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Dynamic Changes in Sarcoplasmic Reticulum Structure in Ventricular Myocytes

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The fidelity of excitation-contraction (EC) coupling in ventricular myocytes is remarkable, with each action potential evoking a \([\text{Ca}^{2+}]_i\) transient. The prevalent model is that the consistency in EC coupling in ventricular myocytes is due to the formation of fixed, tight junctions between the sarcoplasmic reticulum (SR) and the sarcolemma where \(\text{Ca}^{2+}\) release is activated. Here, we tested the hypothesis that the SR is a structurally inert organelle in ventricular myocytes. Our data suggest that rather than being static, the SR undergoes frequent dynamic structural changes. SR boutons expressing functional ryanodine receptors moved throughout the cell, approaching or moving away from the sarcolemma of ventricular myocytes. These changes in SR structure occurred in the absence of changes in \(\text{Ca}^{2+}\) during EC coupling. Microtubules and the molecular motors dynein and kinesin 1(Kif5b) were important regulators of SR motility. These findings support a model in which the SR is a motile organelle capable of molecular motor protein-driven structural changes.

1. Introduction

The sarcoplasmic reticulum (SR) is an intracellular organelle that forms an intricate tubular network throughout ventricular myocytes. A critical function of the SR is to store and release \(\text{Ca}^{2+}\). The SR is a structurally diverse organelle that consists of junctional, corbular, and network SR. The network SR is formed by a series of interconnected tubules that span the region between the transverse-tubules (T-tubules). The junctional SR is where this organelle forms specialized junctions with the sarcolemmal T-tubules, which result in the close juxtaposition (\(\approx 15\) nm) of ryanodine-sensitive \(\text{Ca}^{2+}\) channels (RyRs) in the SR and sarcolemmal voltage-gated L-type \(\text{Ca}^{2+}\) channels [1, 2]. Corbular SR is a protrusion from the SR network that expresses RyRs but does not make junctions with the sarcolemma.

During excitation-contraction (EC) coupling, membrane depolarization opens L-type \(\text{Ca}^{2+}\) channels near the junctional SR. This allows a small amount of \(\text{Ca}^{2+}\) to enter the small cytosolic volume that separates the SR and the T-tubule sarcolemma, rising local \([\text{Ca}^{2+}]_i\); beyond a threshold for RyRs activation via the mechanism of \(\text{Ca}^{2+}\)-induced \(\text{Ca}^{2+}\) release (CICR) [3]. Local \([\text{Ca}^{2+}]_i\) elevations resulting from the activation of a small cluster of RyRs are called “\(\text{Ca}^{2+}\) sparks” [4]. Synchronous activation of multiple \(\text{Ca}^{2+}\) sparks by L-type \(\text{Ca}^{2+}\) channels produces a global rise in \([\text{Ca}^{2+}]_i\) that initiates myocardial contraction. The coupling strength between L-type \(\text{Ca}^{2+}\) channels and RyRs is critically dependent on the proximity and stability of the T-tubule/sarcolemmal-SR junction [5, 6].

EC coupling in ventricular myocytes is remarkably reliable. That is, an action potential invariably evokes a whole-cell \([\text{Ca}^{2+}]_i\) transient that results in contraction. Under steady-state conditions, the amplitude of the whole-cell \([\text{Ca}^{2+}]_i\) transient and contraction remains virtually unchanged for long periods of time. A recent study examined the biophysical basis for the fidelity of action potential-evoked \([\text{Ca}^{2+}]_i\); transient and contraction [7]. This study suggested that the probability of \(\text{Ca}^{2+}\) spark activation \(P_s\) by nearby L-type \(\text{Ca}^{2+}\) channels is very high (\(\approx 1\)) during the plateau of the ventricular action potential. Because the \(P_s\) during EC coupling is dependent on the separation between
L-type Ca\(^{2+}\) channels and RyRs, these findings suggest that the T-tubule sarcolemma-SR junction is a functionally stable structure at least over relatively short periods of time (i.e., minutes). At present, however, the SR structural dynamics in living ventricular myocytes have not been examined.

In this study, we used electron, confocal, and total internal reflection fluorescence (TIRF) microscopy to address this fundamental issue. Our data suggest that the SR in neonatal and adult cardiac myocytes forms a vast, yet dynamic network with mobile SR elements that contain functional RyRs. The microtubule network and the molecular motors dynein and kinesin 1 (referred as Kif5b) regulate SR motility. Interestingly, rearrangements in SR structure were more pronounced in neonatal than in adult myocytes. These findings support a model in which the SR is a motile organelle capable of undergoing rapid microtubule-associated molecular motor protein-driven structural reorganization.

