Intracellular Ca dynamics in ventricular fibrillation


Intracellular Ca dynamics in ventricular fibrillation. Am J Physiol Heart Circ Physiol 286: H11006–H11021, 2004. First published January 2, 2004; 10.1152/ajpheart.00123.2003. —In the heart, membrane voltage (V_m) and intracellular Ca (Ca_i) are bidirectionally coupled, so that ionic membrane currents regulate Ca_i cycling and Ca_i affects ionic currents regulating action potential duration (APD). Although Ca_i, reliably and consistently tracks V_m at normal heart rates, it is possible that at very rapid rates, sarcoplasmic reticulum Ca_i cycling may exhibit intrinsic dynamics. Non-voltage-gated Ca_i release might cause local alternations in APD and refactoriness that influence wavebreak during ventricular fibrillation (VF). In this study, we tested this hypothesis by examining the extent to which Ca_i is associated with V_m during VF. Ca_i transients were mapped optically in isolated arterially perfused swine right ventricles using the fluorescent dye rhod 2 AM while intracellular membrane potential was simultaneously recorded either locally with a microelectrode (5 preparations) or globally with the voltage-sensitive dye RH-237 (5 preparations). Mutual information (MI) is a quantitative statistical measure of the extent to which knowledge of one variable (V_m) predicts the value of a second variable (Ca_i). MI was high during pacing and ventricular tachycardia (VT; 1.13 ± 0.21 and 1.69 ± 0.18, respectively) but fell dramatically during VF (0.28 ± 0.06, P < 0.001). Ca_i at sites 4–6 mm apart also showed decreased MI during VF (0.63 ± 0.13 compared with pacing (1.59 ± 0.34, P < 0.001) or VT (2.05 ± 0.67, P < 0.001). Spatially, Ca_i waves usually bore no relationship to membrane depolarization waves during nonreentrant fractionated waves typical of VF, whereas they tracked each other closely during pacing and VT. The dominant frequencies of V_m and Ca_i signals analyzed by fast Fourier transform were similar during VT but differed significantly during VF. Ca_i is closely associated with V_m closely during pacing and VT but not during VF. These findings suggest that during VF, non-voltage-gated Ca_i release events occur and may influence wavebreak by altering V_m and APD locally.

It is well known that the intracellular Ca (Ca_i) transient shapes cardiac action potential (AP) characteristics by modulating Ca-sensitive ionic currents, such as the L-type Ca channel, Na-Ca exchange, and Ca-activated Cl and nonsensitive channels (26). Ca_o overload also predisposes the myocardium to delayed and early afterdepolarizations, which may contribute to the initiation and perpetuation of cardiac arrhythmias. During ventricular fibrillation (VF), diastolic Ca_i may increase to systolic levels (24), potentially triggering spontaneous (i.e., nonvoltage triggered) Ca_i release from the sarcoplasmic reticulum (SR). This raises the possibility that VF may be maintained in part by effects of Ca-sensitive membrane currents on AP propagation (13). Previously, we reported that in simulated cardiac tissue, spiral waves could be destabilized and induced to breakup into a fibrillation-like state as a result of spontaneously developing spatial nonuniformities in Ca_i caused by spontaneous Ca_i-induced Ca_i release (4). If spontaneous Ca_i-induced Ca_i release was prevented, the spiral wave remained intact and did not break up. In this same study, we also demonstrated experimentally that Ca_i release in isolated cardiac myocytes exhibited intrinsic complex dynamics independent of membrane voltage (V_m). When isolated myocytes were paced at rapid rates under AP clamp conditions (so that V_m had a fixed waveform), beat-to-beat Ca_i release developed alternans and more complex periodicities, indicating that Ca_i-induced Ca_i release was being regulated by other factors besides V_m alone. We speculated that during VF, the interaction between Ca_i-release dynamics and AP dynamics might influence, and perhaps enhance, wavebreak in intact cardiac tissue.

Direct assessment of the role of Ca_i dynamics in wave stability during VF in intact tissue is difficult because V_m and Ca_i cycling are highly interdependent and bidirectionally coupled. It is not possible in intact tissue to control one process (i.e., by voltage or Ca_i clamp) to study the other. This limitation necessitates indirect approaches. A first step in understanding whether the two processes may destabilize each other is to determine whether Ca_i cycling is reliably and consistently associated with V_m during VF. If so, then it is less likely that non-voltage-gated Ca_i release occurs and that Ca_i dynamics contribute independently to wavebreak during VF. The purpose of this study was to test this hypothesis by comparing how closely Ca_i and V_m are associated with each other in arterially perfused swine right ventricles (RVs) during VF compared with pacing and ventricular tachycardia (VT).

METHODS

The research protocol was approved by the Institutional Animal Care and Use Committee and followed the guidelines of the American Heart Association.

