muscle RyR (RyR1) has been solved by cryo-electron microscopy (14). Although, at this resolution, the structure may not provide the exact locations of amino acid side chains, it does provide a useful and important structural framework and guidance for structure-function relationship studies of RyR2, especially of the channel domain, where a better resolution was observed. The RyR2 channel domain is one of the disease-causing mutation hot spots, but the mechanisms of disease muta-

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FIGURE 1. Location of the S6 cytoplasmic region in the three-dimensional structure of RyR2. The 3D structure of two RyR2 monomers (30) is shown. The central domain, U motif, CTD, and S6 inner helix are highlighted. The inset is a close-up view showing the cytoplasmic region of the S6 inner helix encompassing residues 4876QQEQVKEDM4884.

Results

Role of the Cytoplasmic Region of the S6 Inner Helix in RyR2 Activation—Structural studies suggest that the cytoplasmic region of the S6 inner helix interacts with the U motif of the central domain (14, 15), but the functional significance of this S6/U motif interaction is unknown. To assess the role of the S6 cytoplasmic region in RyR2 function, we mutated each residue in the S6 cytoplasmic region (4876QQEQVKEDM4884) of the mouse RyR2 (Fig. 1) and determined its impact on the response of RyR2 to caffeine activation. As shown in Fig. 2, the amplitude of Ca2+ release in HEK293 cells expressing the RyR2 WT increased progressively with each repeated addition of caffeine (from 0.05 to 0.5 mM) and then decreased with further additions of caffeine (from 1.0 to 5 mM). This decrease likely resulted from the depletion of intracellular Ca2+ stores by the prior additions of caffeine (0.025 to 0.5 mM) (Fig. 2, A and F). The caffeine response of HEK293 cells expressing the V4880A and K4881A mutants was shifted to the left (Fig. 2, H, I, and L), whereas the caffeine response of HEK293 cells expressing the Q4877A, E4878A, Q4879A, and D4883A mutants was shifted to the right (Fig. 2, B–F). The caffeine responses of the Q4876A, E4882A, and M4884A mutants were similar to that of the WT (Fig. 2, G and J–L). Thus, the V4880A and K4881A mutations enhance the response of RyR2 to caffeine activation, whereas the Q4877A, E4878A, Q4879A, and D4883A mutations inhibit it.

Mutations in the Cytoplasmic Region of S6 Alter the Basal Activity and Ca2+-dependent Activation of [3H]ryanodine Binding—To gain further insights into the role of the S6 cytoplasmic region in RyR2 function, we used [3H]ryanodine binding assays to assess the impact of mutations in the S6 cytoplasmic region on the basal activity of the RyR2 channel and the sensitivity of the channel to Ca2+-dependent activation. Because ryanodine only binds to the open state of the RyR channel, [3H]ryanodine binding has been widely used as an index for RyR channel activity. As shown in Fig. 3, in the near absence of activating Ca2+ (<16 nM), [3H]ryanodine binding to RyR2 WT shows a basal activity of ~3% of the maximum binding. On the other hand, the level of [3H]ryanodine binding to the Q4876A, V4880A, K4881A, and M4884A mutants in the near absence of activating Ca2+ is a close-up view showing the cytoplasmic region of the S6 inner helix encompassing residues 4876QQEQVKEDM4884.

Among the mutations in the S6 cytoplasmic region that enhance RyR2 activity, V4880A displays the most dramatic impact on [3H]ryanodine binding (Fig. 3). Importantly, V4880A is a naturally occurring RyR2 mutation associated with catecholaminergic polymorphic ventricular tachycardia (16). To further understand the mechanism of action of this disease-causing RyR2 mutation, we assessed the impact of V4880A on
FIGURE 2. Effect of mutations in the cytoplasmic region of S6 on caffeine activation of RyR2. A–E and G–K, HEK293 cells were transfected with RyR2 WT (A), Q4877A (B), E4878A (C), Q4879A (D), E4883A (E), Q4876A (G), V4880A (H), K4881A (I), E4882A (J), or M4884A (K). The fluorescence intensity of the Fluo-3-loaded transfected cells before and after repeated additions of increasing concentrations of caffeine (0.025–5 mM) was monitored continuously. F and L, Ca\textsuperscript{2+} release-cumulative caffeine concentration relationships in HEK293 cells transfected with RyR2 WT and mutants that shift the caffeine response curve to the right (F) or mutants that have little effect on caffeine response or shift the caffeine response curve to the left (L). The amplitude of each caffeine peak was normalized to that of the maximum peak for each experiment. Data shown are mean ± S.E. (n = 3–4; *, p < 0.05 versus WT).