2. Methods

2.1. Isolation of Ventricular Myocytes. Animals were handled according to the guidelines of the University of Washington Institutional Animal Care and Use Committee. Hearts were obtained from adult male Sprague Dawley rats or neonatal mice (<72 hours postpartum) euthanized by a lethal intraperitoneal injection of pentobarbital or decapitation, respectively. Single adult and neonatal ventricular myocytes were enzymatically isolated and kept in short-term culture as previously described [8–10].

2.2. Plasmid Construction and Production of Adenovirus Expressing tRFP-SR and Kif5b. The tagged red fluorescent protein (tRFP) was genetically modified to contain av N-terminal calreticulin signal sequence and a C-terminal ER retention signal, KDEL [11]. A cartoon of our tRFP-SR construct is presented in Figure 2(a). Human Kif5b was a generous gift of Dr. Ronald Vale (University of San Francisco, CA, USA). We constructed a dominant version of Kif5b as previously described [12] and fused blue fluorescent protein (BFP) to the N-terminus of Kif5b-WT and Kif5b-DN. Adenoviruses expressing tRFP-SR, Kif5b-WT, and Kif5b-DN were generated using the ViraPower Adenoviral Expression System kit in accordance with manufacturer’s instructions (Invitrogen, Carlsbad, CA, USA). Isolated neonatal and adult cardiac ventricular myocytes were infected with tRFP-SR, Kif5b-WT, or Kif5b-DN, and experiments were conducted 48 hours postinfection unless otherwise indicated.

2.3. Confocal, Total Internal Reflection Fluorescence Imaging and Analysis. For all experiments, cells were perfused with a physiological saline solution containing in mM: KCl 5, NaCl 140, MgCl\(_2\) 1, Glucose 10, HEPES 10 and CaCl\(_2\) 2, pH 7.4 with NaOH. All experiments were performed at room temperature (22–25°C). Images were acquired using a confocal or total internal reflection fluorescence (TIRF) microscope. Confocal imaging was performed using a Bio-Rad Radiance 2000, Nikon Swept Field, or Nikon A1R-S1 confocal microscopes with a Nikon 60x oil immersion lens (NA = 1.4). The TIRF microscope has a PlanApo (60x; NA = 1.45) oil immersion lens, an Andor iXON CCD camera, and a TILL photonics TIRF illuminator. Images were collected every 10 seconds, unless otherwise indicated. The potential for focus drift during time-lapse microscopy was minimized using Nikon’s Perfect Focus System, which provides continuous, real-time focus drift correction caused by thermal and mechanical conditions. Analysis was performed on cells that showed no signs of focus drift.

Confocal and TIRF images were quantitatively analyzed using National Institutes of Health ImageJ software. Correlative SR bouton distances, velocities, and trajectory mapping were performed using the Manualtracking plug-in. To measure change in tRFP-SR fluorescence, we identified prominent peaks using a top-hat filter with a 5x5 pixel window. Thresholds for the top-hat filter were determined per image to generate a full cell profile. A site was counted as having changed if at least one of the frames of a trace exceeded 3 standard deviations (SD) above or below the mean. To measure SR structural change of adult ventricular myocytes, we used an analysis for quantifying change in transverse-tubule structure [13]. Briefly, tRFP-SR myocytes were processed using fast Fourier transform (FFT) to convert two-dimensional images into frequency domain and its corresponding power spectrum (Figure 7(b)) allowing us to determine the spatial regularity of SR structure and the absolute value of the magnitude of the first harmonic (SR\(_{\text{Power}}\)) via Gaussian fitting with pCLAMP 10 software (Axon Instruments). The graphs shown in Figure 7(c) were fit with either a single Gaussian or sum of two Gaussian functions using Graphpad Prism v5.0a (GraphPad, La Jolla, CA, USA).

To measure changes in [Ca\(^{2+}\)]\(_i\), ventricular myocytes were loaded with the free-acid or AM form of the fluorescent Ca\(^{2+}\) indicator Fluo-4 as previously described [4]. [Ca\(^{2+}\)]\(_i\) signals were imaged using confocal or TIRF microscopy. For the experiments in which a TIRF microscope was used, the perforated patch-clamp technique was used to control membrane potential. The patch pipette solution was composed in mM: amphotericin B (30 μg/mL), penta-potassium form of fluo-4 (100 μM), CaCl\(_2\) 1, HEPES 10, MgATP5, CaCl\(_2\) 2.94, EGTA 10, and pH 7.2 with CsOH. TIRF images were acquired at 30–60 Hz using TILL Image software. Confocal images were acquired using a Nikon swept field or Olympus FV1000 confocal microscope. TIRF and confocal images were analyzed with custom software written in IDL language (Research Systems, Boulder, CO, USA). Briefly, background-subtracted fluorescence signals were normalized by dividing the fluorescence (F) intensity at each time point by the resting fluorescence (F\(_0\)) intensity. Fluorescence signals were calibrated using the “pseudo-ratio” equation [4]:

