Differential Integration of Ca^{2+}-Calmodulin Signal in Intact Ventricular Myocytes at Low and High Affinity Ca^{2+}-Calmodulin Targets*

Qiujing Song1, Jeffrey J. Saucerman1, Julie Bossuyt2, §, and Donald M. Bers1, ¶

From the 1Department of Physiology, Loyola University Chicago, Maywood, Illinois 60153, the 2Department of Biomedical Engineering, University of Virginia, Charlottesville, Virginia 22908, and the 4Department of Pharmacology, University of California at Davis, Davis, California 95616

Cardiac myocyte intracellular calcium varies beat-to-beat and calmodulin (CaM) transduces Ca^{2+} signals to regulate many cellular processes (e.g. via CaM targets such as CaM-dependent kinase and calcineurin). However, little is known about the dynamics of how CaM targets process the Ca^{2+} signals to generate appropriate biological responses in the heart. We hypothesized that the different affinities of CaM targets for the Ca^{2+}-bound CaM (Ca^{2+}-CaM) shape their actions through dynamic and tonic interactions in response to the repetitive Ca^{2+} signals in myocytes. To test our hypothesis, we used two fluorescence resonance energy transfer-based biosensors, BsCaM-45 (K_d = \sim 45 \text{ nM}) and BsCaM-2 (K_d = \sim 2 \text{ nM}), to monitor the real time Ca^{2+}-CaM dynamics at low and high affinity CaM targets in paced adult ventricular myocytes. Compared with BsCaM-2, BsCaM-45 tracks the beat-to-beat Ca^{2+}-CaM alterations more closely following the Ca^{2+} oscillations at each myocyte contraction. When pacing frequency is raised from 0.1 to 1.0 Hz, the higher affinity BsCaM-2 demonstrates significant elevation of diastolic Ca^{2+}-CaM binding compared with the lower affinity BsCaM-45. Biochemically detailed computational models of Ca^{2+}-CaM biosensors in beating cardiac myocytes revealed that the different Ca^{2+}-CaM binding affinities of BsCaM-2 and BsCaM-45 are sufficient to predict their differing kinetics and diastolic integration. Thus, data from both experiments and computational modeling suggest that CaM targets with low versus high Ca^{2+}-CaM affinities (like CaM-dependent kinase versus calcineurin) respond differentially to the same Ca^{2+} signal (phasic versus integrating), presumably tuned appropriately for their respective and distinct Ca^{2+} signaling pathways.

Calcium is a well recognized diffusible messenger in many cell types (1). Indeed, Ca^{2+} regulates multiple biological responses in cardiomyocytes, where cytosolic Ca^{2+} oscillates in the range of 0.1–1 \text{ \mu M} on a beat to beat basis (2). Calmodulin (CaM), a ubiquitous Ca^{2+}-sensing protein (3), is an important Ca^{2+} signal transducer in the heart. It is well established that the activated form of CaM, Ca^{2+} -CaM, modulates the function of various downstream targets (4) including ion channels (5–7) and enzymes such as CaM-dependent protein kinase II (CaMKII) (8), myosin light chain kinase (MLCK) (9), nitric-oxide synthase (10), and CaM-dependent phosphatase calcineurin (CaN) (11), which play central roles in diverse cellular processes. For example, CaMKII is implicated in regulation of the excitation-contraction coupling (8, 12) and apoptosis in the heart (13, 14), whereas CaN dephosphorylates transcription factors to regulate gene expression related to cardiac hypertrophy and heart failure (11). Despite the wide appreciation of the importance of Ca, CaM, and CaM targets on cardiac myocyte function, an intriguing question remains unsolved about how CaM and its target proteins differentiate Ca^{2+} signals to ensure the appropriate cellular responses in the heart.

The concentration of free CaM in adult ventricular myocytes is \sim 50–100 \text{ nM}, which is only 1–2% of the total myocyte CaM (15). Thus, the pool of free CaM is limited given the abundance of CaM targets in muscle cells (16). As a result, there is likely intense competition among CaM targets for CaM activation in response to Ca^{2+} signals in cardiomyocytes. Consequently, the action of different CaM targets in processing Ca^{2+} signals may be at least partially determined by their respective affinities for Ca^{2+}-CaM. There is an extensive range of affinities for Ca^{2+}-CaM among CaM targets (K_d values vary from \sim 0.1 to \sim 100 \text{ nM}) (4). For instance, CaN has a particularly high affinity (K_d = \sim 0.1 \text{ nM}) (17) for Ca^{2+}-CaM, whereas the affinity of CaMKII for Ca^{2+}-CaM is much lower (K_d = \sim 50 \text{ nM}) (18). It has been shown that CaN responds to sustained, low amplitude Ca^{2+} signals in lymphocytes and transduces these signals into the activation of nuclear transcriptional factor NFAT (19). In skeletal muscle, activation of CaM caused by sustained Ca^{2+} signals (evoked by motoneuron stimulation) and subsequent nuclear translocation of NFAT has been suggested to modulate muscle type-specific gene expression (20). On the other hand, CaMKII

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1To whom correspondence should be addressed: Dept. of Pharmacology, University of California at Davis, Genome Bldg. 3513, 451 E. Health Science Dr., Davis, CA 95616. Tel.: 530-752-6517; Fax: 530-752-7710; E-mail: dmbers@ucdavis.edu.

2The abbreviations used are: CaM, calmodulin; FRET, fluorescence resonance energy transfer; CaMKII, CaM-dependent protein kinase II; CaN, calcineurin; Ca^{2+}-bound CaM, Ca^{2+}-CaM; MLCK, myosin light chain kinase; YFP, yellow fluorescent protein; CFP, cyan fluorescent protein; HEK, human embryonic kidney; Iso, isoproterenol; F-CaM, fluorescent CaM.
is activated preferentially by transient, high amplitude Ca$_{2+}$ spikes and can undergo autophosphorylation switching it to a prolonged Ca$_{2+}$ independent active state (21). It is found that activation of CaMKII can produce a memory with frequency and amplitude in neurons (22, 23) and skeletal muscle (24). However, there is limited understanding of the potential role of CaMKII and those of the high affinity targets (e.g. CaN), may play in decoding the Ca$_{2+}$ signals in the cardiomyocyte.