FIGURE 3. Effect of the S6 inner helix mutations on [3H]ryanodine binding to RyR2. A and B, [3H]ryanodine binding to cell lysate prepared from HEK293 cells expressing RyR2 WT and the S6 mutants Q4877A, E4878A, Q4879A, E4882A, and D4883A (A) or the S6 mutants Q4876A, V4880A, K4881A, and M4884A (B) was carried out at various Ca\textsuperscript{2+} concentrations (0.2 nM to 0.1 mM), 800 mM KCl, and 5 nM [3H]ryanodine. The amounts of [3H]ryanodine binding at various Ca\textsuperscript{2+} concentrations were normalized to the maximal binding (100%). C, mutating the cytoplasmic region of the S6 inner helix of RyR2 affects the basal level of [3H]ryanodine binding to RyR2 in the near absence of activating Ca\textsuperscript{2+} (<20 nM). D, the EC\textsubscript{50} values of Ca\textsuperscript{2+} activation of [3H]ryanodine binding to RyR2 WT and the S6 mutants. The data points shown are mean ± S.E. from three to five separate experiments (*, p < 0.05 versus WT).
RyR2 gating using single channel recordings in planar lipid bilayers. As shown in Fig. 5, a single RyR2 WT channel exhibited very few opening events at 45 nM cytosolic Ca\(^{2+}\) (Fig. 5, A and C). However, at the same cytosolic Ca\(^{2+}\) concentration, single V4880A mutant channels showed a markedly increased frequency of openings with an event frequency of 4.68 ± 1.00 events/s (Fig. 5, B and C), which is significantly higher than that of single RyR2 WT channels (1.01 ± 0.56 events/s) (p < 0.01). The V4880A mutation also increased the open probability (Po) (Fig. 5D) and decreased the mean closed time (Tc) (Fig. 5F), but had no effect on mean open time (To) (Fig. 5E) of single RyR2 channels. These data suggest that the V4880A mutation destabilizes the closed state of the RyR2 channel, leading to enhanced channel openings in the near absence of activating Ca\(^{2+}\), which is consistent with the increased basal level of \(^{3}\)H]ryanodine binding (Fig. 3).

The V4880A Mutation Increases the Sensitivity of Single RyR2 Channels to Cytosolic Ca\(^{2+}\) Activation—RyR2 can be activated by cytosolic or luminal Ca\(^{2+}\). To determine whether the V4880A mutation affects the activation of RyR2 by cytosolic Ca\(^{2+}\), we carried out a single-channel analysis of the RyR2 WT and V4880A mutant channels in the presence of various cytosolic Ca\(^{2+}\) concentrations and in the near absence of luminal Ca\(^{2+}\) (~45 nM). As shown in Fig. 6, single RyR2 WT channels were activated by cytosolic Ca\(^{2+}\) with an EC\(_{50}\) of ~0.25 μM. On the other hand, single V4880A mutant channels were activated by cytosolic Ca\(^{2+}\) with an EC\(_{50}\) of ~0.15 μM. Thus, the V4880A mutation enhances the cytosolic Ca\(^{2+}\) activation of RyR2.