Tissue preparation. Eleven farm pigs (weighing 25–32 kg) of either sex were anesthetized with 20 mg/kg iv thiopental sodium. The chest was opened with a median sternotomy, the hearts were removed.
and the RV was perfused through the right coronary artery as described previously (10). The isolated RV was placed with the endocardial side up in a tissue bath. In addition to continuous arterial perfusion, the entire tissue was also superfused with 37°C oxygenated Tyrode solution at a flow rate of 20 ml/min. The composition of the solution was as follows (in mmol/l): 125.0 NaCl, 4.5 KCl, 0.5 MgCl₂, 0.54 CaCl₂, 1.2 Na₂HPO₄, 24.0 NaHCO₃, and 5.5 glucose with 50 mg/l albumin; pH 7.35. Extracellular [Ca] was reduced to decrease motion artifacts because no excitation-contraction uncouplers were used to suppress contraction. Bipolar electrodes, a pacing electrode, and pseudo-ECG electrodes were attached to the endocardial surface. A pair of defibrillation coil electrodes (Guidant) were placed on either side of the tissue bath and were connected to a HVS-02 external defibrillator (Ventritex).

In all isolated RV tissues, spontaneous VT or VF occurred during the isolation procedure (presumably induced by the combination of transient ischemia and mechanical manipulation) and persisted with stable characteristics after arterial perfusion was achieved unless defibrillated. Ca⁺ transients during VT and VF were mapped while Vm was simultaneously recorded either optically or with microelectrodes. Biphasic shocks of 1.5-3.0 J were used to defibrillate the RV. The RV was then paced with electrical stimuli of 2-ms duration and twice diastolic threshold current at 400 ms.

Transmembrane AP recording. In five preparations, Vm was recorded with a standard glass microelectrode filled with 3 mol/l KCl and digitized at 3.13 kHz with 12-bit accuracy (Axon Instruments). The microelectrode was coupled with an Ag-AgCl wire leading to amplifiers with a high input impedance and variable-capacity neutralization (Am-2 and ME-3221, Warner Instruments). Microelectrode impalements were made in a region injected with the fluorescent Ca indicator rhod 2 AM to measure the Ca transient. Ca⁺ was recorded optically using a charge-coupled device (CCD) camera as described in the next section.

Optical mapping of Ca⁺ and voltage. In addition to the five preparations locally injected with rhod 2 AM, an additional five isolated, perfused swine RVs were double stained with the voltage-sensitive dye RH-237 in addition to rhod 2 AM by arterial perfusion. The double-stained heart was excited with a solid-state laser (532 nm, Verdi, Coherent), and Vm and Ca⁺ fluorescence were recorded optically by separate CCD cameras (CA-D1-0128T, Dalsa, Ontario, Canada) using a 690-nm long-pass filter for RH-237 and a 585 ± 20-nm filter for rhod 2. The two CCD cameras were carefully aligned to image the same region. To calibrate the alignment, we placed a reference grid in the optical field to provide fiduciary points, which were used to calculate the correct positions after stretching and rotational corrections. After realignment, we ascertained the positional accuracy between the two camera images to be ±1 mm. Data were acquired at an acquisition rate of 2.3 ms/frame (435 frames/s). Spatial resolution was 128 × 128 pixels over 30 × 30 mm², corresponding to 0.25 mm² tissue/pixel.

After acquisition, fluorescence signals were baseline subtracted. The RH-237 signal was inverted to show increased Vm in the upward direction. A moving median temporal filter of 5 data points (sampled at 2.3 ms/point) was applied, after which the signals at each pixel were normalized on a scale from 0 to 256 (with the average of the lowest 5 values assigned 0 and the average of the 5 maximal values assigned 256). The signal at each pixel was then spatially averaged with the signals of eight neighboring pixels to improve the signal-to-noise ratio. The overall recording area after spatial filtering was therefore estimated at 0.75 mm², with a 1-mm uncertainty between the relative positions of the Vm and Ca⁺ recording sites as noted above. Data were displayed using MATLAB (Math Works; Natick, MA), ORIGIN (Microcal Software; Northampton, MA), and customized (Wavefinder) software. In spatial maps, both Vm and Ca⁺ were color coded from red (highest voltage) to blue (lowest). Wavelets were identified using our previously described depolarization wavefront and repolarization wavebreak detection algorithm (11). Points in which depolarization and repolarization met were defined as wavebreak points. Reentry was defined as wavefront rotation around a wavebreak point completing a 360° cycle (although a stationary center of rotation was not required).

The level of cross-talk between the optically measured voltage and Ca dyes was assessed by staining preparations with only one dye and then measuring the optical signals at both wavelengths from the dual CCD cameras (Fig. 1). The absolute level of fluorescence detected in the Ca channel when the preparation was stained with RH-237 but not rhod 2 averaged <1% of the signal compared with when rhod 2 was present. Modulation of the Ca⁺ signal (i.e., the difference between maximum and minimum levels of fluorescence at each pixel) under these conditions was also <1% of the modulation in the voltage channel (Fig. 1). Conversely, with rhod 2 present but no RH-237 staining, the signal averaged <1% of the signal when RH-237 was...
present. Thus cross-talk between the optical voltage and Ca signals was insignificant.