$$\left[ \frac{\text{Ca}^{2+}}{i} \right] = \frac{K_d(F/F_0)}{K_d/[\text{Ca}^{2+}]_{\text{rest}}} + 1 - (F/F_0),$$

where F is the fluorescence intensity, F\(_0\) is the resting fluorescence, K\(_d\) is the dissociation constant of Fluo4 (1100 nM), and [Ca\(^{2+}\)]\(_{\text{rest}}\) is resting Ca\(^{2+}\) concentration (150 nM).
2.4. Ryandine Labeling. Live-cell labeling of ryandine receptors was accomplished by incubating neonatal myocytes with 100 nM BODIPY-FL-X ryandine (Invitrogen) for 10 minutes at room temperature in Hanks Balanced Salt Solution (HBSS; Invitrogen). Excess BODIPY-FL-X ryandine was removed by washing cells 3 times, 10 minutes each, with HBSS prior to imaging. For adult ventricular myocytes, the concentration of BODIPY-FL-X-ryandine was reduced to 50 nM, and the incubation period was reduced to 5 minutes at room temperature.

2.4.1. Electron Microscopy and Immunogold Labeling. Neonatal ventricular myocytes infected with tRFP-SR were fixed with 2.5% glutaraldehyde in PBS (10 mM phosphate buffer/150 mM NaCl, pH 7.4) for 30 minutes followed with 0.1% NaBH4 in 0.1 M PB (phosphate buffer, pH 7.4) for 15 minutes to inactivate aldehydes. Cells were then fixed with 0.5% osmium tetroxide in PB for 15 minutes, subjected to graded dehydration and flat embedded in a mixture of Spur resin and Eponate 12 resin (Ted Pella, Redding, CA, USA). Ultrathin sections (70 nm) were poststained in 4% uranyl acetate and lead citrate. The samples were viewed on a JEOL JSM 6300F scanning electron microscope with digital image capture. Immunogold labeling of tRFP-SR was performed on tRFP-SR-infected neonatal mouse myocytes grown on coverslips using silver enhancement as previously described [14, 15] and prepared for electron microscopy as described above. The primary antibody, rabbit anti-tRFP (Evrogen, Joint Stock, Russia), was used in 1:5000 dilution and the secondary antibody, ultrasmall gold-conjugated F(ab)2 fragment of goat anti-rabbit IgG (Aurion, Electron Microscopy Services, Hatfield, PA, USA) was used in a 1:100 dilution.

2.4.2. Statistics. Statistical significance was assessed by ANOVA analysis of all data points with pairwise comparisons completed as a posttest, with significance defined as P value less than 0.05. Data are presented as mean ± SEM. The asterisk (*) symbol is used in the figures to illustrate a significant difference between groups.

3. Results

3.1. SR Ca2+ Release Is the Principal Contributor to [Ca2+]i Variance during EC Coupling. We imaged action potential-evoked [Ca2+]i transients in adult and neonatal ventricular myocytes loaded with the fluorescent indicator fluo-4 AM (Figure 1). Action potentials were evoked via field stimulation at a frequency of 1 Hz. The mean [Ca2+]i transient amplitude was 508 ± 28 and 705 ± 67 nM in adult (n = 9) and neonatal (n = 12) ventricular myocytes, respectively. The amplitude of these [Ca2+]i transients did not change over a 5-minute period (Figure 1(a), left panel). Indeed, the coefficient of variation (i.e., standard deviation ÷ mean) of the [Ca2+]i transient peak amplitude over this period of time was 0.12 ± 0.2 and 0.18 ± 0.03 in adult and neonatal ventricular myocytes, respectively (P < 0.05) (Figure 1(a), right panel).

To determine the time course and origin of beat-to-beat variations in [Ca2+]i, we generated signal-averaged [Ca2+]i records (Figure 1(b)) and their associated variance (σ2) (Figure 1(c)) from neonatal and adult ventricular myocytes before and after the application of the SERCA pump inhibitor thapsigargin (1 μM). The variance of the [Ca2+]i transient under control conditions (σ2 transient) is defined by the following equation:

\[
\sigma^2_{\text{transient}} = \sigma^2_{\text{SR}} + \sigma^2_{\text{influx}},
\]

where σ2SR and σ2influx are the variance of SR Ca2+ release and Ca2+ influx, respectively.σ2 transient and σ2influx can be experimentally determined from the [Ca2+]i transients before and after application of thapsigargin. Thus, σ2 SR = σ2 transient − σ2 influx. Our analysis indicate that σ2 transient increased during the onset of the [Ca2+]i transient. Consistent with the coefficient of variation data above, peak σ2 transient was larger in neonatal (20,111 ± 234 nM2) than in adult (5066 ± 74 nM2) ventricular myocytes. Thapsigargin significantly reduced mean peak [Ca2+]i; transient amplitude to 225 ± 9 and 420 ± 39 nM of adult and neonatal ventricular myocytes. Furthermore, σ2 influx was 8963 ± 39 nM2 and 509 ± 9.01 nM2 in neonatal and adult ventricular myocytes. Thus, σ2SR is 4557 nM2 in adult myocytes and 11148 nM2 in neonatal myocytes. These data suggest that the majority of the σ2 transient in adult (∼90%) and neonatal myocytes (∼55%) originates from variations in Ca2+ release from the SR.