The V4880A Mutation Increases the Luminal Ca\(^{2+}\) Sensitivity of Single RyR2 Channels—We next assessed whether the V4880A mutation also affects the sensitivity of RyR2 to activation by luminal Ca\(^{2+}\). Single RyR2 WT and V4880A mutant channels were first inactivated by reducing the cytosolic Ca\(^{2+}\) concentration to ~45 nM using EGTA. The luminal Ca\(^{2+}\) concentration was then increased stepwise from 45 nM to 40 mM. As shown in Fig. 7, single RyR2 WT channels were barely active at luminal Ca\(^{2+}\) concentrations of less than 1 mM in the near absence of cytosolic Ca\(^{2+}\) (45 nM). However, under the same conditions, single V4880A mutant channels were activated by luminal Ca\(^{2+}\) at 2.5 μM (Fig. 7C). Therefore, the V4880A mutation also increases the luminal Ca\(^{2+}\) activation of RyR2.

The V4880A Mutation Enables the Propensity for Store Overload-induced Ca\(^{2+}\) Release in HEK293 Cells—We have shown previously that enhanced luminal and/or cytosolic Ca\(^{2+}\) activation of RyR2 increases the propensity for spontaneous release of Ca\(^{2+}\) from the SR, leading to diastolic Ca\(^{2+}\) overload and severe cardiac dysfunction in vivo.
FIGURE 6. **Effect of V4880A on the cytosolic Ca$^{2+}$ activation of single RyR2 channels.** A and B, single-channel activities of RyR2 WT (A) and the V4880A (B) mutant were recorded in a symmetrical recording solution containing 250 mM KCl and 25 mM Hepes (pH 7.4). The Ca$^{2+}$ concentration on the cytoplasmic (cyto) and luminal face of the channel was first adjusted to 45 nM. The cytosolic Ca$^{2+}$ concentration was then increased from 45 nM to various levels by addition of aliquots of CaCl$_2$ solution. C, the relationships between open probability and pCa of single RyR2 WT (solid circles) and the V4880A mutant (open circles). The data points shown are mean ± S.E. from four RyR2-WT and 11 V4880A single channels.

FIGURE 7. **Effect of V4880A on the luminal Ca$^{2+}$ activation of single RyR2 channels.** A and B, single-channel activities of RyR2 WT (A) and V4880A mutant (B) were recorded in a symmetrical recording solution containing 250 mM KCl and 25 mM Hepes (pH 7.4). The Ca$^{2+}$ concentration on the cytoplasmic (cyto) and luminal face of the channel was first adjusted to 45 nM. The luminal Ca$^{2+}$ concentration was then increased to various levels by addition of aliquots of CaCl$_2$ solution. C, the relationships between open probability and luminal pCa of single RyR2 WT (solid circles) and the V4880A mutant (open circles). The data points shown are mean ± S.E. from five RyR2-WT and four V4880A single channels.
Ca$_2^{+}$ release during store Ca$_2^{+}$ overload, a phenomenon we referred to as store overload-induced Ca$_2^{+}$ release (SOICR) (17, 18). Thus, it is possible that the V4880A mutation that enhances both cytosolic and luminal Ca$_2^{+}$ activation of RyR2 may also increase the propensity for SOICR. To test this possibility, we monitored SOICR in HEK293 cells expressing the RyR2 WT or the V4880A mutant using single-cell Ca$_2^{+}$ imaging. Caff, caffeine. C, percentages of RyR2 WT (220 cells) and V4880A (171 cells) cells that display Ca$^{2+}$ oscillations at various extracellular Ca$^{2+}$ concentrations. Data shown are mean ± S.E. (n = 3; **, p < 0.01 versus WT).

