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Mechanisms Underlying Heterogeneous Ca²⁺ Sparklet Activity in Arterial Smooth Muscle

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In arterial smooth muscle, single or small clusters of Ca²⁺ channels operate in a high probability mode, creating sites of nearly continual Ca²⁺ influx (called “persistent Ca²⁺ sparklet” sites). Persistent Ca²⁺ sparklet activity varies regionally within any given cell. At present, the molecular identity of the Ca²⁺ channels underlying Ca²⁺ sparklets and the mechanisms that give rise to their spatial heterogeneity remain unclear. Here, we used total internal reflection fluorescence (TIRF) microscopy to directly investigate these issues. We found that tsA-201 cells expressing L-type Cav α 1.2 channels recapitulated the general features of Ca²⁺ sparklets in cerebral arterial myocytes, including amplitude of quantal event, voltage dependencies, gating modalities, and pharmacology. Furthermore, PKC α activity was required for basal persistent Ca²⁺ sparklet activity in arterial myocytes and tsA-201 cells. In arterial myocytes, inhibition of protein phosphatase 2A (PP2A) and 2B (PP2B; calcineurin) increased Ca²⁺ influx by evoking new persistent Ca²⁺ sparklet sites and by increasing the activity of previously active sites. The actions of PP2A and PP2B inhibition on Ca²⁺ sparklets required PKC activity, indicating that these phosphatases opposed PKC-mediated phosphorylation. Together, these data unequivocally demonstrate that persistent Ca²⁺ sparklet activity is a fundamental property of L-type Ca²⁺ channels when associated with PKC. Our findings support a novel model in which the gating modality of L-type Ca²⁺ channels vary regionally within a cell depending on the relative activities of nearby PKC α , PP2A, and PP2B.

INTRODUCTION

During the myogenic response (Bayliss, 1902), smooth muscle lining the walls of resistance arteries respond to increased intravascular pressure by undergoing gradual depolarization, thus increasing the open probability of dihydropyridine-sensitive, voltage-gated L-type Ca²⁺ channels located in the sarcolemma of arterial smooth muscle cells (Harder et al., 1987; Fleischmann et al., 1994; Rubart et al., 1996; Knot and Nelson, 1998). Increased opening of L-type Ca²⁺ channels causes greater Ca²⁺ influx that culminates in a global rise in intracellular Ca²⁺ ([Ca²⁺]_i) and arterial constriction. Accordingly, by regulating the activity of L-type Ca²⁺ channels, smooth muscle cells are able to modulate [Ca²⁺]_i, arterial diameter, and therefore blood flow.

We recently observed single and small clusters of seemingly coupled Ca²⁺ channels (presumably L-type) operating in a high activity gating mode that created local areas of nearly continual Ca²⁺ influx termed “persistent Ca²⁺ sparklet” sites (Navedo et al., 2005). On the basis of these findings it was proposed that steady-state Ca²⁺ influx in arterial smooth muscle occurs through persistent Ca²⁺ sparklet sites in combination with random, infrequent openings of solitary L-type Ca²⁺ chan-

nels (Fleischmann et al., 1994; Rubart et al., 1996). At present, however, this “persistent Ca²⁺ sparklet model” remains largely untested.

The goal of this study was to address four fundamental, yet unresolved, issues raised by this provocative model. First, we investigated the molecular identity of the channels that underlie persistent Ca²⁺ sparklets. This is of particular importance because the conclusion that persistent Ca²⁺ sparklets are produced by the opening of single, or small clusters, of L-type Ca²⁺ channels was based largely on pharmacological evidence (i.e., sensitivity to dihydropyridines), which is equivocal. Second, we identified the minimal molecular components required for persistent Ca²⁺ sparklet activity. Third, we investigated the mechanisms underlying dynamic, regional variations in Ca²⁺ sparklet activity. Fourth, we examined the molecular identities of the signaling molecules involved in the regional modulation of persistent Ca²⁺ activity.

Our data indicate that expression of PKC α and L-type Ca²⁺ (Cav α 1.2) channels was sufficient to reproduce the basic features of persistent Ca²⁺ sparklet activity in a

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The online version of this article contains supplemental material.

Abbreviations used in this paper: CsA, cyclosporine A; OA, okadaic acid; PDBu, phorbol 12, 13-dibutyrate; PKC β i, PKC β inhibitor; PP2A, protein phosphatase 2A; TIRF, total internal reflection fluorescence; WT, wild-type.

heterologous expression system. This provides the first direct demonstration that Cav α 1.2 channels underlie persistent Ca $^{2+}$ sparklets in smooth muscle. Accordingly, our data indicate, for the first time, that persistent Ca $^{2+}$ sparklet activity is a fundamental feature of L-type Ca $^{2+}$ channels (with PKC α), which suggests the intriguing possibility that persistent Ca $^{2+}$ sparklets may be a general mechanism underlying steady-state Ca $^{2+}$ entry in excitable cells. Finally, our data support the novel concept that subcellular compartmentalization of Ca $^{2+}$ influx via L-type Ca $^{2+}$ channels is determined by the local balance between PKC α and opposing phosphatase (protein phosphatase 2A and 2B) activities.