3.2. Network, Junctional, and SR Boutons in Living Ventricular Myocytes. Changes in the spatial organization of the SR could be a potential contributor to variations in Ca2+ release from this organelle during EC coupling. To investigate this issue, we infected ventricular myocytes with an adenoaviral vector expressing the tag red fluorescent protein (tRFP-SR) [16] with a calreticulin entrance sequence in its N-terminal and a four-amino-acid sequence (KDEL) in its C-terminal that signals its retention in the SR lumen [11] (Figure 2(a)). Confocal imaging revealed that tRFP-SR fluorescence was confined to the nuclear envelope and a vast, seemingly interconnected tubular SR network that extended throughout neonatal and adult ventricular myocytes (Figures 2(b)-2(c)).

In neonatal ventricular myocytes, the width of these SR tubules at 50% fluorescence amplitude ranged from 0.6 to 3 μm with a mean of 0.8 ± 0.1 μm (n = 120 sites) (Figure 2(b), green). In contrast, adult ventricular myocytes expressing tRFP-SR did not show the convoluted, irregular reticular pattern observed in neonatal myocytes (Figures 2(b)-2(c)). Instead, as previously reported by others [1, 17], the SR forms terminals (i.e., junctional SR) with a periodic distribution of about 1.8 μm along the longitudinal axis of the cell. The diameter of these terminals ranged from 0.8 to 1.2 μm (Figure 2(c), green). The junctional SR at opposite ends of each sarcomere was interconnected by “network” SR.

We observed multiple bright structures protruding from the SR network of neonatal and adult ventricular myocytes (Figures 2(b)-2(c)). We refer to these structures as corbular SR boutons [1, 18]. Corbular SR boutons varied in length.
Figure 1: Beat-to-beat fidelity of EC coupling in ventricular myocytes. (a) Average peak amplitude of action potential evoked (1 Hz) global Ca\(^{2+}\) transients (nM) from adult (closed circles) and neonatal (open circles) ventricular myocytes measured at 1 minute interval for five minutes and corresponding coefficient of variation among the adult and neonatal myocyte population. Dashed line represents average peak [Ca\(^{2+}\)]\(_i\) signal over 5 minutes. (b) Representative [Ca\(^{2+}\)]\(_i\) transient of adult and neonatal myocytes. (c) Representative [Ca\(^{2+}\)]\(_i\) signal variance of adult and neonatal ventricular myocytes and the distribution of peak amplitude [Ca\(^{2+}\)]\(_i\) variance (nM\(^2\)) of adult ventricular myocytes in the presence and absence of the SR Ca\(^{2+}\) pump inhibitor thapsigargin (1 μM).

Consistent with our confocal data, EM images showed multiple Ω-shaped boutons within the SR and rough ER of non-permeabilized myocytes (boxes in Figure 2(d)). tRFP-SR-associated immunogold particles were also found to be broadly distributed through the perinuclear, network, and junctional, corbular SR of permeabilized myocytes (Figure 2(e) inset; arrows indicate immunogold particles).
Figure 2: Imaging of network, junctional, and corbular SR boutons in living and fixed ventricular myocytes. (a) A cartoon of the tRFP-SR construct. (b) Confocal image of representative living neonatal ventricular myocyte expressing tRFP-SR. (c) Confocal images of representative living adult ventricular myocyte. Insets show zooms of a corbular SR bouton (red box) and SR network (green box) in these cells. Graphs show fluorescence intensity profile in the region in the image marked by arrows. (d) Electron microscopy image of non-permeabilized neonatal ventricular myocytes infected with tRFP-SR (scale bar = 1 μm; N = nucleus). Insets depict invaginations of the endoplasmic/sarcoplasmic reticulum (scale bar = 100 nm). Panel Dii shows an example of how the dimensions of SR boutons were made. (e) Immunogold labeling of tRFP-SR (arrows of insets) from a permeabilized neonatal ventricular myocyte infected with tRFP-SR depicting invaginations of the endoplasmic/sarcoplasmic reticulum (scale bar = 500 nm and 100 nm (insets); N = nucleus).
Analysis of these EM images indicated that corbular SR had a height and maximum width of 25 ± 5 nm and 1.1 ± 0.1 μm (n = 50), respectively. The diameter at the “neck” of the corbular SR, where they connect to the SR network, was 156 ± 5 nm. Figure 2(d)ii shows the location of these structures in an electron micrograph from a representative myocyte.