The V4880A Mutation Reduces the Threshold for SOICR—SOICR is thought to occur when the store Ca$^{2+}$ load exceeds a threshold level (SOICR threshold) (17–23). To assess whether the V4880A mutation affects the SOICR threshold, we measured the endoplasmic reticulum (ER) luminal Ca$^{2+}$ dynamics using the ER luminal Ca$^{2+}$ sensing protein D1ER (22, 24). As shown in Fig. 9, HEK293 cells expressing RyR2 WT displayed spontaneous ER Ca$^{2+}$ oscillations upon elevating extracellular Ca$^{2+}$ from 0 to 2 mM. We determined the SOICR activation threshold (FSOICR) at which SOICR occurred and the SOICR termination threshold (Ftermi) at which SOICR terminated. Consistent with its increased propensity for SOICR, the V4880A mutation reduced the SOICR activation threshold (79% versus 93% in the WT) (p < 0.05) (Fig. 9C). The V4880A mutation also reduced the SOICR termination threshold (53% versus 62% in the WT) (p < 0.05) (Fig. 9D). The fractional Ca$^{2+}$ release during SOICR (activation threshold — termination threshold) was calculated by subtracting the termination threshold from the activation threshold. The maximum FRET signal F$_{\text{max}}$ is defined as the FRET level after tetracaine treatment. The minimum FRET signal F$_{\text{min}}$ is defined as the FRET level after caffeine treatment. F, the store capacity was calculated by subtracting F$_{\text{min}}$ from F$_{\text{max}}$. Data shown are mean ± S.E. (n = 4; *, p < 0.05 versus WT; NS, not significant).

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threshold) in V4880A mutant cells (26%) was also reduced compared with that in RyR2 WT cells (32%) (p < 0.05) (Fig. 9F). There was no significant difference in the store capacity (Fmax − Fmin) between RyR2 WT and the V4880A mutant cells (Fig. 9F). It should be noted that SOICR did not occur in control HEK293 cells expressing no RyR2 and that SOICR was not affected by the IP3R inhibitor xestospongic C (23), indicating that SOICR is mediated by RyR2. Together, these data indicate that the V4880A mutation enhances the propensity for SOICR by reducing the SOICR threshold.

Discussion

RyRs are the largest known ion channels, important for a number of cellular processes ranging from muscle contraction to neuronal function (1–3). Despite their pivotal significance, the structure-function relationships of RyRs are poorly understood, in part because of their gigantic size and the lack of atomic resolution structures. Yan et al. (14) have recently made a major advance in our understanding of the structure-function relationships of especially the ion-conduction pathway of RyR by solving the 3D structure of RyR at a near-atomic resolution (3.8 Å). Based on this structure, the ion-conducting pathway of RyR is formed by four elongated S6 inner helices. The N-terminal region of the S6 helix is embedded in the membrane, where it forms the central pore cavity, whereas the middle region of the S6 helix, corresponding to the helix bundle crossing region located at the border zone between the membrane and cytoplasm, encompasses the constriction point or the ion gate of the channel (Fig. 1). These structural features are consistent with our previous functional studies demonstrating the critical role of the S6 helix bundle crossing region in the regulation and gating of RyR2 (25–29). However, the functional significance of the C-terminal (cytoplasmic) region of the S6 inner helix is unknown.

The C-terminal region of S6 is located in the cytoplasm, where it interacts with the U motif of the central domain of RyR. It has recently been proposed that the central domain of RyR acts as a transducer that converts the conformational changes in the large cytosolic assembly to the channel pore domain (14, 15, 30). Because the intracellular gate of RyR is formed by the S6 helices, movements of the cytoplasmic region of S6 would directly affect the opening and closure of the gate located in the helix bundle crossing region of S6. Hence, interaction between the cytoplasmic region of S6 and U motif would be expected to be critical for controlling the gating of the channel, but functional evidence is lacking. To assess the functional significance of this S6/U motif interaction, we mutated each residue in the S6 cytoplasmic region (Q4876QQEQVKEDM4884) (Fig. 1). We found that the Q4876A, V4880A, K4881A, and M4884A mutations increase the activity of RyR2, whereas the Q4877A, E4878A, Q4879A, and D4883A mutations located in the cytoplasmic region of S6 would destabilize the open state of the channel. Hence, mutations located in the S6/U motif interface would destabilize the closed state of RyR2. In support of this view, we indeed found that the Q4876A, V4880A, K4881A, and M4884A mutations located in the S6/U motif interface enhanced basal activity and increased the sensitivity of RyR2 to activation. Further, the V4880A mutation increased the opening event frequency of single RyR2 channels in the near absence of activating Ca2+ and reduced the mean closed time but had no effect on the mean open time of single RyR2s. These data are consistent with the notion that the V4880A mutation destabilizes the closed state of the channel.