Computer modeling. We carried out computer simulations in a two-dimensional (2-D) tissue model using the following differential equation with a no-flux boundary for transmembrane voltage ($V$):

$$\frac{\partial V}{\partial t} = -I_{\text{ion}}C_m + D\left(\frac{\partial^2 V}{\partial x^2} + \frac{\partial^2 V}{\partial y^2}\right)$$

where $t$ is time, $C_m$ is membrane capacitance, $D$ is the isotropic diffusion coefficient, and $x$ and $y$ are spatial coordinates. The ionic currents ($I_{\text{ion}}$) in the above equation were taken from two different AP models. First, we used the ventricular AP model developed by Fox et al. (8), in which we increased the maximum conductance of L-type Ca current by 50% to produce sustained spiral wave break up. In this model, the Ca transient is passively triggered by voltage. Second, we used the same modifications to the Fox et al. model but substituted a new formulation of Ca cycling developed by us. In this new model, Ca cycling is dynamically active by itself and oscillates independently of voltage in certain parameter regimes. The equations are as follows:

$$\frac{dCa}{dt} = I_{\text{rel}} + I_{\text{leak}} - I_{\text{up}} - \left(\frac{A_{\text{Ca}}Cs}{2FV_{\text{moy}}}(I_{\text{LCa}} + I_{\text{Ca}} + I_{\text{Ca}} - 2I_{\text{CaS}})\right)$$  

$$\frac{dCa_{\text{SR}}}{dt} = -\left(\frac{A_{\text{Ca}}Cs}{2FV_{\text{moy}}}(I_{\text{LCa}} + I_{\text{Ca}} + I_{\text{Ca}} - 2I_{\text{CaS}})\right)$$

$$\frac{dc}{dt} = \frac{(w_c - w)}{\tau_c}$$

$$w_c = \frac{1}{1 + k_cCa}$$

$$\tau_c = \frac{w_cCa_2}{k_{\text{off}}}$$

$$I_{\text{rel}} = \frac{v_1Ca_2}{Ca_2 + k_{\text{up}}}(Cs - Ca)$$

$$I_{\text{leak}} = \frac{v_2(Ca - Ca)}{\beta}$$

$$I_{\text{CaS}} = \frac{v_4Ca_2}{Ca_2 + k_{\text{up}}}$$

where Ca is the Ca concentration in the myoplasm, $I_{\text{rel}}$ is SR Ca release current, $I_{\text{leak}}$ is SR Ca leak current, $A_{\text{Ca}}$ is capacitive membrane area, $F$ is Faraday’s constant, $V_{\text{moy}}$ is myocardial volume, $I_{\text{LCa}}$ is L-type Ca current, $I_{\text{Ca}}$ is Ca background current, $I_{\text{CaS}}$ is SR Ca pump current, $I_{\text{rel}}$ is Ca release channel current, $w_c$ is the steady state of $w$, $\tau_c$ is the time constant for $w$, $k_c$ is a rate constant for $w$ and equals $4 \text{ mmol}^{-1} \text{sec}^{-1}$, $k_{\text{off}}$ is a rate constant for the $w$ gate and equals 0.0105 ms$^{-1}$, $v_1$ is the maximum conductance of $I_{\text{rel}}$ and equals 0.02, $k_{\text{up}}$ is a rate constant for the activation gate of $I_{\text{rel}}$ and equals 0.25 mmol, $\beta$ is a constant for the volume factor and equals 0.02, $v_2$ is the maximum conductance of $I_{\text{leak}}$ and equals $1.5 \times 10^{-4} \text{ ms}^{-1}$, and $v_4$ is the maximum conductance of $I_{\text{CaS}}$ and equals 0.1 ms$^{-1}$.

Data analysis. All data are presented as means ± SE. In the five hearts locally injected with rhod 2 AM, mutual information (MI) was used to assess the statistical dependency of Ca on $V_{\text{m}}$. MI is a nonlinear measure of the statistical dependence between two variables ($x$ and $y$); that quantifies how much knowing the value of $x$ reduces our uncertainty in the value of $y$. MI is a useful measure of the nonlinear relationship between two variables and is more sensitive than correlation, which can only find linear relationships. Given a voltage signal $V$ and a synchronous calcium signal Ca, we decimated the signals by first estimating the time it takes for the autocorrelation function of the signal to drop to zero. We then resampled the signal at an interval slightly greater than this “autocorrelation time,” thus generating statistically independent points. MI, like most measures, requires such statistically independent data points. Given a resampled time series $V_t$ and a synchronous calcium signal $Ca_n$ ($n = 1, \ldots, N$), we formed the scatterplot $\{(V_n, Ca_n)\}$, coarse grained into a $K \times K$ grid, with columns $C_1, \ldots, C_K$ and rows $R_1, \ldots, R_K$. The basic idea is that the probability of a point falling in a box $B_{ij}$ is equal to the product of its falling in row $R_i$ times the probability of its falling in column $C_j$, provided that rows and columns are independent. MI uses the discrepancy between the two as a measure of the nonindependence of row and column, i.e., voltage and calcium. The definition is as follows: let $#R_i$ be the number of points in row $R_i$, $#C_j$ the number of points in column $C_j$, and $#B_{ij}$ the number of points in box $B_{ij}$. We then define the probabilities as $P(R_i) = #R_i/N$, $P(C_j) = #C_j/N$, and $P(B_{ij}) = #B_{ij}/N$. MI is then defined as follows:

$$MI = \sum_{i,j} P(B_{ij}) \times \log \frac{P(B_{ij})}{P(R_i)P(C_j)}$$

For our case, this results in:

$$MI_{V_Ca} = \sum_{V(t),Ca(t)} P(V(t),Ca(t)) \log \frac{P(V(t)|Ca(t))}{P(V(t))P(Ca(t))}$$

where $P(V(t)|Ca(t))$ is the joint probability density for $V_m$ and Ca, resulting from values $V_m(t)$ and $Ca(t)$. $P(V(t))$ and $P(Ca(t))$ are the individual probability densities for $V_m$ and Ca. The probability densities are calculated from the data as above. Because a time lag between two variables can affect MI, we calculated MI using a range of lag times ($\tau$) from 0 to 225 ms [which exceeds the average cycle length (CL) during VT and VF]. We report the MI data by indicating the mean MI value as well as the minimum and maximum MI values over the full range of $\tau$.

The statistical significance of differences in MI among pacing, VT, and VF was assessed from the mean MI values by Student’s t-test using the Bonferroni correction for multiple comparisons. In addition, to assess whether MI values were significantly different from randomness, we used a bootstrapping method (6) in which the mean MI for the actual data was compared with the MI obtained after reshuffling the $x$, time series ($V_m$) and calculating MI for the reshuffled $x$ values and the original $y$ values ($Ca$). This procedure was iterated 100 times, and if the actual MI exceeded 95 of 100 of the MIs for the randomized time series, the MI was considered to be significant at the $P < 0.05$ level, independent of assumptions about probability distributions. MI between $V_m$ and Ca was calculated for Ca signals recorded both near ($< 1 \text{ mm}$) and far (4–6 mm) from the microelectrode $V_m$ recording site. MI between the two Ca signals at the near and far sites was also calculated to assess the spatial dependence of Ca during pacing, VT, and VF.

Fast Fourier transform (FFT) spectra were derived from optical recordings as described elsewhere (25), with the dominant frequency defined as the highest peak in the 3- to 20-Hz range of the FFT spectrum. FFT spectra were determined from a recording duration of 2.3 s in all cases. Wavebreak points in voltage maps from real and simulated tissue were detected using customized software (16).

RESULTS

Relationships between local Ca and local $V_{\text{p}}$. Figure 2, A–C, shows $V_{\text{p}}$ (black traces) and Ca transients from a heart loaded with rhod 2 AM by local injection during pacing (400-ms CL), VT, and VF, respectively. Ca was recorded both near (within 1 mm, red traces) and far (4–6 mm, blue traces) from the microelectrode $V_m$ recording site. The superimposed
traces illustrate that Cai is closely associated with \( V_m \) closely during pacing and VT but not during the majority of VF, although brief periods of synchrony were observed. During VF, not only did Cai fail to track \( V_m \), but nearby Cai signals also did not track each other closely. To quantify statistically how closely Cai at near and far sites tracked \( V_m \), MI was calculated as described in METHODS. Figure 3A shows the value of MI for simultaneous recordings of \( V_m \) and Cai during pacing, VT, and VF \( (n = 5 \text{ in each group}) \) plotted against lag time \( \tau \) (because MI is sensitive to the phase delay between the \( V_m \) and Cai traces). Figure 3B summarizes the mean value of MI for each experiment, with the minimum and maximum values indicated by the vertical bars. MI between \( V_m \) and Cai was much higher during pacing or VT than during VF, at both the near and far Cai recording sites. MI was also significantly higher during VT compared with pacing because of the longer diastolic interval during pacing. This reflects the fact that \( V_m \) and Cai have a greater influence on each other during systole than during diastole. During diastole, the decay of the Cai transient is primarily regulated by SR Ca uptake, which is independent of \( V_m \). Conversely, Cai appears to follow peak \( V_m \) because \( V_m \) is dominated by the large inward rectifier K conductance, against which Cai-sensitive currents such as Na-Ca exchange have only minor influence. Because the same relatively stable value of diastolic \( V_m \) is associated with many values of Cai, the Cai transient trails off, MI is low during diastole, decreasing the overall value of MI during the full cardiac cycle. This point reflects the robustness of MI as a sensitive quantitative measure of the extent to which Cai and \( V_m \) are associated. Moreover, this point emphasizes that the decrease in MI during VF is highly significant, because diastole is even shorter during VF, which would tend to increase MI if Cai were passively driven by \( V_m \).