Persechini and co-workers (25–29) have recently developed fluorescence resonance energy transfer (FRET)-based biosensors allowing real time studies on the interaction of CaM with CaM targets in response to Ca$_{2+}$ signals in intact cells. These biosensors work by detecting the changes of FRET between two green fluorescent protein variants, a donor (enhanced CFP) and an acceptor (enhanced YFP) connected by a linker, which is derived from the Ca$_{2+}$-CaM-binding domain of the smooth muscle MLCK. Site mutations of the linker sequence confer different Ca$_{2+}$-CaM affinities to the biosensors (25). Two biosensors (BsCaM-45, $K_d = \sim 45 \text{ nM}$ versus BsCaM-2, $K_d = \sim 2 \text{ nm}$) are of particular interest because their apparent Ca$_{2+}$-CaM affinities are similar to those of the low affinity targets (e.g. CaMKII) and those of the high affinity targets (e.g. CaN), respectively. Our initial report using BsCaM-45 demonstrated the feasibility of detecting dynamic beat-to-beat Ca$_{2+}$-CaM changes in adult ventricular myocytes (30). The BsCaM-2 sensor, with higher Ca$_{2+}$-CaM affinity, has been used to measure Ca$_{2+}$-CaM changes in human embryonic kidney (HEK) cells (25) and endothelial cells (28). Here, we compare BsCaM-2 and BsCaM-45 biosensors to test the hypothesis that the different Ca$_{2+}$-CaM affinities of CaM targets may dictate their dynamic and tonic binding to Ca$_{2+}$-CaM in adult ventricular myocytes.

**EXPERIMENTAL PROCEDURES**

**Materials—**Di-8-Anepps, Flu-o-4AM, and Alexa Fluor 488 conjugate calmodulin were from Molecular Probes. High purity calmodulin were from Calbiochem. All other chemicals were from Sigma.

**Construction of Adenoviral Vectors Encoding Biosensors—**
The BsCaM-2 plasmid (29), a kind gift from Dr. A. Persechini (University of Missouri-Kansas City, MO), was incorporated in adenoviruses to ensure the high infection efficiency in the terminally differentiated adult ventricular myocytes. The commercially available AdEasy™ adenoviral vector system (Qbiogene, Inc., Carlsbad, CA) was used to construct the adenoviral vector encoding the BsCaM-2 biosensor following the manufacturer’s instructions. The BsCaM-45 adenoviral vector was previously constructed in our lab (30).

**HEK293 Cell Transfection—**HEK293 cells were cultured in Dulbecco’s modified Eagle’s medium with 5% fetal bovine serum and penicillin/streptomycin for 24 h and then transiently transfected with expression plasmids for BsCaM-2 and BsCaM-45 using the mammalian transfection kit (Stratagene). The cells were cultured for an additional 24 h prior to experiments.

**In Vitro Fluorescence Measurements—**Fluorescence measurements of BsCaM-2 and BsCaM-45 were performed using a spectrofluorometer (SLM, model LS-8100). Excitation and emission slits were set at 4 nm. An excitation wavelength of 430 nm was used, and dual photon counting emission detectors were set at 480 nm ($F_{480}$) and 530 nm ($F_{530}$), respectively. The cytosolic fraction of the transfected HEK cells were diluted in Ca$_{2+}$-free internal solution (in 100 mM potassium aspartate, 30 mM KCl, 20 mM HEPES, 5 mM ATP-DiTris, 5 mM MgCl$_2$, 5 mM NaCl, 3 mM KH$_2$PO$_4$, 5 mM sodium pyruvic acid, 0.5 mM K$_4$-BAPTA, pH 7.2, at 23 °C) with the biosensor concentration –0.8–4.8 nm. Standard Ca$_{2+}$ solutions were added to achieve free Ca$_{2+}$ of 50 μM according to the calculation with the MaxChelator program. Incremental addition of CaM was used to reach various concentrations as indicated in the experiments. CaM binding affinity ($K_d$) of biosensors were estimated using nonlinear regression analysis (Prizm 4.0, GraphPad Software, Inc., San Diego, CA).

**Myocyte Isolation and Adenoviral Infection—**Adult rabbit ventricular myocytes were isolated as previously described (31). All of the procedures were performed in accordance with the Guide for the Care and Use of Laboratory Animals and approved by the Loyola University Chicago Institutional Animal Care and Use Committee. Myocytes were seeded on laminin-coated culture inserts in serum-free PC-1™ medium (Lonza Group Ltd.) supplemented with penicillin (400 units/ml) and streptomycin (400 μg/ml). After 1 h of incubation in 5% CO$_2$ at 37 °C for cell attachment, the nonadherent myocytes were washed off. Then the myocytes were infected for 2 h at multiplicity of infection of 10–100 with adenovirus expressing BsCaM-2 or BsCaM-45 and subsequent fresh medium replacement with one further time change before the experiment −36–48 h later. Adenovirus expressing CaM (gift from Dr. David Yue, Johns Hopkins University) was used to co-infect cells to boost the FRET signals as previously reported (30).