MATERIALS AND METHODS

Isolation of Arterial Myocytes

Rats (Sprague-Dawley; \approx 250 g) as well as wild type and PKC α knockout mice (\approx 25 g) (Braz et al., 2004) were euthanized in strict accordance to the regulations of the University of Washington Institutional Animal Care and Use Committee using a lethal dose of sodium pentobarbital (100 mg/kg, intraperitoneally). Myocytes were dissociated from cerebral arteries using standard enzymatic techniques described in detail elsewhere (Amberg and Santana, 2003). After dissociation, cells were maintained in a nominally Ca $^{2+}$ -free Ringer's solution until used. Thapsigargin (1 μ M) was included in all solutions to eliminate Ca $^{2+}$ release from intracellular stores during experimentation.

Heterologous Expression of Cav α 1.2 and PKC α in tsA-201 Cells

Cultures of tsA-201 cells were maintained in Dulbecco's modified essential Media supplemented with 10% fetal bovine serum, L-glutamine (2 mM), and a 1% streptomycin and penicillin solution. Cells were transiently transfected with the pcDNA clones of Cav α 1.2, Cav β 3, Cav α 2 δ 1 (a gift from D. Lipscombe, Brown University, Providence, RI), and the enhanced green fluorescent protein using Lipofectamine 2000. In some experiments, tsA-201 cells were transfected with Cav α 1.2 and accessory subunits as well as PKC α tagged with the enhanced green fluorescent protein (provided by J. Exton, Vanderbilt University, Nashville, TN). Successfully transfected cells were identified on the basis of enhanced green fluorescent protein fluorescence.

Electrophysiology

We used the conventional whole-cell patch-clamp technique to control membrane voltage using an Axopatch 200B amplifier. During experiments, cells were continuously superfused with a solution with the following constituents (in mM): 140 NMDG, 5 CsCl, 1 MgCl $_2$, 10 glucose, 10 HEPES, and 2 or 20 CaCl $_2$ adjusted to pH 7.4. NMDG concentration was 120 mM when 20 mM CaCl $_2$ was used. Pipettes were filled with a solution composed of (in mM) 87 Cs-aspartate, 20 CsCl, 1 MgCl $_2$, 5 MgATP, 10 HEPES, 10 EGTA, and 0.2 Fluo-5F or Rhod-2 adjusted to pH 7.2 with CsOH. A voltage error of 10 mV attributable to the liquid junction potential was corrected for. In some experiments, Ca $^{2+}$ currents were recorded and later analyzed using pCLAMP 9.0 software. In these experiments, currents were sampled at 20 kHz and low pass filtered at 2 kHz. All experiments were performed at room temperature (22–25°C).

Total Internal Reflection Fluorescence (TIRF) Microscopy

Ca $^{2+}$ sparklets were recorded using a through-the-lens TIRF microscope built around an inverted Olympus IX-70 microscope equipped with an Olympus PlanApo (60X, numerical aperture =

1.45) oil-immersion lens and an XR Mega 10 intensified CCD camera (Solamere Technology Group). To monitor [Ca $^{2+}$] $_i$, cells were loaded with the calcium indicators Fluo-5F or Rhod-2. Rhod-2 was used in all experiments in which the enhanced green fluorescent protein was expressed. Excitation of Fluo-5F and Rhod-2 was achieved with the 488- or 568-nm line of an argon or krypton laser, respectively (Dynamic Lasers). Excitation and emission light was separated with the appropriate set of filters. Images were acquired at 30–90 Hz.

Background-subtracted fluorescence signals were converted to concentration units using the "F $_{max}$ " equation (Maravall et al., 2000):

$$[Ca^{2+}] = K_d \frac{F/F_{max} - 1/R_f}{1 - F/F_{max}},$$

where F is fluorescence, F $_{max}$ is the fluorescence intensity of Fluo-5N or Rhod-2 in the presence of a saturating free Ca $^{2+}$ concentration, K $_d$ is the dissociation constant of the fluorescence indicator used (Fluo-5N = 1100 nM; Rhod-2 = 600 nM), and R $_f$ (Fluo-5N = 210; Rhod-2 = 150) is this indicator's F $_{max}$ /F $_{min}$. F $_{min}$ is the fluorescence intensity of Fluo-5N or Rhod-2 in a solution where the Ca $^{2+}$ concentration is 0. K $_d$ and R $_f$ values for Fluo-5N and Rhod-2 were determined in vitro using standard methods (Woodruff et al., 2002). F $_{max}$ was determined at the end of the experiments by exposing cells to a solution to which the Ca $^{2+}$ ionophore ionomycin (10 μ M) and 20 mM external Ca $^{2+}$ had been added.

Ca $^{2+}$ sparklets were detected and defined for analysis using an automated algorithm written in IDL language. Ca $^{2+}$ sparklets had an amplitude equal to or larger than the mean basal [Ca $^{2+}$] $_i$ plus three times its standard deviation. For a [Ca $^{2+}$] $_i$ elevation to be considered a sparklet, a grid of 3 \times 3 contiguous pixels had to have a [Ca $^{2+}$] $_i$ value at or above the amplitude threshold. These detection criteria for Ca $^{2+}$ sparklets are similar to those used by other investigators (Cheng et al., 1999; Demuro and Parker, 2004, 2005).