To assess the degree to which MI between Cai and \( V_m \) was significant, MI for actual data was compared with MI obtained after reshuffling the \( V_m \) time series and calculating MI for the reshuffled \( V_m \) values and the original Cai values. This procedure is expected to destroy any MI that exists between the \( V_m \) and Cai signals and was iterated 100 times for each episode of pacing, VT, or VF. For each pacing and VT episode, the actual MI value was greater than the 100th percentile of all 100 shuffled MI values for both the near and far Cai recording sites. This indicates that reshuffling destroyed, at the \( P < 0.01 \) level, the MI between Cai and \( V_m \). In contrast, for VF, the actual MI between Cai and \( V_m \) at near and far sites fell at the 26th, 45th, 55th, 86th, and 97th percentile of the 100 shuffled MI values for 5 episodes of VF and between the 14th, 22nd, 52nd, 61st, and 91st percentile with respect to the far Cai site. Thus the MI of the real data during VF was not statistically different (at the \( P > 0.05 \) level) from the randomly shuffled data in all but 9 of 10 cases. This suggests that the bidirectional influence of \( V_m \) on Cai and Cai on \( V_m \) becomes highly interactive and complex during VF, consistent with the idea that non-voltage-gated Cai release occurs and has a strong influence on local \( V_m \), which affects wave propagation.

Figure 3C addresses the spatial coupling of Cai during pacing, VT, and VF by calculating MI between Cai recorded at the near \((<1 \text{ mm})\) and far \((4–6 \text{ mm})\) sites. MI was high during pacing and VT but decreased significantly during VF \((P < 0.0001)\). To estimate the spatial range over which MI decreased during VF, we compared MI among three sites that were \(<1 \text{ mm}\) apart (near), \(2–3 \text{ mm}\) apart (intermediate), or \(4–6 \text{ mm}\) (far) (Fig. 3C). For five VF episodes, the mean MI between Cai traces averaged 0.25 ± 0.01 for adjacent sites, 0.25 ± 0.01 for intermediate sites, and 0.26 ± 0.01 for far sites. Note that even for nearby sites, MI between Cai traces was significantly lower than that during either pacing or VT.

Five additional preparations were dually loaded with RH-237 and rhod 2 AM via the arterial perfusate to permit simultaneous optical mapping of \( V_m \) and Cai during episodes of pacing, VT, and VF. Figure 2, \( E \) and \( F \), shows representative simultaneous optically recorded \( V_m \) and Cai traces from the same site during VT and VF, respectively. Similar to preparations in which \( V_m \) was recorded with microelectrodes (Fig. 2, \( B–D \)), Cai was closely associated with \( V_m \) during VT (Fig. 2E) but not VF (Fig. 2F). Even though in the latter example peak Cai appears to follow peak \( V_m \), Cai appears to follow peak \( V_m \) during VF, MI was much lower than during VT. It is unlikely that the Cai signal during VF represented amplified noise, because the absolute values of fluorescence of the Cai signal before any signal processing, as well as the difference between the minimal and maximal
values, were similar during VT and VF episodes. To calculate MI between \( V_m \) and \( C_a \), four sites with good signals were selected from each quadrant of the mapped area in five episodes of VT and five episodes of VF (Fig. 3D). The mean MI between \( V_m \) and \( C_a \) averaged 0.66 ± 0.06 (range 0.31–1.28) during VT episodes and decreased to 0.28 ± 0.01 (range 0.21–0.38) during VF. Despite the wide variation in MI at different sites during VT evident in Fig. 3D, the differences between MI in VT and VF were highly significant using either the mean, maximum, or minimum values of MI (\( P < 0.0001 \) for all 3 cases). These findings are consistent with the experiments in which \( V_m \) was measured using a microelectrode. The lower average MI value during VT when \( V_m \) was recorded optically may be related to the lower signal-to-noise ratio or spatial averaging of the optical signal compared with the microelectrode \( V_m \) signal.

Finally, to confirm that non-voltage-gated \( C_a \) dynamics could account for the decrease in MI during VF, we performed simulations in 2-D homogeneous cardiac tissue using two different AP models. In one model, the \( C_a \) transient was exclusively triggered by \( V_m \) depolarization. In the other model, \( C_a \) cycling was dynamically active on its own in addition to being triggered by voltage (see METHODS). Parameters were set to the spiral wave breakup regime in both models, characterized by multiple meandering wavelets resembling VF (Fig. 4, A and B). The means ± SD of CL (168 ± 70 vs. 190 ± 93 ms), AP duration (123 ± 50 vs. 133 ± 41 ms), and diastolic interval (49 ± 56 vs. 50 ± 60 ms) were similar during simulated VF for the passive and active \( C_a \) cycling cases, respectively. Figure 4, C and D, shows representative traces of \( V_m \) and \( C_a \) at the same site, with \( C_a \) passively tracking \( V_m \) in the passive \( C_a \) cycling model (Fig. 4C) and \( C_a \) in a self-oscillatory regime in the dynamic \( C_a \) cycling model (Fig. 4D). Note the similarity of the traces in the latter case to the experimental recordings in Fig. 2C. Figure 4E shows the calculated MI between \( V_m \) and \( C_a \) as a function of \( \tau \) between the \( V_m \) and \( C_a \) traces. When \( C_a \) passively tracked \( V_m \), MI during VF was high (~1.5), but when \( C_a \) and \( V_m \) were both dynamically active, MI decreased to a much lower value (~0.5), similar to values measured during experimental VF. Moreover, the incidence of wavebreak increased significantly when \( C_a \) cycling was dynamically active, from 3.85 ± 1.89 to 7.23 ± 2.35 wavebreaks/frame (\( P < 0.01 \)). Thus localized non-voltage-gated \( C_a \) release promoted wavebreak during simulated VF in this model.