**Confocal Microscopy Imaging—**Inserts containing the cultured myocytes were mounted on the stage of an inverted microscope (Zeiss, LSM5 Pascal) equipped with a 40×1.4 NA water immersion objective lens. The superfusate for experiments was Tyrode solution (140 mM NaCl, 4 mM KCl, 1 mM MgCl$_2$, 2 mM CaCl$_2$, 10 mM HEPES, and 10 mM glucose, pH 7.4, at 23 °C). CFP emission fluorescence at 485 ± 15 nm was measured by confocal microscopy with argon laser excitation at 458 nm, indicating infection and localization. Di-8-Anepps, a dye staining the T-tubules of myocytes, was excited with argon laser excitation at 488 nm with emission at >560 nm to avoid fluorescence signal cross-talk. YFP excitation was at 514 nm, and emitted fluorescence was measured at >530 nm. Acceptor photobleaching was performed by YFP excitation at 514 nm (laser at power of 90% for 45 s). Image-J software was used for image analysis.

**Dynamic Fluorescence Measurements in Ventricular Myocytes—**The cells were superfused with normal Tyrode solution (23 °C) or in the presence of 1 μM isoproterenol (Iso). The cells were field stimulated at 0.1, 0.25, 0.5, 0.75, and 1 Hz. Beat-to-beat fluorescence signals of CFP ($F_{480}$) and YFP ($F_{530}$) from the infected cells were monitored using a standard epifluorescence microscope, attached to two photomultiplier tubes allowing simultaneous detection of CFP and YFP emission. Customized filters included: a CFP exciter 430 ± 10 nm, a 505-nm long pass dichroic filter to separate CFP signal from...
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YFP signals, a CFP emitter 480 ± 10 nm, and a YFP emitter 535 ± 15 nm (Chroma, Brattleboro, VT). FRET signals were presented as the ratio of \(F_{480}\) to \(F_{530}\) (\(F_{480}/F_{530}\) or \(R\)), indicating the \(\text{Ca}^{2+}\)-\(\text{CaM}\) interaction with the biosensors. For intracellular \(\text{Ca}^{2+}\) measurements ([\(\text{Ca}^{2+}\)]), the cells were loaded for 20 min with the membrane-permeant fluorescent \(\text{Ca}^{2+}\) indicator Fluo-4 AM (6 \(\mu\)M). Fluo-4 was excited at 480 ± 5 nm, and emission was measured using a 535 ± 20 nm bandpass filter. [\(\text{Ca}^{2+}\)] signals are presented as background-subtracted normalized fluorescence (\(F/F_0\)) where \(F\) is the fluorescence intensity and \(F_0\) is the resting fluorescence recorded under steady state conditions. All of the fluorescence signals were recorded with the Axon pClamp8 software and analyzed using custom software designed by Dr. Eckard Picht.

F-CaM Binding Experiments in Permeabilized Ventricular Myocytes—After settling in Tyrode solution, some myocytes were switched to internal solution (50 nm free calcium) containing saponin (50 \(\mu\)g/ml) for 20 s to permeabilize the sarclemma (15). Internal solutions with various levels of F-CaM as indicated were immediately washed in, and the cells were incubated at room temperature for at least 4 h to reach steady state. Confocal images of F-CaM were recorded using a laser excitation wavelength of 488 nm and the emission wavelength of 500–530 nm. Note that the F-CaM signal could be calibrated based on the known bath concentration and fluorescence (15). The apparent CaM binding affinity of permeabilized cell was estimated using nonlinear regression analysis with Prism 4.0 (Graph-Pad Software, Inc., San Diego CA).

Statistics—Pooled data are represented as the means ± s.e. Statistical comparisons were made using repeated two-way analysis of variance, paired, and unpaired Student’s \(t\) test where applicable. \(p < 0.05\) was considered significant.

RESULTS

CaM-dependent FRET Responses of \(\text{Ca}^{2+}\)-CaM Biosensors—The goal of the current study is to test the hypothesis that the different affinities of CaM targets for \(\text{Ca}^{2+}\)-CaM alter the transduction of \(\text{Ca}^{2+}\) signals to appropriate cellular processes in adult cardiomyocytes via the \(\text{Ca}^{2+}\)-CaM-CaM target signaling axis. To accomplish this, two versions of FRET-based biosensors with different affinities for \(\text{Ca}^{2+}\)-CaM were established (Fig. 1A). In the absence of \(\text{Ca}^{2+}\)-CaM binding to the linker of the biosensors, excitation of CFP at 430 nm generates FRET between the CFP and YFP. Upon binding of \(\text{Ca}^{2+}\)-CaM to the biosensors, the CFP and YFP move apart, and FRET is reduced because of its strong dependence on the CFP to YFP distance. Reduced FRET is shown as an increased ratio of fluorescence emission of CFP at 480 nm (\(F_{480}\)) to YFP at 530 nm (\(F_{530}\)). Thus, the ratio of \(F_{480}/F_{530}\) is a read-out of \(\text{Ca}^{2+}\)-CaM interaction with the biosensors (Fig. 1B). To ensure that \(\text{Ca}^{2+}\)-CaM-dependent FRET changes reflect the expected biosensor affinities, the \(F_{480}/F_{530}\) signals for BsCaM-2 and BsCaM-45 were assessed in the cytosolic fraction of transfected HEK cells upon sequential addition of CaM in the presence of 50 \(\mu\)M Ca. In the absence of \(\text{Ca}^{2+}\)-CaM, the minimal \(F_{480}/F_{530}\) ratio for BsCaM-2 is 0.542 ± 0.042 (\(n = 5\)) versus BsCaM-45 (0.529 ± 0.058, \(n = 5\)). At saturating \(\text{Ca}^{2+}\)-CaM, the maximal ratio (\(R_{\text{max}}\)) of BsCaM-2 is 1.094 ± 0.072 versus BsCaM-45 (0.965 ± 0.102), indicating a similar dynamic range of the two sensors. The apparent affinity for \(\text{Ca}^{2+}\)-CaM (as \(K_d\)) assessed by FRET reduction for BsCaM-2 and BsCaM-45, was 2.5 ± 0.2 nM (Fig. 1C) and 46.3 ± 3.8 nM (Fig. 1D), respectively.