By simultaneously recording single Ca $^{2+}$ channel currents and Ca $^{2+}$ sparklets in arterial myocytes, we recently reported that at -70 mV and with 20 mM external Ca $^{2+}$, a single Ca $^{2+}$ channel current of \approx 0.5 pA produced a Ca $^{2+}$ sparklet of \approx 37 nM (Navedo et al., 2005). As shown in Fig. S1 (see online supplemental material, available at <http://www.jgp.org/cgi/content/full/jgp200609519/DC1>), an "all-points" histogram from representative [Ca $^{2+}$] $_i$ records obtained from arterial myocytes had multiple, clearly separated peaks and could be fit with the following multi Gaussian function:

$$N = \sum_{j=1}^n a_j * \exp \left[-\frac{([Ca^{2+}]_i - jq)^2}{2jb} \right],$$

where a and b are constants and [Ca $^{2+}$] $_i$ and q are intracellular Ca $^{2+}$ and the quantal unit of Ca $^{2+}$ influx, respectively. Using this analysis, and consistent with our previous study, we obtained a q value of 34 nM in the all-points histogram, a value that is similar to that obtained previously (see below). This analysis provides further support to the hypothesis that Ca $^{2+}$ sparklets are quantal in nature and that the size of Ca $^{2+}$ sparklet depends on the number of quanta activated.

Analogous to single-channel data analysis, we determined the activity of Ca $^{2+}$ sparklets by calculating the nP $_s$ of each sparklet site, where n is the number of quantal levels and P $_s$ is the probability that a quantal Ca $^{2+}$ sparklet event is active. To do this, we used the single channel analysis module of pCLAMP 9.0. First, [Ca $^{2+}$] $_i$ from previously identified sparklet sites were imported into this program and a baseline defined. To estimate nP $_s$, Ca $^{2+}$ sparklet events were detected using pCLAMP's "threshold detection

analysis” using no duration constraints and a unitary Ca^{2+} elevation of 38 nM as a starting point for event detection (note that the amplitude of the unitary event was not fixed). Traces were then fitted with these initial parameters. Each one of the events detected with this analysis were then cross-referenced with the original image stack to verify that they met the amplitude and spatial criteria described above. Only Ca^{2+} influx events that met the spatial and amplitude criteria were used to estimate nP_s for each experimental condition. An example of this type of analysis is shown in the online supplemental material (Fig. S2).

Amplitude histograms were constructed using the amplitudes of the detected Ca^{2+} sparklet events. The resulting histogram was fitted with the multicomponent Gaussian function described above, which allowed us to obtain an estimate of the amplitude of quantal Ca^{2+} sparklets under varied experimental conditions. It is important to note that the observation of multiple peaks and similar q values in our all-points and event histograms (e.g., Figs. 1–3 and Fig. S1) indicate that the use of the event histogram for our quantal analysis is appropriate.

As previously reported (Navedo et al., 2005), Ca^{2+} sparklet activity was bimodal in arterial myocytes and tsA-201 cells expressing $\text{Cav}\alpha 1.2$ and $\text{PKC}\alpha$, with sites of low activity ($nP_s = 0.07 \pm 0.01$) and sites of high activity ($nP_s = 0.73 \pm 0.7$). Based on this behavior, we grouped Ca^{2+} sparklets into three categories; silent (by default has an nP_s of 0), low (nP_s between 0 and 0.2), and high (nP_s higher than 0.2). Note that a silent Ca^{2+} sparklet site represents a site that is ordinarily inactive, but can be activated by an agonist.

Chemicals and Statistics

All PKC inhibitors were acquired from Calbiochem. Cell culture media and supplements were from Life Technologies. Lipofectamine 2000 was purchased from Invitrogen; all other chemicals were from Sigma-Aldrich. Data are presented as mean \pm SEM. Two-sample comparisons were made using Student’s t test. For datasets containing more than two groups, an ANOVA was used. A P value of <0.05 was considered significant. The asterisk (*) symbol used in the figures denotes a significant difference between groups.

Online Supplemental Material

The online supplemental material (Figs. S1 and S2, available at <http://www.jgp.org/cgi/content/full/jgp.200609519/DC1>) provides an example of our Ca^{2+} sparklet activity analysis.

RESULTS

$\text{Cav}\alpha 1.2$ Channels Produce Ca^{2+} Sparklets

All experiments in this study were performed in the presence of the SERCA pump inhibitor thapsigargin (1 μM) to eliminate Ca^{2+} release from intracellular stores. TIRF microscopy was used to image Ca^{2+} sparklet events with high spatial and temporal resolution. To increase the driving force for Ca^{2+} entry, experiments were performed in the presence of 20 mM external Ca^{2+} while cells were held at -70 mV. Analogous to single-channel data analysis, we determined the activity of Ca^{2+} sparklets by calculating the nP_s of each sparklet site as described in the Materials and Methods.