We have shown previously that the S6 helix bundle crossing region is critical for channel activation by luminal and cytosolic Ca2+ (29). Mutations in the helix bundle crossing region reduced the open duration and open probability of single RyR2 channels. Hence, it is possible that the Q4877A, E4878A, Q4879A, and D4883A mutations located in the cytoplasmic region of S6 may affect Ca2+ activation or destabilize the open state of RyR2. Further studies are required to test these possibilities.

Recently, Des Georges et al. (34) identified a Ca2+ binding site located in the interface between the central domain and the carboxyl-terminal domain (CTD) of RyR1. Based on this structural information, one can speculate that binding of Ca2+ at the interface between the central domain and CTD may cause an outward movement of the S6 helices, leading to dilation of the S6 helix bundle and, thus, opening of the gate. The V4880A mutation, which destabilizes the closed state of the channel, may reduce the free energy that is required to open the gate and, thus, makes the channel more sensitive to activation by Ca2+. Des Georges et al. (34) also identified an ATP binding site located in the interface between the S6-CTD junction and the U motif. Interestingly, the interface between the cytoplasmic region of S6 and U motif constitutes part of the binding pocket for ATP. Based on our data that S6/U-motif interaction is important for stabilizing the closed state of the channel, ATP, a physiological agonist of RyRs, may enhance RyR2 channel activity by promoting destabilization of the closed state of the channel. Hence, the S6/U motif interface represents an important regulatory element of RyR channel gating.

The channel domain is one of the hot spots of disease-causing RyR2 mutations (4, 5). The mechanisms underlying the action of most of the disease mutations located in the channel domain are largely unknown. The V4880A mutation located in the cytoplasmic region of S6 is a disease-causing mutation (16). We found that the V4880A mutation markedly increases the...
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frequency of spontaneous channel openings in the absence of activating Ca\(^{2+}\), indicating that this mutation destabilizes the closed state of the channel. The V4880A mutation also sensitizes the RyR2 channel to activation by cytosolic or luminal Ca\(^{2+}\) and enhances the propensity for spontaneous Ca\(^{2+}\) release in HEK293 cells. Interestingly, the Val\(^{4880}\) residue is located in the S6/U motif interface, where its hydrophobic side chain may interact with the hydrophobic side chains of residues Ile\(^{4173}\), Phe\(^{4174}\), and Val\(^{4177}\) of the U motif (Fig. 10). Thus, like the disease-causing N-terminal RyR mutations that weaken domain interactions in the N-terminal region of RyR (6, 8, 12), the disease-causing V4880A mutation may destabilize the closed state of the channel by weakening the hydrophobic interactions between the S6 helix and the U motif of the Central domain.

In summary, our data demonstrate that the cytoplasmic region of the S6 inner helix of RyR2 is an important determinant of channel stability and activation. Disease-causing RyR2 mutations in this region enhance the propensity for spontaneous Ca\(^{2+}\) release and the susceptibility to cardiac arrhythmias by weakening the interaction between S6 and the U motif and destabilizing the closed state of the channel. Stabilizing the S6/U motif interaction may present a therapeutic approach for suppressing enhanced channel activity.

Experimental Procedures

Construction of RyR2 Mutations Located in the Cytoplasmic Region of the S6 Inner Helix—The S6 RyR2 mutations (Q4876A, Q4877A, E4878A, Q4879A, V4880A, E4882A, D4883A, and M4884A) were generated by the overlap extension method using PCR (25, 35). Briefly, the NruI-NotI (in the vector) fragment containing the Q4876A, Q4877A, E4878A, Q4879A, V4880A, E4882A, D4883A, or M4884A mutation was obtained by overlapping PCR and used to replace the corresponding WT fragment in the full-length mouse RyR2 cDNA in pcDNA5. All mutations were confirmed by DNA sequencing.