3.3. The SR of Ventricular Myocytes Is Motile. Having found a strategy to monitor SR ultrastructure in living ventricular myocytes, we tested the hypothesis that the SR of neonatal and adult ventricular myocytes is a structurally inert organelle. To do this, time-lapse confocal microscopy was performed using neonatal and adult ventricular myocytes expressing tRFP-SR (Figures 3 and 4). Contrary to our hypothesis, we found a significant level of mobility and reorganization within the SR of ventricular myocytes. Corbular SR boutons moved frequently and for relatively long distances (see Movie 1 in Supplementary Material available online at doi: 10.1155/2011/382586).

Figures 3(a)–3(d) shows two-dimensional confocal images from representative neonatal and adult ventricular myocytes expressing tRFP-SR at various time points after the beginning of imaging. Motile corbular SR boutons are identified in each image with circles of varied colors. We tracked the position of the epicenter of each corbular SR bouton over time. The lines connected to each circle illustrate the trajectories of each of the corbular SR boutons in these images. We found that corbular SR boutons moved exclusively within the SR network of neonatal and adult ventricular myocytes. Furthermore, our analysis suggests that the trajectories and velocities of corbular SR boutons were highly variable. Accordingly, individual corbular SR boutons did not move at a constant speed, with periods of rapid mobility often followed by periods of slowed movements or even stasis (Figures 3(b)–3(d)). For example, while the corbular SR bouton i (green) in the neonatal myocyte in Figures 3(a)–3(b) only moved 0.8 μm at a maximum velocity of 0.06 μm/s, corbular SR bouton ii (orange) in the same cell moved a total of ≈10 μm at a maximum velocity of 0.18 μm/s over a period of 300 seconds.

Motile corbular SR boutons were also observed in the SR of adult ventricular myocytes, albeit with a lower frequency than in neonatal myocytes (Figures 3(c)–3(d)). An example of a motile corbular SR bouton in an adult myocyte is shown in Figure 3(c). The green line shows the trajectory of this specific corbular SR bouton. Like neonatal corbular SR boutons, the velocity and distance traveled by this adult corbular SR bouton was variable. It moved with a maximum velocity of 0.06 μm/s and traveled ≈8 μm over a period of 300 seconds (Figure 3(d)).

Motile corbular SR boutons moved faster in neonatal (0.045 ± 0.01 μm/s, n = 250) than in adult ventricular myocytes (0.020 ± 0.004 μm/s, n = 50; P < 0.05) (Figure 3(e)). The maximum velocity reached by corbular SR boutons was 0.6 μm/s. Neonatal and adult neonatal SR boutons traveled distances of up to 30 μm in a period of 300 seconds (Figure 3(f)). The distance traveled for a corbular SR bouton in a continuous movement event (called “step distance”) in adult and neonatal myocytes ranged from 0.3 to 4.2 μm (Figure 3(g)).

We found that changes in SR structure were not limited to motile SR boutons but included changes in the structure of the SR network of neonatal and ventricular myocytes. An example of this is shown in Figures 4(a)–4(b) and Movie 2 in Supplemental Material. Site i in Figure 4(a) is an area of the SR network of a neonatal ventricular myocyte, where an SR tubule bifurcates into two branches. Time-lapse confocal microscopy of this site showed the formation of a new connection between two adjacent SR branches. In site ii, we were able to capture the sequence of events that likely lead to the formation of these SR connections. The corbular SR bouton marked by the red arrow elongated finally fusing with a neighboring section of SR network. Figure 4(b) shows a similar set of events, but in an adult ventricular myocyte.

We also detected fluctuations of tRFP-SR fluorescence presumably resulting from changes in SR structure in these cells. An example of this type of phenomenon is shown in Figure 4(c) (see also Movie 3 in Supplementary Material). Surface plots of sites i and ii in this adult ventricular myocyte revealed dynamic fluctuations in tRFP-SR fluorescent suggesting structural changes in the junctional SR.

We used TIRF microscopy to resolve subtle changes in SR morphology within the subsarcolemmal space of neonatal and adult ventricular myocytes (Figure 5). In our TIRF microscope, excitation is limited to an evanescent field of about 100 nm from the coverslip. Thus, the axial resolution of the TIRF microscope (∼0.1 μm) is nearly an order of magnitude higher than our confocal microscope (∼1.1 μm). In neonatal myocytes, we observed multiple motile SR projections into the subsarcolemmal space per cell (Figure 5(a); see also Movie 4 in Supplementary Material). Interestingly, these SR structures moved laterally as well as in and out of the evanescent field, suggesting that in neonatal myocytes the SR is forming and dissolving SR-sarcotemia junctions on a regular basis. A similar observation was made using adult ventricular myocytes (Figure 5(b)). Collectively, these confocal and TIRF data suggest that the SR is a dynamic organelle that frequently forms and dissolves SR-sarcotemia junctions in neonatal and adult ventricular myocytes.