First, we tested the hypothesis that L-type Ca^{2+} channels underlie Ca^{2+} sparklet activity in arterial myocytes. If L-type Ca^{2+} channels underlie Ca^{2+} sparklets, then Ca^{2+} sparklets should be observed in a heterologous expression system expressing these channels. Thus, we

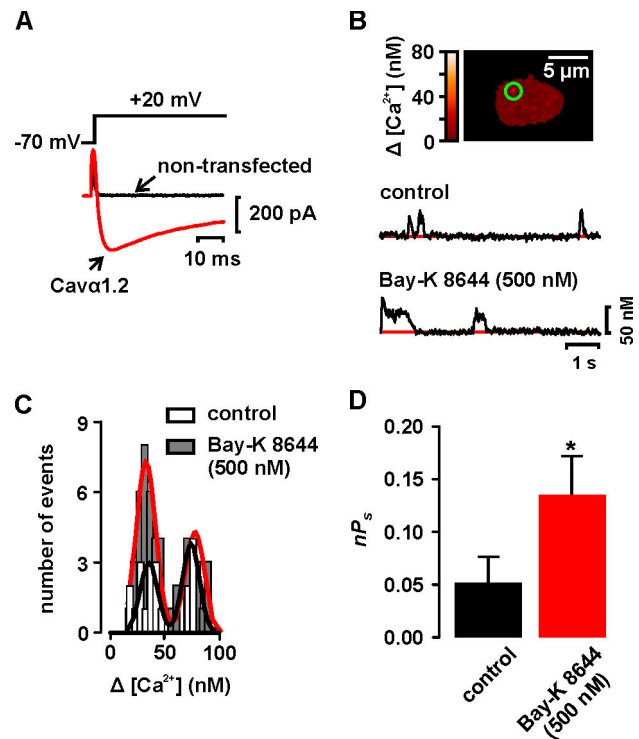


Figure 1. Ca^{2+} sparklets in a heterologous expression system. (A) Ca^{2+} current evoked by a depolarization from -70 to $+20$ mV in a representative nontransfected (control) cell and cell expressing $\text{Cav}\alpha 1.2$. (B) Image of a cell transfected with $\text{Cav}\alpha 1.2$ only. The traces below the images show the time course of $[\text{Ca}^{2+}]_i$ in the sites marked by the green circle before and after the application of 500 nM Bay-K 8644. (C) Amplitude histogram of Ca^{2+} sparklets in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ before and after Bay-K 8644 treatment. The black and red lines are the best fit to the control ($q = 37$ nM) and Bay-K 8644 ($q = 37$ nM) data, respectively, with the multicomponent Gaussian function described in the Materials and Methods section. (D) Bar plot of the mean \pm SEM of the nP_s before and after Bay-K 8644 application.

examined Ca^{2+} sparklet activity in tsA-201 cells expressing the L-type Ca^{2+} channel pore-forming $\text{Cav}\alpha 1.2$ and accessory $\text{Cav}\beta 3$ and $\text{Cav}\alpha 2\delta 1$ subunits. $\text{Cav}\alpha 1.2$ channels are the predominant L-type Ca^{2+} channels expressed in arterial smooth muscle (Koch et al., 1990; Sinnegger-Brauns et al., 2004). To prevent any potential effect of endogenous PKC activity on $\text{Cav}\alpha 1.2$ function, these experiments were performed using the PKC inhibitory peptide (100 μM) in the pipette solution.

To verify that tsA-201 cells transfected with $\text{Cav}\alpha 1.2$, $\text{Cav}\beta 3$, and $\text{Cav}\alpha 2\delta 1$ expressed functional channels, we depolarized these cells from -70 to $+20$ mV. As shown in Fig. 1 A, this protocol evoked robust Ca^{2+} currents in these cells. Ca^{2+} currents were not detected in nontransfected cells ($n = 25$). In addition, in tsA-201 cells expressing $\text{Cav}\alpha 1.2$, Ca^{2+} sparklets were observed at -70 mV (Fig. 1 B). Ca^{2+} sparklets were never observed in nontransfected cells ($n = 25$). As shown in the amplitude histogram in Fig. 1 C, the distribution of