Generation of Stable, Inducible HEK293 Cell Lines Expressing RyR2 WT and the V4880A Mutant—Stable, inducible HEK293 cell lines expressing RyR2 WT and V4880A were generated using the Flp-In T-REx Core Kit from Invitrogen. Briefly, Flp-In T-REx HEK293 cells were co-transfected with the inducible expression vector pcDNA5/FRT/TO containing the WT or mutant RyR2 cDNA and the pOG44 vector encoding the Flp recombinase in 1:5 ratios using the Ca\(^{2+}\) phosphate precipitation method. The transfected cells were washed with PBS 24 h after transfection, followed by a change into fresh medium for 24 h. The cells were then washed again with PBS, harvested, and plated onto new dishes. After the cells had attached (~4 h), the growth medium was replaced with a selection medium containing 200 μg/ml hygromycin (Invitrogen). The selection medium was changed every 3–4 days until the desired number of cells were grown. The hygromycin-resistant cells were pooled, aliquoted (1 ml), and stored at ~80 °C. These positive cells are believed to be isogenic because the integration of RyR2 cDNA is mediated by the Flp recombinase at a single FRT site.

Caffeine-induced Ca\(^{2+}\) Release in HEK293 Cells—The free cytosolic Ca\(^{2+}\) concentration in transfected HEK293 cells was measured using the fluorescence Ca\(^{2+}\) indicator dye Fluo-3 AM (Molecular Probes). HEK293 cells grown on 100-mm tissue culture dishes for 18–20 h after subculture were transfected with 12–16 μg of RyR2 WT or the S6 inner helix mutant cDNAs. Cells grown for 18–20 h after transfection were washed four times with PBS and incubated in Krebs-Ringer-Hepes (KRH) (125 mM NaCl, 5 mM KCl, 1.2 mM KH\(_2\)PO\(_4\), 6 mM glucose, 1.2 mM MgCl\(_2\), 2 mM CaCl\(_2\), and 25 mM HEPES (pH 7.4)) buffer without MgCl\(_2\) and CaCl\(_2\) at room temperature for 40 min and at 37 °C for 40 min. After being detached from culture dishes by pipetting, cells were collected by centrifugation at 1000 rpm for 2 min in a Beckman TH-4 rotor. Cell pellets were loaded with 10 μM Fluo-3 AM in high glucose Dulbecco’s modified Eagle medium at room temperature for 60 min, followed by washing with KRH buffer plus 2 mM CaCl\(_2\) and 1.2 mM MgCl\(_2\) (KRH+ buffer) three times and resuspended in 150 μl of KRH+ buffer plus 0.1 mg/ml BSA and 250 μM sulfinpyrazone. The Fluo-3 AM-loaded cells were added to 2 ml (final volume) KRH+ buffer in a cuvette. The fluorescence intensity of Fluo-3 AM at 530 nm was measured before and after repeated additions of various concentrations of caffeine (0.025–5 mM) in an SLM-Aminco series 2 luminescence spectrometer with 480 nm excitation at 25 °C (SLM Instruments). The peak levels of each caffeine-induced Ca\(^{2+}\) release were determined and normalized to the highest level (100%) of caffeine-induced Ca\(^{2+}\) release for each experiment.