Detection of Biosensor Expression and FRET in Adult Ventricular Myocytes—Adult rabbit ventricular myocytes were cultured for 36–48 h to obtain sufficient expression of the biosensors without detectable cellular toxicity. The expression of both BsCaM-2 and BsCaM-45 sensors exhibited striated patterns as shown in the images of Fig. 2. To test whether this striated localization of biosensors is at Z-lines versus M-lines, myocytes were stained with the membrane indicator Di-8-Anepps to identify transverse tubules, which occur at the Z-lines (Fig. 2B).
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The line plot (Fig. 2D) of the superimposed images (Fig. 2C) shows that Di-8-Anepps and CFP signals of the biosensors overlapped spatially and that the distance between striations was about 1.9 \( \mu \text{m} \) (the sarcomere length). This suggests that the biosensors are relatively concentrated at Z-lines.

A critical hallmark of FRET is an increase in donor (CFP) fluorescence upon bleaching of the acceptor (YFP) (32). Therefore, acceptor YFP photo-bleaching experiments were performed with quiescent myocytes superfused with normal Tyrode’s solution. Fig. 2 (E–H) illustrates that photobleach of YFP (with the 514-nm laser line; note the \(~80\%\) decrease in YFP fluorescence) increased CFP donor fluorescence by \(~30\%\) in a rabbit ventricular myocyte expressing BsCaM-2. Comparable results were found in cells expressing BsCaM-45 (Fig. 2, I–L).

As shown in Fig. 2M, the mean values for increased CFP fluorescence after YFP photobleach are 32 \( \pm \) 1.3\% \((n = 8)\) and 29 \( \pm \) 1.5\% \((n = 8)\) for BsCaM-2 and BsCaM-45, respectively. Therefore, both BsCaM-2 and BsCaM-45 sensors have similar FRET efficiency in the infected myocytes.

Frequency-dependent Diastolic Ca\textsuperscript{2+}-CaM Integration in Paced Ventricular Myocytes—A unique character of Ca\textsuperscript{2+} signals in adult ventricular myocyte is the rhythmic Ca\textsuperscript{2+} oscillation occurring with each heartbeat (2). We expected that the FRET based biosensors would detect the beat-to-beat Ca\textsuperscript{2+}-CaM changes in cardiomyocytes. Indeed, using epifluorescence dual emission microscopy in field-stimulated intact adult ventricular myocytes, we found dynamic alterations of the emission of both CFP (\(540/530\)) with supplemental CaM via adenoviral infection, which was previously found to overcome the endogenous decline in CaM expression during cell culture (30). The simultaneous yet opposite changes of F\textsubscript{480/530} in cells expressing either BsCaM-2 (Fig. 3, A–C) or BsCaM-45 (Fig. 3, D–F) with supplemental CaM via adenoviral infection, which was previously found to overcome the endogenous decline in CaM expression during cell culture (30). The simultaneous yet opposite changes of F\textsubscript{480/530} in cells expressing either BsCaM-2 (Fig. 3, A–C) or BsCaM-45 (Fig. 3, D–F) with supplemental CaM via adenoviral infection, which was previously found to overcome the endogenous decline in CaM expression during cell culture (30). The simultaneous yet opposite changes of F\textsubscript{480/530} in cells expressing either BsCaM-2 (Fig. 3, A–C) or BsCaM-45 (Fig. 3, D–F) with supplemental CaM via adenoviral infection, which was previously found to overcome the endogenous decline in CaM expression during cell culture (30).
The presence of cytochalasin D, an inhibitor of contraction (33) (not shown), corroborating the dynamic FRET signal as a reliable indicator of Ca^{2+}-CaM change in the paced cells.

The frequency of Ca^{2+} signals triggered by different extracellular stimuli has been suggested to influence the resulting biologic reactions in several cell types such as neurons (22, 23) and skeletal muscle (9, 20, 24, 34). Along this line, we studied the changes of FRET signals of BsCaM-2 and BsCaM-45 in response to pacing at various rates (which drives oscillations of [Ca^{2+}]). As the pacing rate increased from 0.1 to 1 Hz, there was a clear frequency-dependent $F_{480}/F_{530}$ signal integration in BsCaM-2 sensor as shown by the significant elevation of diastolic $F_{480}/F_{530}$ levels (Fig. 3C). In contrast, there was only a tendency of diastolic $F_{480}/F_{530}$ integration of BsCaM-45 at 1 Hz (Fig. 3F). A similar slight integration of diastolic Ca^{2+} signal detected by the Fluo-4 AM Ca^{2+} indicator was observed in parallel experiments using control cells without biosensor adenoviral infection (not shown). Average diastolic Ca^{2+}-CaM signal levels of myocytes paced at increasing frequencies are summarized in Fig. 3G. The diastolic $F_{480}/F_{530}$ level of the higher affinity BsCaM-2 sensor was increased at 1 Hz (125% ± 2.9% versus 0.1 Hz, $n = 10$, $p < 0.05$), whereas the lower affinity sensor BsCaM-45 (like CaMKII) turned on and off almost completely each beat with only a slight diastolic increase at 1.0 Hz (103% ± 3.1%, versus 0.1 Hz, $n = 10$, $p > 0.05$). The BsCaM-2 amplitude of $F_{480}/F_{530}$ oscillations ($R_{\text{cytosolic}} - R_{\text{diastolic}}/ (R_{\text{max}} - R_{\text{min}})$), was ~2.5-fold larger than that of BsCaM-45 in beating myocytes (Fig. 3H), despite similar dynamic ranges of Ca^{2+}-CaM-dependent FRET ($R_{\text{max}} - R_{\text{min}}$) for the sensors (Fig. 1, C and D). As the pacing frequency increased, BsCaM-45 signal amplitude tended to increase, but not significantly so for BsCaM-2 (Fig. 3H).