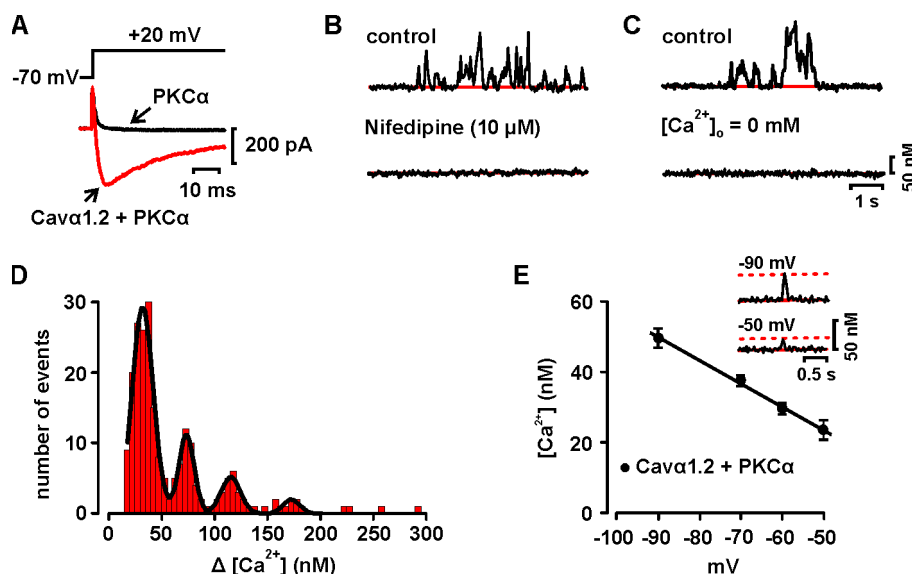


Figure 2. Ca^{2+} sparklets in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ and $\text{PKC}\alpha$. (A) Ca^{2+} current evoked by a depolarization from -70 to $+20$ mV in representative cells expressing $\text{PKC}\alpha$ or $\text{PKC}\alpha$ and $\text{Cav}\alpha 1.2$. (B) Time course of $[\text{Ca}^{2+}]_i$ in Ca^{2+} sparklet site before and after the application of $10 \mu\text{M}$ nifedipine. (C) Time course of $[\text{Ca}^{2+}]_i$ in Ca^{2+} sparklet site under control conditions and after perfusion with a solution without Ca^{2+} . (D) Amplitude histogram of Ca^{2+} sparklets in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ and $\text{PKC}\alpha$. The black line is the best fit to the data ($q = 36$ nM) with the multicomponent Gaussian function described in the Materials and Methods section. (E) Voltage dependence of quantal Ca^{2+} sparklet amplitude in tsA-201 cells. Solid line is a linear fit to the data. The inset shows representative quantal Ca^{2+} sparklets recorded from tsA-201 cells expressing $\text{Cav}\alpha 1.2$ channels at -90 and -50 mV. The dotted lines mark the quantal level for Ca^{2+} sparklets at -90 and -50 mV.

Ca^{2+} sparklet amplitudes in tsA-201 cells was modal. Indeed, the data could be fit with a multicomponent Gaussian function with a quantal unit of Ca^{2+} elevation of 37.9 nM ($\chi^2 = 0.76$). Interestingly, the amplitude of quantal Ca^{2+} sparklets in these cells is similar to that reported in arterial myocytes (≈ 38 nM) under identical experimental conditions (Navedo et al., 2005). This analysis suggests that, as with Ca^{2+} sparklets recorded in arterial myocytes, Ca^{2+} entry via heterologously expressed $\text{Cav}\alpha 1.2$ channels is quantal in nature and that the size of Ca^{2+} sparklet depends on the number of quanta activated.

Detailed analysis revealed that the activity of Ca^{2+} sparklets sites in these cells had a mean nP_s value of 0.05 ± 0.02 ($n = 9$) (Fig. 1 D). This nP_s value is similar ($P > 0.05$) to that of low activity Ca^{2+} sparklet sites reported in rat arterial myocytes (see also new data from rat and mouse myocytes presented below) (Navedo et al., 2005). Indeed, it is important to note that high activity, persistent Ca^{2+} sparklet sites ($nP_s > 0.2$) were never observed in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ only.

In rat arterial myocytes, Bay-K 8644 increases Ca^{2+} influx, at least in part, by increasing persistent Ca^{2+} sparklet activity (Navedo et al., 2005). However, the mechanisms by which Bay-K 8644 increases persistent Ca^{2+} sparklet activity are unclear. We investigated whether Bay-K 8644 could induce persistent Ca^{2+} sparklet activity in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ (Fig. 1, B–D). Application of 500 nM Bay-K 8644 recruited new

Ca^{2+} sparklet sites and increased the activity of previously active sites. Indeed, Bay-K 8644 increased the average Ca^{2+} sparklet activity (i.e., nP_s) from 0.05 ± 0.03 to 0.13 ± 0.04 ($n = 6$, $P < 0.05$; Fig. 1 D) without increasing the amplitude of quantal Ca^{2+} sparklets (control = 37.0 nM vs. Bay-K 8644 = 37.6 nM; Fig. 1 C). Interestingly, even in the presence of 500 nM Bay-K 8644 we did not detect high nP_s , persistent Ca^{2+} sparklet sites in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ only. These data suggest that although $\text{Cav}\alpha 1.2$ channel can produce low activity Ca^{2+} sparklets sites, expression of these channels alone is not sufficient to produce persistent Ca^{2+} sparklet activity under control conditions or after Bay-K 8644 treatment.

$\text{Cav}\alpha 1.2$ and $\text{PKC}\alpha$ Are Required for Persistent Ca^{2+} Sparklet Activity

We recently observed that PKC activity is required for persistent Ca^{2+} sparklet activity in rat arterial myocytes (Navedo et al., 2005). Thus, we examined Ca^{2+} sparklet activity in tsA-201 cells expressing $\text{Cav}\alpha 1.2$ channels and $\text{PKC}\alpha$ (Fig. 2). We used $\text{PKC}\alpha$ in these experiments because this isoform is highly expressed in cerebral artery myocytes (but see below) (Pang et al., 2002; Wickman et al., 2003). Depolarization from the holding potential of -70 mV to $+20$ mV evoked large Ca^{2+} currents in tsA-201 expressing $\text{Cav}\alpha 1.2$ channels and $\text{PKC}\alpha$ (Fig. 2 A). Ca^{2+} currents were not observed in cells expressing $\text{PKC}\alpha$ alone ($n = 28$).