[^H]Ryanodine Binding—HEK293 cells were grown to 95% confluence in a 75-cm\(^2\) flask, dissociated with PBS, and plated in 100-mm tissue culture dishes at ~10% confluence 18–20 h before transfection with RyR2 WT and the S6 inner helix...
mutant cDNAs. After transfection for 24 h, the cells were harvested and lysed in lysis buffer containing 25 mM Tris, 50 mM HEPES (pH 7.4), 137 mM NaCl, 1% CHAPS, 0.5% egg phosphatidylcholine, 2.5 mM DTT, and a protease inhibitor mixture (1 mM benzamidine, 2 μg/ml leupeptin, 2 μg/ml pepstatin A, 2 μg/ml aprotinin, and 0.5 mM PMSF) on ice for 60 min. Cell lysate was obtained after removing the unsolubilized materials by centrifugation twice in a microcentrifuge at 4 °C for 30 min each. Equilibrium [3H]ryanodine binding to cell lysates was performed as described previously (36) with some modifications. [3H]Ryanodine binding was carried out in a total volume of 300 μl of binding solution containing 30 μl of cell lysate, 800 mM KCl, 25 mM Tris, 50 mM Hapes (pH 7.4), 5 nm [3H]ryanodine, and CaCl2, to set free [Ca2+] from pCa 9.89 to pCa 4 and a protease inhibitor mixture at 37 °C for 20 min. Free Ca2+ concentrations were calculated using the computer program of Fabiato and Fabiato (37). The binding mixture was diluted with 5 ml of ice-cold washing buffer containing 25 mM Tris (pH 8.0) and 250 mM KCl and immediately filtered through Whatman GF/B filters presoaked with 1% polyethylenimine. The filters were washed three times, and the radioactivity associated with the filters was determined by liquid scintillation counting. Non-specific binding was determined by measuring [3H]ryanodine binding in the presence of 50 μM unlabeled ryanodine. All binding assays were done in duplicate.

Western Blotting—HEK293 cells grown for 24 h after transfection with RyR2 WT and the S6 inner helix mutant cDNAs were washed with PBS plus 2.5 mM EDTA and harvested in the same solution by centrifugation for 8 min at 700 × g in an IEC Centra-CL2 centrifuge. The cells were then washed with PBS without EDTA and centrifuged again at 700 × g for 8 min. The PBS-washed cells were solubilized in a lysis buffer containing 25 mM Tris, 50 mM Hapes (pH 7.4), 137 mM NaCl, 1% CHAPS, 0.5% soy bean phosphatidylcholine, 2.5 mM DTT, and a protease inhibitor mixture (1 mM benzamidine, 2 μg/ml leupeptin, 2 μg/ml pepstatin A, 2 μg/ml aprotinin, and 0.5 mM PMSF). This mixture was incubated on ice for 1 h. Cell lysate was obtained by centrifuging twice at 16,000 × g in a microcentrifuge at 4 °C for 30 min to remove unsolubilized materials. The RyR2 WT and mutant proteins were subjected to SDS-PAGE (5% gel) (38) and transferred onto nitrocellulose membranes at 100 V for 2 h at 4 °C in the presence of 0.01% SDS (39). The nitrocellulose membranes containing the transferred proteins were blocked for 60 min with 1% casein blocker (Bio-Rad). The blocked membrane was incubated with anti-RyR antibody (34C) (Thermo Scientific, MA3-925, lot no. PG200294, 1:1000 dilution) and then incubated with secondary anti-mouse IgG (heavy and light) antibodies conjugated to horseradish peroxidase (1:20,000 dilution). After washing for 10 min three times, the bound antibodies were detected using an enhanced chemiluminescence kit from Pierce. The intensity of each band was determined from its intensity profile obtained by ImageQuant LAS 4000 (GE Healthcare/Life Sciences), analyzed by ImageJ software, and normalized to that of β-actin (40).

Single-channel Recordings—Recombinant RyR2 WT and V4880A mutant channels were purified from cell lysate prepared from HEK293 cells transfected with the RyR2 WT or the V4880A mutant cDNA by sucrose density gradient centrifugation as described previously (18, 36). Heart phosphatidylethanolamine (50%) and brain phosphatidylserine (50%) (Avanti Polar Lipids), dissolved in chloroform, were combined and dried under nitrogen gas and resuspended in 30 μl of n-decane at a concentration of 12 mg of lipid/ml. Bilayers were formed across a 250-μm hole in a Delrin partition separating two chambers. The trans chamber (800 μl) was connected to the head stage input of an Axopatch 200A amplifier (Axon Instruments, Austin, TX). The cis chamber (1.2 ml) was held at virtual ground. A symmetrical solution containing 250 mM KCl and 25 mM Hapes (pH 7.4) was used for all recordings unless indicated otherwise. A 4-μl aliquot (~1 μg of protein) of the sucrose density gradient-purified recombinant RyR2 WT or the V4880A mutant channels was added to the cis chamber. Spontaneous channel activity was always tested for sensitivity to EGTA and Ca2+. The chamber to which the addition of EGTA inhibited the activity of the incorporated channel presumably corresponds to the cytosolic side of the Ca2+ release channel.