Relationship of \( C_a \) waves to membrane depolarization waves. In the five preparations dually loaded with RH-237 and rhod 2 AM, simultaneous high-resolution spatiotemporal optical mapping of \( V_m \) and \( C_a \) was performed during episodes of pacing (3 hearts), VT (3 hearts), and VF (3 hearts). During pacing (not shown) and VT (Fig. 5), \( C_a \) waves closely tracked the spread of membrane depolarization after a delay averaging 32 ± 20 ms. During the majority of VF when multiple fractionated wavefronts were present in the mapped area, \( C_a \) waves bore no consistent relationship to voltage waves (Fig. 6A). For example, at 0 ms, membrane depolarization was propagating in the upper right corner into a region in which \( C_a \) was already elevated, as though the membrane depolarization had been preceded by \( C_a \) release. Similar findings were observed in 15 nonreentrant waves during VF in 3 hearts. Similar non-voltage-gated \( C_a \)-release events were also observed in simulated VF when \( C_a \) cycling in the cell AP model was in a self-oscillatory mode (Fig. 4B, yellow oval) but not when \( C_a \) cycling was passively driven by \( V_m \) (Fig. 4A). Occasionally, however, cycles of complete reentry (360° loop) were observed during VF (~15% of all wavefronts (25)). In these cases, the \( C_a \) wave tracked the membrane depolarization wave after a brief time delay, similar to pacing and VT. Figure 6B shows an example in which the membrane depolarization wave had a center of rotation in the lower right corner and rotated clockwise. The \( C_a \) wave showed an identical pattern but delayed (at 0 ms, the membrane depolarization wave is vertical at 12 o’clock and the \( C_a \) wave is at 10 o’clock). A similar pattern...
was observed in 15 other reentrant waves during VF in 3 hearts.

Figure 7 compares the spatial dominant frequency (DF) maps of the $V_m$ and Ca signals during 2.3-s recordings of VT in five hearts and VF in five hearts. During VT (Fig. 7A), the DFs of the $V_m$ (DF$_{V_m}$) and Ca (DF$_{Ca}$) signals were closely matched over most of the mapped surface, as illustrated best in the difference maps (DF$_{V_m-Ca}$) obtained by subtracting the DF of the $V_m$ and Ca signals at each pixel. In the VT episodes analyzed, the average absolute value of DF$_{V_m-Ca}$ for all pixels was 1.07 ± 0.66 Hz. In contrast, during VF (Fig. 7B), the absolute value of DF$_{V_m-Ca}$ averaged 5.02 ± 1.22 Hz ($P = 0.00022$ compared with VT), reflecting that Ca was no longer passively tracking $V_m$ alone.

DISCUSSION

Under normal physiological conditions, regulation of contractile force by Ca in the beating heart is tightly controlled by $V_m$, such that the systolic Ca transient tracks the AP reliably and consistently with each beat. However, the Ca-induced Ca release system in the heart is an excitable subsystem capable of exhibiting its own dynamics independent of $V_m$, such as propagating Ca waves due to regenerative Ca-induced Ca release from the SR in the setting in Ca overload (3, 17, 19, 27) and from the endoplasmic reticulum in other tissues (2). In the heart, such Ca waves have even been shown to propagate intercellularly from myocyte to myocyte at 0.3–5.5 mm/s (20).

Because Ca affects $V_m$ via various Ca-sensitive ionic currents, and conversely because $V_m$ affects both Ca-release and Ca-removal processes, these two excitable systems are bidirectionally coupled. This coupling raises the possibility that under pathophysiological conditions, they may destabilize each other sufficiently to influence wavebreak (13). Computer simulations support this scenario. Chudin et al. (4) showed that a stable reentrant spiral wave became destabilized enough to break up into a fibrillation-like state when spontaneous Ca-induced Ca release was allowed to operate in the model.