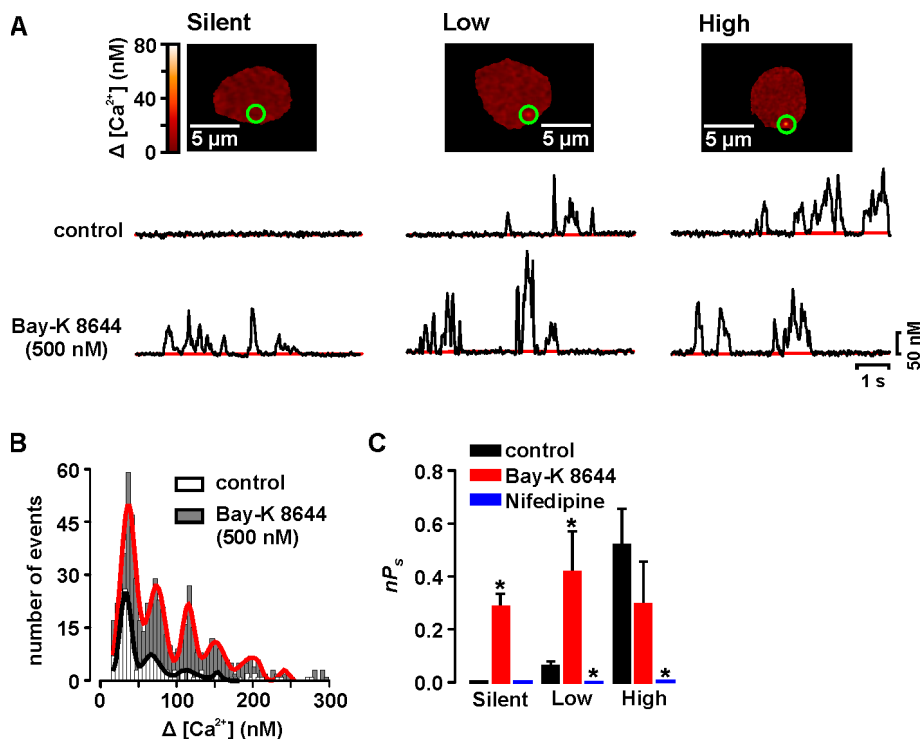


Figure 3. Cav α 1.2 and PKC α are required for persistent Ca $^{2+}$ sparklet activity. (A) Image of tsA-201 cells expressing PKC α and Cav α 1.2 with silent, low, and high activity Ca $^{2+}$ sparklet sites. The traces below the images show the time course of [Ca $^{2+}$] $_i$ in the sites marked by the green circle before and after the application of 500 nM Bay-K 8644. (B) Amplitude histogram of Ca $^{2+}$ sparklets in tsA-201 cells expressing Cav α 1.2 and PKC α before and after Bay-K 8644 treatment. The black and red lines are the best fit to the control ($q = 38$ nM) and Bay-K 8644 ($q = 37$ nM) data, respectively, with a multi-component Gaussian function. (C) Bar plot of the mean \pm SEM nP_s under control conditions and after application of Bay-K 8644 or nifedipine.

Accordingly, Ca $^{2+}$ sparklet sites were frequently observed at -70 mV in tsA-201 cells expressing Cav α 1.2 channels and PKC α , but never in cells expressing PKC α alone ($n = 28$). To provide further support to the hypothesis that Ca $^{2+}$ sparklets are produced by Ca $^{2+}$ influx via Cav α 1.2 channels and not Ca $^{2+}$ release from a thapsigargin-insensitive intracellular Ca $^{2+}$ store in tsA-201 cells, we examined the effects of the L-type Ca $^{2+}$ channel blocker nifedipine, which blocks sparklets in arterial myocytes (Navedo et al., 2005), and removing external Ca $^{2+}$ on Ca $^{2+}$ sparklets. As shown in Fig. 2 B, nifedipine completely eliminated Ca $^{2+}$ sparklet activity ($n = 15$). Furthermore, we found that perfusion of an external solution with 0 Ca $^{2+}$ (without nifedipine) rapidly eliminated Ca $^{2+}$ sparklet activity in tsA-201 cells expressing Cav α 1.2 channels ($n = 10$; Fig. 2 C). Together with our observation that Ca $^{2+}$ sparklets are only detected in tsA-201 cells expressing Cav α 1.2, these findings support the hypothesis that Ca $^{2+}$ sparklets in tsA-201 cells are produced by Ca $^{2+}$ influx events through plasma membrane Cav α 1.2 channels.