Dynamic Properties of Ca^{2+}-CaM Signals Detected in Living Cardiomyocytes—To better appreciate the different dynamic properties of the biosensors in response to repetitive Ca^{2+} signals, the normalized $F_{480}/F_{530}$ signals of BsCaM-2, BsCaM-45, and Ca^{2+} transients were superimposed with expanded time scales (Fig. 4A). When cells were paced at 0.1 Hz, the Ca^{2+}-CaM signals of both BsCaM-2 and BsCaM-45 have a rising phase closely following the Ca^{2+} transient, whereas they reached the peak slower than the Ca^{2+} transient (Fig. 4C). With respect to the signal decline at 0.1 Hz, BsCaM-2 and BsCaM-45 have slower decay compared with the Ca^{2+} transient, which was demonstrated by their $\tau$ (time constant of decay, s): BsCaM-2, 2.09 ± 0.25; BsCaM-45, 0.48 ± 0.03; and Ca^{2+} transient, 0.36 ± 0.02. Therefore, at 0.1 Hz, Ca^{2+}-CaM bound to BsCaM-2 more tightly and thus presumably dissociated slower than for BsCaM-45. This was also true when the cells were paced at 1 Hz as shown in Fig. 4B, $\tau$ (s): BsCaM-2, 0.75 ± 0.07; BsCaM-45, 0.34 ± 0.01; and Ca^{2+} transient, 0.24 ± 0.04. Notably, as the pacing frequency increased from 0.1 Hz to 1.0 Hz, both BsCaM-2 and BsCaM-45 revealed an accelerated Ca^{2+}-CaM signal decay in association with the enhanced $[Ca^{2+}]_{i}$, decline (Fig. 4D).

Effects of $\beta$-Adrenergic Stimulation on the Beat-to-beat Changes of Ca^{2+}-CaM Signals—Cardiac Ca^{2+} oscillations are regulated not only by pacing frequency but also by $\beta$-adrenergic stimulation, which modulates Ca^{2+} signal amplitude and accelerates Ca^{2+} transient decline via multiple pathways, including Ca^{2+}-CaM targets (35). Therefore, the responses of biosensors with different Ca^{2+}-CaM affinities to $\beta$-adrenergic stimulation were explored by pacing cells in the presence of 1 $\mu$M Iso. The amplitude of Ca^{2+}-CaM signals of both sensors were significantly increased upon Iso stimulation at all pacing frequencies (Fig. 5, A–D, e.g. at 0.5 Hz, BsCaM-2: 186% ± 15% versus control, $p < 0.05$; BsCaM-45: 236% ± 28% versus control, $p < 0.05$). Representative $F_{480}/F_{530}$ traces showed that the frequency-dependent diastolic elevation of Ca^{2+}-CaM signal in BsCaM-2 was sustained with Iso stimulation (Fig. 5, A and B), whereas Iso-stimulated BsCaM-45 kinetics still exhibited more modest diastolic elevation at increased rates (Fig. 5, C and D). Quantitative comparisons of the kinetics of BsCaM-2, BsCaM-45, and Ca^{2+} transients at 0.5 Hz pacing rate with and without $\beta$-adrenergic stimulation are shown in Fig. 5 (E and F). In control cells, BsCaM-2 and BsCaM-45 had slower decay compared with the Ca^{2+} transient (Fig. 5E), which is consistent with our findings.
for 0.1 and 1 Hz described above. Upon application of Iso, the rates of decline of BsCaM-2, BsCaM-45, and Ca$^{2+}$/H$_{110001}$ transient were all enhanced (Fig. 5G). However, their relative positions in the superimposed tracings was still largely maintained (Fig. 5F) compared with control (Fig. 5E).

Mathematical Model of CaM Biosensor Dynamics in Ventricular Myocytes—To better understand the biochemical basis for differential activation of BsCaM-2 and BsCaM-45, we developed a mathematical model of the interaction of these biosensors with CaM (Fig. 6A), which we embedded in a previously described detailed model of Ca$^{2+}$/H$_{110001}$ handling in the adult rabbit ventricular myocyte (36).

Reactions of Ca$^{2+}$/H$_{110001}$ binding to CaM are modeled as previously (37), with sequential binding of two Ca$^{2+}$/H$_{110001}$ first to the C-terminal sites of CaM and then two Ca$^{2+}$/H$_{110001}$ to the N-terminal sites at lower affinity. The remaining reactions were based on biochemical data of CaM interactions with MLCK (whose CaM-binding domain is the linker for the biosensors used here) or when available, BsCaM-2 itself. For BsCaM-45, similar parameters are used assuming that the decreased affinity is due to an increased off rate of Ca$_4$CaM with unaltered on rate. We assume that FRET only occurs with Ca$_4$CaM bound to BsCaM. Because we overexpressed CaM in our experiments (which restores endogenous CaM levels or increases ~2-fold) (30), we also simulated conditions with an extra 6 M CaM. Simulations at either CaM level produced very similar kinetic results but with lower amplitude signals at lower [CaM]. Complete model equations and parameter values along with their sources from the biochemical literature are provided in the supplemental data. Note that none of these parameters were curve fit or adjusted to mimic the BsCaM myocyte data obtained in our experiments.

Simulations of BsCaM kinetics during 1 Hz pacing showed that BsCaM-45 FRET signal decays much more quickly than BsCaM-2 (Fig. 6B), remarkably consistent with what we
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FIGURE 7. Model-predicted diastolic integration of Ca\textsuperscript{2+}-CaM signals for BsCaM-2 and BsCaM-45 with increasing pacing rates. A, simulation of BsCaM-2 (left panels) and BsCaM-45 (right panels) kinetics in cytosolic (upper panels) and dyadic cleft (lower panels) compartments in response to increasing pacing rates from 0.1 to 1 Hz. B, comparison of cytosolic model predictions with experimental results of diastolic integration for BsCaM-2 and BsCaM-45. Diastolic integration is quantified as the magnitude of diastolic signal compared with the systolic signal at 1 Hz: \( \frac{R_{\text{dia}} - R_{\text{syst,1}}}{R_{\text{syst,1}}} \), where \( R_{\text{dia}} \) is the diastolic ratio at the indicated pacing rate, \( R_{\text{syst}} \) is the ratio without pacing, and \( R_{\text{syst,1}} \) is the systolic ratio at the pacing rate of 1 Hz.