3.4. Motile Corbular SR Boutons Express Functional Ryanodine Receptors. We investigated whether the corbular SR boutons and junctional SR observed with confocal and TIRF microscopy contained functional RyRs (Figure 6). To do this, we labeled living neonatal and adult ventricular myocytes expressing tRFP-SR with BODIPY-Ryanodine (100 nM neonatal, 50 nM adult) (Figures 6(a) and 6(c)). The Pearson’s coefficient between tRFP-SR with BODIPY-Ryanodine signals in neonatal and adult ventricular myocytes was 0.69 and 0.80, respectively. We performed simultaneous time-lapse confocal microscopy imaging of our neonatal ventricular myocytes labeled with BODIPY-RyR. The representative two-dimensional confocal images of the cell presented in Figure 6(a) are shown at various time points (Figure 6(b), see also Movie 5 in Supplementary Material). We found
that BODIPY-RyR labeled structures moved with corbular SR boutons and along the tracks of the tRFP-SR labeled SR network at varying speeds over time. As described previously, we tracked the position and distance traveled of the epicenter of each corbular SR bouton labeled with BODIPY-RyR and compared them to unlabeled control cells. Labeled and unlabeled SR boutons traveled similar distances over a period of 300 seconds. These data suggest that tRFP-SR with BODIPY-Ryanodine colocalize in neonatal and adult ventricular myocytes and ryanodine receptors are expressed in diverse structures (i.e., network, junctional, and boutons) and are mobile within the SR of ventricular myocytes.

Next, we tested the hypothesis that motile SR boutons express functional RyRs. Consistent with this, we recorded spontaneous Ca$^{2+}$ sparks from motile SR boutons (Figures 6(d)-6(e)). For example, the corbular SR bouton marked with a green dot in Figure 6(d) moved at a velocity of up to 0.004 $\mu$m/s and multiple Ca$^{2+}$ sparks were recorded from this site. As shown in Figure 6(e), Ca$^{2+}$ sparks were also recorded from corbular SR boutons in adult myocytes. In combination with the data above, these findings suggest that mobile corbular SR boutons express functional ryanodine receptors in ventricular myocytes.

3.5. The Microtubule Network and Microtubule-Associated Motors Dynein and Kif5b Regulate SR Motility in Ventricular Myocytes. We investigated the mechanisms underlying SR motility in ventricular myocytes...
motility in ventricular myocytes. Previous studies have implicated the microtubule network and the molecular motor dynein in the regulation of ER structure and motility in nonmuscle cells [19–22]. Thus, we examined the effects of the microtubule disruptor nocodazole (10 μM) on the SR of neonatal and adult ventricular myocytes expressing tRFP-SR. We found that incubation with 10 μM nocodazole for 2 hours induced significant SR fragmentation in neonatal ventricular myocytes (Figure 7(a)). Nocodazole also seemed to alter SR organization in adult ventricular myocytes (Figures 7(b) and 7(c)). To quantify changes in SR-structure in adult ventricular myocytes, where the SR is organized in a well-defined periodic pattern, we implemented an analysis similar to the one used by Wei et al. [13], to monitor transverse tubule periodicity in ventricular myocytes. Briefly, this analysis involves a Fast Fourier Transform (FFT) analysis of tRFP-SR images, which are converted into a frequency domain that quantifies periodicity. The first harmonic, which centers at 1.7 μm, is the strongest periodicity of the SR-network based upon the spacing between each harmonic (Figure 7(b)). The magnitude of the 1st harmonic, or SRPower, provides a quantitative index for SR structural periodicity. In this analysis, a decrease in SRPower would represent a decrease in SR periodicity and hence its structural integrity. We found that under control conditions the histogram of SRPower values could be fit with a single Gaussian function with a center at 1.70 μm (n = 17). However, in nocodazole-treated cells the SRPower was shifted towards lower SRPower values than under control conditions. Indeed, the SRPower of nocodazole treated cells could be fit with the sum of two Gaussian functions with centers at 0.88 and 1.53 μm (n = 30) (Figure 7(c)). These data suggest that microtubules play a critical role in the maintenance of the structural integrity of the SR network in ventricular myocytes.