The purpose of this study was to seek indirect evidence in real cardiac tissue to corroborate this scenario by examining how tightly Ca is associated with $V_m$ during VF. We show that the Ca transient tracks $V_m$ closely during pacing and VT as...
Fig. 6. Snapshots of $V_m$ and $Ca^{2+}$ during VF using the same display format as in Fig. 5. A: typical pattern during VF, with multiple $V_m$ waves and a complex distribution of wavebreak points (far right panel). The $Ca^{2+}$ wave pattern and wavebreak points bear no consistent relationship to the $V_m$ wave pattern and wavebreak points. B: episode of clockwise reentry (white arrows) during VF, which was observed only occasionally and in this example lasted for 7 rotations. The $Ca^{2+}$ wave tracked (after a delay) the reentrant $V_m$ wave, and the distribution of wavebreak points (yellow arrows) was similar.

Fig. 7. Spatial distribution of dominant frequency (DF; between 3 and 20 Hz) of optically recorded $V_m$ and $Ca^{2+}$ signals during VT (A; 5 episodes) and VF (B; 5 episodes). Top and middle, DF maps of $V_m$ (DF$_V$) and $Ca^{2+}$ (DF$_{Ca}$) on the grayscale indicated; bottom, corresponding difference DF maps (DF$_V$-DF$_{Ca}$) with red and green representing positive and negative differences, respectively.
expected, but not for the majority of time during VF. During VF, MI between simultaneously recorded traces of \( V_m \) and \( C_a \) was significantly lower than during either pacing or VT. Although MI, which quantifies the extent to which knowledge of one variable's value \( (V_m) \) predicts the value of a second variable \( (C_a) \), is a statistical measure that does not imply causality, the decreased MI during VF quantitatively indicates that the highly predictable tracking of \( C_a \) to \( V_m \) during pacing and VT is disturbed during VF, as is qualitatively apparent when traces of \( V_m \) and \( C_a \) during VF were superimposed (Fig. 2). A plausible mechanism is that the intrinsic \( C_a \) dynamics, reflected by non-voltage-gated \( C_a \)-release events, cause \( V_m \) and \( C_a \) to interact in an increasingly less predictable fashion. This conjecture is further directly supported by simultaneous spatiotemporal maps of \( V_m \) and \( C_a \). These maps showed that when fractionated multiple wavefronts were present during VF, \( C_a \) waves bore no consistent relationship to membrane depolarization, as they did during pacing and VT. With multiple fractionated wavefronts, \( C_a \) waves could either precede or follow membrane depolarization, suggesting that both AP-dependent and voltage-independent \( C_a \) release events were occurring. Only occasionally during VF, when complete reentrant cycles occurred in the mapped area \([\sim 15\% \text{ of the time (25)}]\), did \( C_a \) waves closely track membrane depolarization (Fig. 6). The analysis of DFs of \( V_m \) and \( C_a \) signals during VF also showed much greater differences during VF than during VT. By virtue of the bidirectional coupling between \( C_a \) and \( V_m \), these observations indicate that \( C_a \) dynamics influence wave behavior in a complex manner during VF. However, they do not prove that this complex mutualism is an independent cause of wavebreak during VF. It is just as possible that increased spatial \( C_a \) heterogeneity due to non-voltage-gated \( C_a \) release could prevent wavebreak as promote it. However, in the computer simulation, the net effect of non-voltage-gated \( C_a \) release was to increase the incidence of wavebreak.

Possible mechanisms underlying dissociation of \( C_a \) and \( V_m \) during VF. The mechanism by which \( C_a \) is no longer associated with \( V_m \) in a consistent and reliable fashion during multiple wavefront VF remains speculative, but several possibilities exist. At a general level, theoretical studies have shown examples in which feedback between two or more coupled excitable systems induces unstable chaotic behavior (7, 14, 21). Wavebreak promoted by \( C_a \)-induced \( C_a \) release could be an example of this, because the AP and \( C_a \)-induced \( C_a \) release are coupled excitable systems each capable of exhibiting complex dynamics.

At the cellular level, we speculate that the failure of \( C_a \) to faithfully track \( V_m \) during VF might be explained by spontaneous \( C_a \)-induced \( C_a \) release becoming dominant over \( C_a \) release triggered by the L-type \( C_a \) current during the AP. Because \( C_a \) transients due to spontaneous \( C_a \)-induced \( C_a \) release are localized events that propagate only slowly from myocyte to myocyte [0.3–5.5 mm/s (20)], this speculation is consistent with our observation that during VF, \( C_a \) transients at sites even <1 mm apart had a significant drop in MI compared with \( C_a \) transients during pacing or VT (Fig. 3C).