FIGURE 8. Calibration of F-CaM binding in permeabilized adult cardiomyocytes. A, the fluorescence of different [F-CaM] in internal solution (50 nM free Ca\textsuperscript{2+}) was recorded by confocal microscopy and linear regression was shown (n = 10). B, F-CaM binding affinity in myocytes, which were permeabilized and followed by incubation of [F-CaM] in internal solution for 4–6 h. Fluorescence was taken after reaching the F-CaM binding steady status in the cells (n = 8–10), and the bound F-CaM (\( \mu \text{M} \)) was plotted according to A. Standard nonlinear regression analysis was performed to obtain the apparent \( K_d \).

observe experimentally without any curve fitting (Fig. 6, C and D). Interestingly, predicted overall decay rates of both BsCaM-2 and BsCaM-45 (1.4 and 2.3 s\textsuperscript{-1}, respectively) were significantly faster than the rate constants for Ca\textsubscript{4}CaM dissociation from BsCaM-Ca\textsubscript{4}CaM complex (0.05 and 1.5 s\textsuperscript{-1}, respectively). This indicates that Ca\textsuperscript{2+} dissociation from the complex (rather than Ca\textsubscript{4}CaM dissociation) may be the dominant BsCaM deactivation pathway in the beating myocyte. Simulations in which Ca\textsuperscript{2+} dissociation was blocked produced greatly slowed inactivation, consistent with this interpretation. Ca\textsuperscript{2+} dissociation has been previously shown to be a significant inactivation pathway in experiments and models of purified CaM targets including MLCK (38, 39), CaMKII (40), and CaN (41).

Next, we simulated BsCaM-2 and BsCaM-45 dynamics in response to a staircase of increasing pacing rates (0.1–1 Hz). BsCaM-2 exhibited significantly greater diastolic integration at increasing pacing rates compared with BsCaM-45 (Fig. 7A, upper panels), in a manner quite consistent with the experimental data (Figs. 3 and 7B) given that no curve fitting was used. This demonstrates that the biochemical mechanisms included in our model are sufficient to qualitatively predict differential signal integration and kinetics by BsCaM-2 and BsCaM-45.

However, the amplitudes of model-predicted cytosolic BsCaM signals were smaller than what was observed experimentally. Given the localization data in Fig. 2A, some BsCaM biosensors may be located in regions of high local [Ca\textsuperscript{2+}] signals (42). To determine whether BsCaM localization may explain the remaining discrepancy between model and experiment, we also simulated BsCaM-2 and BsCaM-45 in the dyadic cleft (Fig. 7A, lower panels), which predicted fully activated biosensors but with lesser diastolic integration and no gradual increase in systolic FRET ratio at higher pacing rates (Fig. 7A, lower panels). Although the experimental data is not fully explained by either BsCaM completely in the cytosol or completely in the dyadic cleft, it seems likely that some combination of these occurs. Another possible explanation for the smaller predicted amplitudes of BsCaM may be that free CaM may increase more than the expected from total CaM expression measurements (endogenous to 2-fold) (30). That is, if myocyte free CaM-binding sites are nearly saturated under basal endogenous conditions, then a doubling of total [CaM] may increase free [CaM] much more than 2-fold. This higher free [CaM] could then greatly enhance the ability of the sensors to compete for CaM in our experiments. Note that without CaM co-expression, FRET transients were not readily detectable (30).

CaM Binding in Permeabilized Ventricular Myocytes—To test the overall CaM binding affinity in cardiac myocytes, we titrated permeabilized myocytes with fluorescent CaM (F-CaM) at 50 nM free [Ca\textsuperscript{2+}], using confocal microscopy (Fig. 8). The total [CaM] in the permeabilized myocyte can be calibrated by the fluorescence intensity of pixels in the extracellular space (Fig. 8A). We previously demonstrated (15) that when
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specific CaM sites were saturated (blocked) with nonfluorescent CaM, the apparent [F-CaM] in the myocyte space was nearly the same as [F-CaM] in the bath myocyte space (within 10%) over a wide range of [F-CaM]. We used this relationship (Fig. 8A) to calibrate the [F-CaM] in images where bath [F-CaM] is known and the cellular fluorescence is measured (Fig. 8B).

At 50 nM [Ca\(^{2+}\)], the apparent \(K_a\) for CaM with nonsoluble myocyte proteins is 31 ± 5.5 nM. This indicates that at the 62–100 nM free [CaM] and 6 μM total [CaM] in ventricular myocytes (15, 30), that endogenous CaM-binding sites would be 67–76% saturated (and ~99% of total CaM is bound). If we then double the total CaM (e.g. from 6–12 μM), the endogenous sites (with 31 nM \(K_a\)) would be saturated, and free [CaM] could rise to 3–4 μM (i.e. >40-fold increase). Although it is surprising that endogenous CaM-binding sites seem near saturation at 50 nM [Ca\(^{2+}\)], this may help explain why biosensor signal amplitude is higher than expected from the model.

We also estimated the sensor concentration in myocytes using two methods. First, we used semi-quantitative Western blots of myocyte lysates versus known amounts of green fluorescent protein (correcting for two green fluorescent proteins/sensor and myocyte protein concentration, 112 mg/ml myocyte). Second, we compared the fluorescence intensity (in confocal images) of known [YFP] with YFP signals in BsCaM expressing myocytes. Both methods yielded values ~0.7 μmol/liter myocyte volume. This would add to endogenous Ca\(^{2+}\)-CaM buffering and help explain why exogenous CaM was required for robust signals.