Next, we tested the hypothesis that inhibition of the microtubule-associated molecular motor proteins dynein and Kif5b would decrease SR motility. We investigated whether inhibition of the retrograde molecular motor, dynein with erythro-9-(2-hydroxy-3-nonyl) adenine (EHNA) [23] would decrease SR motility in neonatal ventricular myocytes (Figures 8(a) and 8(c)). Confocal images were collected at
Figure 5: TIRF imaging of submembrane SR motility in neonatal and adult ventricular myocytes. (a) TIRF images of representative neonatal ventricular myocyte expressing tRFP-SR. (b) TIRF image of representative adult ventricular myocyte expressing tRFP-SR. Images to the right are two dimensional images (top row) and surface plots of the region delineated by the white.

various time points over a period of 120 minutes in the presence or absence of 30 μM EHNA. Figure 8(a) shows a representative neonatal myocytes before (control) and after the application of EHNA. We mapped the trajectories of corbular SR boutons under control conditions and in the presence of EHNA with colored lines. Note that although corbular SR boutons were observed after incubation with EHNA, motility was significantly reduced (Figure 8(a)). Indeed, EHNA significantly decreased the total distance traveled by neonatal corbular SR boutons by ≈57% from 6.27 ± 0.44 μm (control, n = 6) to 2.69 ± 0.34 μm (EHNA, n = 16) (P < 0.05) (Figure 8(c)).

To examine whether the anterograde microtubule-associated molecular motor protein Kif5b alters SR dynamics, we generated two adenovirus constructs of wild type and a dominant-negative form of Kif5b lacking the motor domain [12] fused to BFP at the N-terminus (Kif5b-WT and Kif5b-DN) and infected ventricular myocytes simultaneously with Kif5b-BFP and tRFP-SR adenoviruses (Figure 8(b)). The Pearson’s coefficient between tRFP-SR and Kif5b-WT-BFP or Kif5b-DN-BFP was 0.82 and 0.81, respectively (Figure 8(b)). These data suggest that the molecular motor Kif5b associates with the diverse structures of the SR network (i.e., network, junctional, and boutons) in ventricular myocytes. Importantly, we found that expression of Kif5b-DN-BFP significantly decreased the total distance traveled by neonatal corbular SR boutons by ≈60% from 7.01 ± 0.31 μm (control, n = 6) to 2.78 ± 0.2 μm (Kif5b-DN-BFP, n = 6) (P < 0.05) (Figure 8(c)).

Finally, we investigated the effects of the dynein inhibitor EHNA on junctional SR in adult ventricular myocytes (Figure 8(d)). We focused on the junctional SR and not on SR boutons because, as noted above, the latter are less frequently observed in adult than in neonatal myocytes. For this analysis, we calculated the percent of junctional SR sites throughout the cell that, over time, underwent fluorescence fluctuations that exceeded 3 standard deviations above or below the mean local tRFP-SR fluorescence. Junctional SR sites were identified automatically with the use of a 5 × 5 top-hat detection filter. Using this analysis, we found that, under control conditions, 4.7 ± 0.2% (n = 20,000 sites from 6 cells) of isolated sites experienced change in tRFP-SR fluorescence in the junctional SR that exceeded the detection threshold. Consistent with the neonatal data above, application of 30 μM EHNA decreased the number of fluctuating junctional SR sites by 45 ± 0.1% (P < 0.05). Collectively, these data suggest that the microtubule network and the molecular motors dynein and Kif5b play a critical role in SR motility of ventricular myocytes.

4. Discussion

This study provides the first characterization of SR structure in living neonatal and adult ventricular myocytes. The main findings of this study were fourfold. First, the SR forms a vast network that is structurally diverse and dynamic. Second, we discovered that corbular SR boutons move throughout
the cell and express functional RyR. Third, by simultaneously imaging \([\text{Ca}^{2+}]_i\) and tRFP-SR, we discovered that the amplitude of the evoked whole-cell \([\text{Ca}^{2+}]_i\) transient remains unchanged even though the SR is undergoing significant remodeling. These data suggest that a static SR network is not required for stable EC coupling in ventricular myocytes. Fourth, SR structure and motility are dependent on an intact microtubule network and the retrograde and anterograde microtubule-associated motors dynein and Kif5b.

Electron microscopy and immunofluorescence approaches have been extensively employed to study SR structure in cardiac muscle [1, 24–26]. These studies have led to the formulation of a model of the SR and its involvement in EC coupling. In this model, the SR is comprised of interconnected longitudinal tubules that terminate in dilated sacs that come into close apposition (≈15 nm) to the sarcolemma. RyRs are expressed in this junctional SR. Together, the junctional SR, RyRs, and nearby sarcolemmal L-type Ca\(^{2+}\) channel form a signaling unit called a “couploin”. The formation of SR-sarcolemmal junctions that underlies the couploin is proposed to be a multistep process [27, 28]. The first event in development of the junctional SR is the “docking” of the SR membrane to the sarcolemma via the protein junctophilin 2. Following the establishment of the SR-sarcolemmal junctions in embryonic myocytes, the proteins calsequestrin, junction, and triadin are targeted and organized within the SR. The arrival of the RyRs and L-type Ca\(^{2+}\) channels and final organization of the macromolecular complex that comprises a Ca\(^{2+}\) release unit within the junctional SR lead to a couploin.
Our data suggest that these SR structures are highly dynamic. This is analogous to what has been described for the endoplasmic reticulum (ER) in nonmuscle cells [20, 29]. The SR is considered to be a specialized derivation of the endoplasmic reticulum (ER) that is dedicated to Ca$^{2+}$ storage and release for the regulation of muscle contraction and EC coupling. The ER of nonmuscle cells is known to have a convoluted structure that is continually reorganizing its structure and moving toward and away from the plasma membrane over time in a variety of living nonmuscle cell models [20, 29]. Our data suggest that like the ER, the SR has the capacity to remodel.