\( C_a \) overload is known to potentiate \( C_a \)-induced \( C_a \) release and also to predispose hearts to VF. Clusin et al. (5) reported that \( C_a \) overload induced VF-like activity in aggregates of cultured chick myocytes, which could be prevented by \( C_a \) channel blockers. In intact rabbit hearts in which VF was induced either electrically or by \( C_a \) overload, Merillat et al. (18) found that \( C_a \) channel blockers or removal of extracellular \( C_a \) abolished VF. However, ryanodine, which depletes SR \( C_a \) (by locking \( C_a \)-release channels into an open subconductance state so that the SR becomes leaky) and prevents \( C_a \) oscillations, did not prevent VF [see also Kusuoka et al. (12)]. Lippi and Billman (9) also found that spontaneous VF induced by \( C_a \) overload was prevented by ryanodine, whereas VF induced by ischemia was not. Saito et al. (23) reported that \( C_a \) dynamics are the main cause of electrical and mechanical alternans in ventricular muscle, and alternans is well known to predispose the heart to VF (22). In summary, these observations suggest that, although intact \( C_a \) cycling is not essential for the maintenance of VF in all settings, it exerts a significant proarrhythmic effect by enhancing initiation of VF by promoting triggered activity. Our simulations suggest that spatially heterogeneous \( C_a \) may also promote maintenance of VF by increasing the incidence of wavebreak (13).

Although technical limitations prevented us from measuring absolute changes in restig \( C_a \) levels in this study, it was shown previously that resting \( C_a \) can increase significantly to systolic levels during VF, depending on the pharmacological setting (24). This previous study in rat and hamster hearts also documented two types of \( C_a \) transients during VF, with either minimal (type 1) or preserved (type 2) amplitude modulation. In our experiments in the pig heart, we only observed type 2 \( C_a \) transients during VF. However, the \( C_a \) signal in the prior study was spatially averaged over a much larger area (38.5 mm²) than in our study (0.45 mm²), so that the lack of modulation of the \( C_a \) transient during type 1 VF may have reflected spatial averaging of locally dysynchronous \( C_a \) signals. This is supported by our observation that \( C_a \) transients at sites <1 mm apart had a significant drop in mutual information compared with pacing or VT (Fig. 3B).

Study limitations. There are several limitations to this study. The finding that \( C_a \) is no longer reliably and consistently associated with \( V_m \) during VF does not directly address the issue of whether \( C_a \) dynamics promote or inhibit wavebreak in VF or whether modifying \( C_a \) dynamics would have antifibrillatory effects. MI is a statistical measure of the relationship between two variables and, if high, indicates an association, but, like all purely statistical measures, does not imply causality. Neither do the simultaneous \( V_m \) and \( C_a \) maps prove that \( C_a \) cycling promotes or prevents additional wavebreaks during VF. Nevertheless, it is interesting to note that in multiple wavefront VF, regional elevations in \( C_a \) often preceded membrane depolarization (Fig. 6A). This suggests that \( C_a \) release was being triggered regionally by non-voltage-gated mechanisms. In our simulations, the net effect of non-voltage-gated \( C_a \) release due to dynamically active \( C_a \) cycling was to increase the incidence of wavebreak, but whether this is also true in real tissue is unproven. Our study also does not shed light into details of the underlying cellular mechanisms: for example, the roles of early or delayed afterdepolarizations, intracellular or intercellular \( C_a \) waves, intercellular \( C_a \) diffusion, or refractoriness of SR \( C_a \)-release channels; the relative importance of specific \( C_a \)-sensitive ionic currents that couple \( C_a \) to \( V_m \); or the role of regional tissue heterogeneities in AP
and Ca-release properties that might promote wavebreak under these conditions.

Our studies were conducted at reduced extracellular [Ca] (0.54 mM) but in the absence of excitation-contraction uncouplers such as diacetyl monoxime, raising the possibility that contraction artifacts may have contaminated the optical signals. Even if a motion artifact contaminated the optical Ca signal, however, there was still very high MI between Vm and “Ca,”. It is difficult to imagine that a contaminating contraction motion would increase MI between the optical Ca trace and the microelectrode Vm signal. During VF, contraction (and therefore potential contamination of optical signals by motion) is minimal, yet the MI between Vm and Ca decreased compared with pacing or VT. This finding indicates that the calculation of MI is fairly robust and makes it unlikely that the decrease in MI during VF was related to motion artifacts. The reduced extracellular [Ca] would be expected, if anything, to ameliorate Ca overload during VF and therefore inhibit the uncoupling of Ca from Vm. However, MI between Vm and Ca still decreased during VF.

Finally, VF was studied in isolated arterially perfused ventricles, whereas VF in in vivo hearts quickly leads to superimposed acute ischemia. Thus our findings are most relevant to the initial phase of VF in the setting of either chronic heart disease or electrically induced VF in the normal heart before acute ischemia sets in. The MI between Vm and Ca during VF may closely resemble that found during VF in the setting of either chronic heart disease or electrically induced VF in the normal heart before acute ischemia sets in. The findings may not apply to VF induced by acute ischemia or other settings (13).

In conclusion, Ca is closely associated with Vm closely during pacing and VT but not during the majority of time in VF. The failure of Ca to passively track Vm during VF may alter local refractoriness and therefore influence wavefronts affecting the maintenance of VF. Strategies to prevent of VF will need to consider Ca cycling dynamics in addition to other factors such as cardiac electrical restitution properties.

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