DISCUSSION

The biosensors used in our study are recombinant proteins that mimic endogenous CaM target proteins binding to Ca\(^{2+}\)-CaM with different affinities (25). We measured FRET signals from these biosensors to assess real time interaction of CaM targets with Ca\(^{2+}\)-CaM in response to the rhythmic Ca\(^{2+}\) oscillations in adult rabbit ventricular myocytes. Our measurements demonstrated that the higher affinity sensor BsCaM-2 (like CaN) binds Ca\(^{2+}\)-CaM more tightly and therefore integrates repetitive Ca\(^{2+}\) signals in a tonic way, whereas the lower affinity sensor BsCaM-45 (like CaMKII) turns on and off more completely by binding to and dissociating from Ca\(^{2+}\)-CaM with each contraction of the ventricular myocyte (i.e. in a phasic way). We could qualitatively predict these different behaviors of BsCaM-2 and BsCaM-45 with a myocyte mathematical model using known biochemical differences in CaM affinity. Our study bridges experiments with different Ca\(^{2+}\)-CaM affinity targets and modeling and implies that the different Ca\(^{2+}\)-CaM affinities of CaM target proteins may provide an important mechanism for differential decoding of Ca\(^{2+}\) signals in myocytes. Considering the broad importance (and complexity) of CaM signaling in cardiac electrophysiology, excitation-contraction coupling, transcriptional regulation, and apoptosis, this study is a critical step in developing a more comprehensive understanding of Ca\(^{2+}\)-CaM-dependent signaling in intact cardiac myocytes.

CaM Binding Integrates at High Affinity Targets (Like CaN) upon Increased Pacing in Myocytes—We showed diastolic accumulation of Ca\(^{2+}\)-CaM bound to the higher affinity BsCaM-2 during field stimulation of myocytes (between 0.1 and 1 Hz), reflecting an integration of repetitive Ca\(^{2+}\) signals. Moreover, the frequency-dependent elevation of diastolic Ca\(^{2+}\)-CaM signal from BsCaM-2 (0.1 Hz versus 1 Hz) was also observed with β-adrenergic stimulation, a physiological stimulus that enhances the amplitude and rate of decline of the Ca\(^{2+}\) transient in myocytes (35). These findings illustrate how high affinity CaM targets like BsCaM-2 (and CaN) can integrate repetitive Ca\(^{2+}\) signals in cardiac myocytes to generate a tonic component of their signaling output. Although it is oversimplified to consider the BsCaM-2 to be an analog of CaN because the apparent Ca\(^{2+}\)-CaM affinity of CaN is ~10-fold higher than that of BsCaM-2 (17), it is reasonable to speculate that the integration of Ca\(^{2+}\) signals by CaN may be even greater (see also Ref. 37). Moreover, the agreement here between kinetics of the BsCaM-2 sensor and the computational model give us confidence that the CaN kinetics predicted in our recent modeling study (37) may be reasonable.

Our findings of CaM signal integration in cardiac myocytes are consistent with prior work on MLCK. Frequency-dependent increases in myosin light chain phosphorylation have been explained by staircasing MLCK activity arising from the kinetic properties of the CaM-MLCK complex. This was first predicted in a mathematical model (9) and later demonstrated with a MLCK biosensor in skeletal muscle (34). Frequency-dependent integration of Ca\(^{2+}\) signals at high Ca\(^{2+}\)-CaM affinity targets (such as CaN and MLCK) has been shown to be of functional importance in Ca\(^{2+}\)-dependent pathways of lymphocytes (19) and skeletal muscle (9, 20, 34). In cardiac muscle, increasing evidence has shown the significance of CaN in regulation of hypertrophy and heart failure (11), where Ca\(^{2+}\) regulation is also often disrupted (43). Interestingly, overexpression of MLCK in cardiomyocytes results in attenuation of cardiac hypertrophy (44). Our finding of Ca\(^{2+}\) signal integration caused by the high Ca\(^{2+}\)-CaM affinity of CaM targets will help develop a comprehensive picture of how cells distinguish between Ca\(^{2+}\) signaling through this pathway versus others.

CaM Binding to Low Affinity Targets (Like CaMKII) Exhibits Modest Integration—In contrast to the BsCaM-2 sensor, the BsCaM-45 sensor showed only a slight tendency toward frequency-dependent diastolic integration whether the cell was exposed to isoproterenol or not. The phasic changes of BsCaM-45 track the beat-to-beat Ca\(^{2+}\) transient much more closely than BsCaM-2. In particular, the decline of the Ca\(^{2+}\)-CaM signal of BsCaM-45 is always faster than that of BsCaM-2, although not as fast as the Ca\(^{2+}\) transient. The different kinetic properties of BsCaM-2 and BsCaM-45 are consistent with their respective Ca\(^{2+}\)-CaM binding affinities, which are similar to that of CaN and CaMKII (especially for the latter). Furthermore, the difference in the time course of BsCaM-2 and BsCaM-45 in response to the intracellular Ca\(^{2+}\) oscillation could indicate their distinctive patterns of activation in beating myocytes. Activation of CaMKII, preferentially by transient and large amplitude local Ca\(^{2+}\) spikes (21), has been implicated in CaMKII signaling at post-synaptic densities in neurons (22),
in mediating local InsP₃ receptor signaling in Ca²⁺-dependent cardiac transcriptional activation via HDAC5 nuclear export (45) and could even participate in frequency-dependent acceleration of relaxation (46, 47). Thus, local high Ca²⁺-CaM signals may preferentially signal activation of low affinity CaM targets such as CaMKII (like BsCaM-45), and this may contribute to the ability of cardiac myocytes to use multiple Ca²⁺-dependent signaling pathways for different processes in the cell simultaneously.