An amplitude histogram of Ca $^{2+}$ sparklets in tsA-201 cells expressing Cav α 1.2 channels and PKC α is shown in Fig. 2 D. The histogram was fitted ($\chi^2 = 0.84$) with a multicomponent Gaussian function with a quantal unit of Ca $^{2+}$ elevation of 36.0 nM, a value that is similar to that of cells expressing Cav α 1.2 alone (see Fig. 1 C above). We also investigated the effects of membrane potential on the amplitude of quantal Ca $^{2+}$ sparklets in tsA-201 cells expressing Cav α 1.2 and PKC α . The amplitude of quantal Ca $^{2+}$ sparklets at the voltages examined

was obtained from the event amplitude histogram of all Ca $^{2+}$ sparklets using the multi-Gaussian analysis described above. The inset in Fig. 2 E shows two [Ca $^{2+}$] $_i$ records from a representative cell with quantal Ca $^{2+}$ sparklets at -90 and -50 mV. Note that the amplitude of these Ca $^{2+}$ sparklets decreased as the driving force for Ca $^{2+}$ entry was decreased by membrane depolarization from -90 to -50 mV. Indeed, as previously reported in rat arterial myocytes (Navedo et al., 2005), the amplitude of quantal Ca $^{2+}$ sparklets in tsA-201 cells decreased linearly over this range of potentials (Fig. 2 E), providing further support to the view that Ca $^{2+}$ sparklets are produced by Ca $^{2+}$ influx via L-type Cav α 1.2 channels in the plasma membrane.

We analyzed the modalities of Ca $^{2+}$ sparklet activity in tsA-201 cells expressing Cav α 1.2 and PKC α (Fig. 3, A and C). Although most of the surface membrane in these cells did not show signs of Ca $^{2+}$ influx at -70 mV (i.e., $nP_s = 0$), there were sites of low and high Ca $^{2+}$ sparklet activity (Fig. 2, C and D). Because cells expressing Cav α 1.2 alone did not have high nP_s sites, the combined nP_s value of cells expressing Cav α 1.2 and PKC α (0.16 ± 0.03 , $n = 12$) was about threefold higher than in cells expressing Cav α 1.2 alone (0.05 ± 0.02 , $n = 6$, $P < 0.05$). These data suggest that Ca $^{2+}$ sparklets in tsA-201 expressing Cav α 1.2 and PKC α and rat arterial myocytes have similar gating modalities.

The effects of Bay-K 8644 on Ca $^{2+}$ sparklets in tsA-201 cells expressing Cav α 1.2 and PKC α were also investigated (Fig. 3). Application of 500 nM Bay-K 8644 increased the number of Ca $^{2+}$ sparklet sites threefold

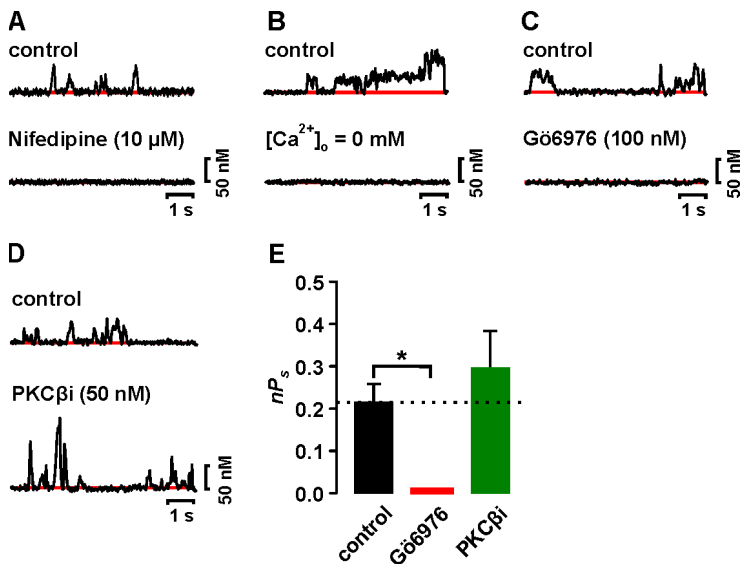


Figure 4. Basal PKC β and PKC ϵ activity is not required for Ca²⁺ sparklet activity in arterial smooth muscle. (A) Time course of $[Ca^{2+}]_i$ in a Ca²⁺ sparklet site before and after the application of 10 μ M nifedipine. (B) Time course of $[Ca^{2+}]_i$ in a Ca²⁺ sparklet under control conditions and after perfusion of a solution without Ca²⁺. Time course of $[Ca^{2+}]_i$ in a Ca²⁺ sparklet site before and after the application of (C) 100 nM Gö6976 or (D) 50 nM PKC β i. (E) Bar plot of the mean \pm SEM nP_s under control conditions and after application of Gö6976 or PKC β i. *, significantly different from control.

($n = 8$). Consistent with this, Bay-K 8644 activated quiescent Ca²⁺ channels in silent sites and increased Ca²⁺ sparklet activity in low nP_s sites ($P < 0.05$), thus increasing the number of high nP_s persistent Ca²⁺ sparklets in tsA-201 cells (Fig. 3, A–C). Bay-K 8644 did not increase Ca²⁺ sparklet activity in high nP_s sites ($P > 0.05$), suggesting maximal channel activity at these sites. Bay-K 8644 increased the number of Ca²⁺ sparklets of all amplitude levels without altering the value of the quantal event (38.4 nM; Fig. 3 B). Importantly, the actions of Bay-K 8644 on Ca²⁺ sparklets in these cells are similar to those reported in arterial myocytes (Navedo et al., 2005). Taken together, these data indicate that Ca²⁺ sparklets in tsA-201 cells expressing Cav α 1.2 and PKC α have similar pharmacology, gating modalities, amplitude of quantal event, and voltage dependencies than sparklets in arterial myocytes. This is consistent with the view that Cav α 1.2 and PKC α are the minimal molecular components required for persistent Ca²⁺ sparklet activity under control conditions and after Bay-K 8644 treatment.