Our results demonstrate that the mean [Ca$^{2+}$], transient amplitude achieves high fidelity yet also displays beat-to-beat variation in [Ca$^{2+}$]. Interestingly, the variance of the [Ca$^{2+}$] attributable to SR Ca$^{2+}$ release was nearly 4-fold larger than adult myocytes. This coincides with a higher degree of SR motility in neonatal than in adult myocytes. Although the exact sources for [Ca$^{2+}$] variance during EC coupling are unclear, it is intriguing to speculate that variations in SR motility of neonatal and adult ventricular myocytes could be a contributing factor.

Similar to studies of the regulation of ER motility [19–22, 29], we demonstrate that the microtubule network regulates SR motility. Although a direct interaction between the microtubule network and the SR has not been shown, it has been postulated to be necessary to mechanically regulate the SR and T-tubule sarcolemma for proper regulation of Ca$^{2+}$ spark activation and termination [30]. Indeed, perturbation of the microtubule network has been shown to alter L-type Ca$^{2+}$ current, [Ca$^{2+}$], transients [30], Ca$^{2+}$ sparks [31], and action potential waveform [32]. Our results suggest intact microtubules are necessary for maintenance of SR ultrastructure.

Kif5b has been implicated in ER motility in nonmuscle cells [29] and in the trafficking of Kv1.5 channels in heterologous cells and rat ventricular myocytes [12]. Likewise, the retrograde motor protein, dynein, has been previously shown to affect the trafficking of Kv1.5 channels in heterologous cells and rat cardiac myocytes [33, 34] and ClC-2 chloride channel in hippocampus [35]. Interestingly, we found that expression of a dominant-negative Kif5b and inhibition of dynein decreased. At present, however, the specific role of Kif5b and dynein in SR motility, couplon formation, and EC coupling is unknown. Future studies should examine the role of microtubules and molecular motor proteins in these important issues.

On the basis of these findings, we propose a new model for the role and regulation of SR structure and motility in EC coupling in ventricular myocytes (Figure 8(e)). In this model, the SR forms a vast interconnected network with distinct microdomains (i.e., junctional domains, corbular domains) throughout the ventricular myocyte. The position of SR structures likely depends on the relative activities of microtubule-associated Kif5b (anterograde motor), and dynein (retrograde motor) regulate SR structure and motility. We propose that the function of these structural rearrangements would include formation of new interconnections with neighboring SR network, formation of new junctions between SR and sarcolemma (i.e., couplons), and
possibly regulation of trafficking of important EC coupling proteins through the SR bouton. An important implication of this model is that the exquisite fidelity of the \([\text{Ca}^{2+}]_i\) transient in ventricular myocytes is not likely produced by the activation of \(\text{Ca}^{2+}\) release from a permanently static SR structure. Rather, it may result from the activation of a temporally averaged number of SR \(\text{Ca}^{2+}\) release units from the formation and dissolution of SR-sarcolemmal junctions that are sensitive to perturbations in the microtubule network such as those observed in cardiac pathology.

5. Limitations

All studies that use cultured ventricular myocytes and pharmacological agents should be interpreted cautiously. For example, cultured adult ventricular myocytes undergo deterioration of their T-tubules and experience electrophysiological changes [36, 37]. Furthermore, it is important to note that although EHNA is known to disrupt the ATPase activity of dynein [23, 35, 38], it also inhibits phosphodiesterases in cardiac myocytes [39]. Thus, we cannot rule
out the possibility that a nodynein molecular motor that is modulated by cAMP levels is involved in SR motility. In spite of these limitations, we contend that novel insights into the bases of SR structure in ventricular myocytes have been obtained in the present experiments. These insights should form the bases of new hypotheses that can be best tested in appropriate animal models.

6. Summary and Conclusions

The present study suggests that normal cardiovascular function is accomplished via complex regulation of SR ultrastructure, [Ca\textsuperscript{2+}]\textsubscript{i}, and contraction. Our study offers a novel conceptual model on the regulation of SR dynamics in ventricular myocytes. Based on this model, we propose that the fidelity of the [Ca\textsuperscript{2+}] transient is not likely produced by the activation of Ca\textsuperscript{2+} release from a permanently static SR but rather may result from the activation of temporally averaged number of SR Ca\textsuperscript{2+} release units forming and dissolving SR-sarcolemmal junctions that are sensitive to perturbations of the microtubule network. Our results and proposal merit further investigation.

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