The direction of single-channel currents was always monitored from the luminal to the cytosolic side of the channel unless mentioned otherwise. Recordings were filtered at 2500 Hz. Data analyses were carried out using the pclamp 8.1 software package (Axon Instruments). Free Ca2+ concentrations were calculated using the computer program of Fabiato and Fabiato (37).

Single Cell Cytosolic Ca2+ Imaging—Cytosolic Ca2+ levels in stable, inducible HEK293 cells expressing RyR2 WT or the V4880A mutant were monitored using single-cell Ca2+ imaging and the fluorescent Ca2+ indicator dye Fura-2/AM as described previously (17, 18). Briefly, cells grown on glass coverslips for 8–18 h after induction (as indicated) by 1 μM tetracycline (Sigma) were loaded with 5 μM Fura-2/AM in KRH buffer (125 mM NaCl, 5 mM KCl, 6 mM glucose, 1.2 mM MgCl2, and 25 mM Hapes (pH 7.4)) plus 0.02% pluronic F-127 and 0.1 mg/ml BSA for 20 min at room temperature (23 °C). The coverslips were then mounted in a perfusion chamber (Warner Instruments) on an inverted microscope (Nikon TE2000-S). The cells were perfused continuously with KRH buffer containing increasing extracellular Ca2+ concentrations (0, 0.1, 0.2, 0.3, 0.5, 1.0, and 2.0 mM). Caffeine (10 mM) was applied at the end of each experiment to confirm the expression of active RyR2 channels. Time-lapse images (0.25 frame/s) were captured and analyzed with Compix Simple PCI 6 software. Fluorescence intensities were measured from regions of interest centered on individual cells. Only cells that responded to caffeine were analyzed. The filters used for Fura-2 imaging were λexcitation = 340 ± 26 nm and 387 ± 11 nm and λemission = 510 ± 84 nm with a dichroic mirror (410 nm).

Single-cell Luminal Ca2+ Imaging—Luminal Ca2+ levels in HEK293 cells expressing RyR2 WT or the V4880A mutant were measured using single-cell Ca2+ imaging and the FRET-based ER luminal Ca2+-sensitive chameleon protein D1ER as described previously (22, 24). The cells were grown to 95% confluence in a 75-cm2 flask, passaged with PBS, and plated in 100-mm-diameter tissue culture dishes at ~10% confluence 18–20 h before transfection with D1ER cDNA using the Ca2+ phosphate precipitation method. After transfection for 24 h, the growth medium was changed to an induction medium con-
taining 1 μg/ml tetracycline. In intact cell studies, after induction for ~22 h, the cells were perfused continuously with KRH buffer containing various concentrations of CaCl2 (0, 1, and 2 mM) and tetracaine (1 mM) for estimating the store capacity or caffeine (20 mM) for estimating the minimum store level by depleting the ER Ca2+ stores at room temperature (23°C). Images were captured with Compix Simple PCI 6 software every 2 s using an inverted microscope (Nikon TE2000-S) equipped with an S-Fluor ×20/0.75 objective. The filters used for D1ER imaging were λexcitation = 436 ± 20 nm for CFP and λexcitation = 500 ± 20 nm for YFP, and λemission = 465 ± 30 nm for CFP and λemission = 535 ± 30 nm for YFP with a dichroic mirror (500 nm). The amount of FRET was determined from the ratio of the light emission at 535 and 465 nm.

Statistical Analysis—All values shown are mean ± S.E. unless indicated otherwise. To test for differences between groups, we used Student’s t test (two-tailed) or one-way analysis of variance with post hoc test. p < 0.05 was considered to be statistically significant.

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