It should also be noted that downstream CaMKII signaling can still exhibit integration or memory, even though BsCaM-45 did not show this appreciably. This is because the affinity of CaMKII for CaM can be increased by CaMKII autophosphorylation, which may prolong the active state of CaMKII versus BsCaM-45. However, we have included that behavior in our computational model (37), and with best estimates of these CaMKII properties CaMKII seems still to activate phasically to high local [Ca²⁺], without much integration on the timescale of the heartbeat. Again our modeling and experimental data suggest that under normal conditions there is little progressive CaMKII autophosphorylation, unless the phosphatases are inhibited (37, 48). This does not preclude functional importance of CaMKII autophosphorylation in the heart, but this may be most important in pathophysiological setting like acidosis, ischemia/reperfusion (13, 49), or heart failure (50, 51). CaMKII can also phosphorylate many targets, and the lifetime and integration of those phosphorylation effects are controlled by a balance of kinase and phosphatase reactions, and these may differ among different targets and in different cellular environments (where local phosphatase and kinase activities vary). This is also true for high affinity CaM targets such as CaN, where the integrated effect of its phosphatase activity on downstream targets (e.g. NFAT) will be dictated by the kinase-phosphatase balance for a given target, location, and ambient conditions. Thus, our characterization here (and in Ref. 37) is an important initial quantitative step toward comprehensive understanding of CaM signaling in cardiac myocytes.

BsCaM Signal Amplitude May Differ Locally—The amplitude of the global BsCaM-45 signals was smaller than those of BsCaM-2 (Fig. 3H). Although the smaller BsCaM-45 signal is not surprising because of the affinity difference, it may reflect a fundamental difference in the way that lower affinity CaM targets (like CaMKII) are activated by Ca. That is, these low affinity Ca²⁺-CaM targets may only be activated in restricted compartments where local [Ca²⁺] is especially high during excitation-contraction coupling (as in the cleft and subsarcolemmal region near ryanodine receptor and L-type Ca²⁺ channels). The fact that only a minor fraction of total cellular BsCaM-45 sensor (or CaMKII) is likely to reside at the cleft or subsarcolemmal region (even if concentrated there) would be consistent with the smaller global BsCaM-45 signal amplitude in both experiments and model. Further experiments aimed at testing the model predictions for the cleft and subsarcolemmal domains would require biosensors like BsCaM-45 to be more appropriately localized and to function more like endogenous CaM targets (e.g. CaMKII).

We further speculate that our inability to detect BsCaM-45 transients without overexpression of CaM is not because signals do not occur but because they may be restricted to sensor in the cleft or subsarcolemmal (<1% of cell volume). Because this represents a small fraction of the sensor in the cell, the signals may remain below our detection sensitivity. When we overexpress CaM, this may disproportionately raise free [CaM] and increase the sensor activation level in the bulk cytosol. This increases signal amplitude to detectable levels but may artificially enhance the cytosolic signal. Clarification of this could come from the same sort of targeted sensor studies suggested above.

Limitations—Our dynamic FRET-based Ca²⁺-CaM measurements in intact ventricular myocytes uncovered differential signal integration of CaM targets because of Ca²⁺-CaM affinity. Furthermore, mechanistic computational modeling of the activation of these CaM targets reveals that the kinetic differences and diastolic integration of BsCaM-2 and BsCaM-45 myocyte signaling can be quantitatively explained by their known differences in biochemistry. These models help us identify not only what can be currently explained but also where there are limitations or gaps in our understanding. One such aspect is the difference in expected signal amplitude between model and experiments, which may be explained by altered CaM availability and important local CaM signaling (as above). With respect to CaM co-expression, our models indicate that the results are qualitatively similar in terms of kinetics regardless of CaM co-expression. Another issue is the diverse localization of endogenous CaM targets. Although our biosensors are relatively enriched at the T-tubule/Z-line (i.e. the dyadic cleft localization), the dynamic Ca²⁺-CaM signals are detected globally from the whole cell. Comparison of the amplitudes of the Ca²⁺-CaM signals in experiments versus simulations suggests that there is likely a combination of cytosolic, subsarcolemmal, and dyadic cleft, which add up to produce the cellular signals. These results point the way for future experimental studies to assess local Ca²⁺-CaM signals (e.g. with subcellular site specific targeting biosensors or imaging) and also with assessment of downstream CaM target activation (e.g. phosphorylation level of CaMKII and CaN targets). Although the BsCaM-2 and BsCaM-45 reporters used here do not share all of the properties of endogenous CaM targets like CaMKII and CaN, FRET biosensors of both CaN activity (52) and CaMKII activity (53) have recently been developed. Future iterations of these biosensors may have sufficient responsiveness for local Ca²⁺-CaM studies with endogenous CaM targets.

In conclusion, we found that BsCaM-2 and BsCaM-45, two sensors mimicking the Ca²⁺-CaM binding at the high versus low affinity CaM targets, are both capable of tracking the beat-to-beat intracellular Ca²⁺ changes, but they respond to the same Ca²⁺ signal in different ways. In particular, the BsCaM-2 (whose Ca²⁺-CaM affinity is similar to those of the high affinity CaM targets including CaN) integrates rhythmic Ca²⁺ signals, whereas the BsCaM-45 (whose Ca²⁺-CaM affinity is similar to those of the low affinity CaM targets like CaMKII) turns on and off more completely at each heartbeat. These results suggest that various CaM targets (e.g. CaN versus CaMKII) may be differentially regulated during the heart-beat via integration, thereby shaping subsequent cellular processes in the heart. Because both diastolic Ca²⁺ and heart rate can be changed in the disease state of the heart, this mechanism can differentially
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tune the downstream CaM-CaM target signaling with pathophysiologic significance.

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