Dynamic Modulation of Ca²⁺ Sparklet Activity Depends on the Relative Activities of PKC α and Opposing Phosphatases

Next, we investigated the mechanisms that underlie dynamic regional variations of persistent Ca²⁺ sparklet activity in arterial myocytes. We tested the hypothesis that regional differences in Ca²⁺ sparklet activity result from regional differences in the relative activities of PKC, which we have shown here to be essential for persistent Ca²⁺ sparklet activity in tsA-201 cells, and nearby opposing phosphatases. To begin, we investigated which of the PKC isoforms expressed in cerebral arterial smooth muscle are required for persistent Ca²⁺ sparklet activity. Although the experiments described above indicate that

expression of PKC α and Cav α 1.2 channels is sufficient to reproduce the basic features of Ca²⁺ sparklets in tsA-201 cells, two recent studies indicate that cerebral artery myocytes express three PKC isoforms: Ca²⁺-dependent PKC α and β and, to a lesser extent, the Ca²⁺-independent PKC ϵ isoform (Pang et al., 2002; Wickman et al., 2003). We used isoform-specific PKC inhibitors to determine which of these three PKC isoforms influence persistent Ca²⁺ sparklet activity in cerebral arterial myocytes.

Like the experiments in tsA-201 cells, all experiments with arterial myocytes were performed with solutions containing the SERCA pump inhibitor thapsigargin (1 μ M) to eliminate Ca²⁺ release from intracellular stores. To verify that Ca²⁺ sparklets were not produced by Ca²⁺ release from an intracellular store insensitive to thapsigargin, Ca²⁺ sparklets were recorded in rat arterial myocytes under control conditions (i.e., 20 mM external Ca²⁺ while the cells were held at -70 mV), in the presence of the dihydropyridine nifedipine and after perfusion of a solution without Ca²⁺. As shown in Fig. 4 A, Ca²⁺ sparklets in arterial myocytes were completely abolished by the application of the dihydropyridine nifedipine (10 μ M; $n = 7$). Furthermore, note that Ca²⁺ sparklets were rapidly abolished by superfusion of a Ca²⁺-free solution (without nifedipine) (Fig. 4 B). Similar results were obtained in 10 independent experiments. Together, these findings indicate that Ca²⁺ sparklets are produced by Ca²⁺ influx via a sarcolemma L-type Ca²⁺ channel in arterial smooth muscle.

Fig. 4 (A and B, top traces) shows the time course of $[Ca^{2+}]_i$ in a Ca²⁺ sparklet site from a representative rat arterial myocyte under control conditions. An amplitude histogram of Ca²⁺ sparklets recorded under these conditions could be fit with a multi-Gaussian function with a quantal unit of Ca²⁺ increase of 37.5 nM (not depicted), which is similar to that in tsA-201 cells

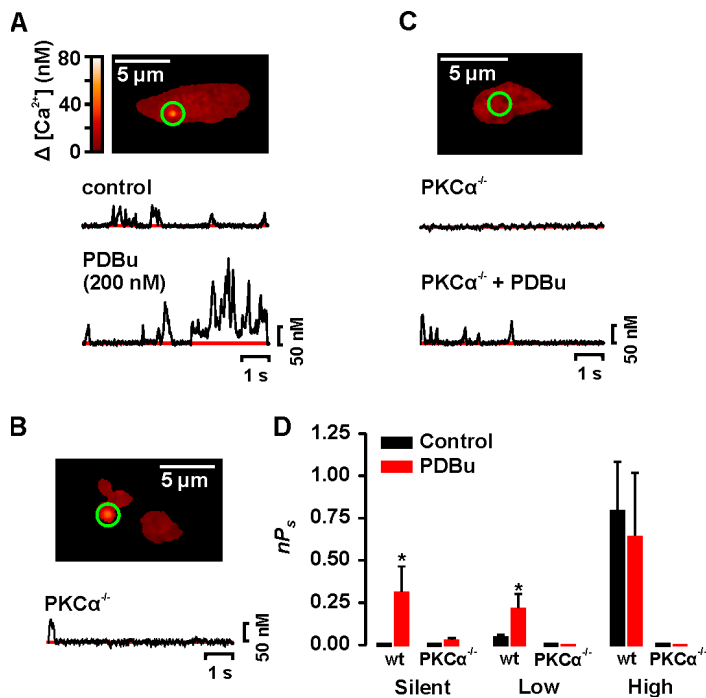


Figure 5. PKC α is required for Ca²⁺ sparklets in arterial smooth muscle. Sample images of typical WT (A) and PKC α ^{-/-} (B and C) mouse arterial myocytes. The traces below the images (A and C) show the time course of [Ca²⁺]_i in the sites marked by the green circle before and after the application of 200 nM PDBu. (D) Bar plot of the mean \pm SEM nP_s under control conditions and after application of PDBu. *, significantly different from control.

(see Fig. 1 C and Fig. 2 D above). We found that the majority of the sarcolemma of rat myocytes was optically silent (i.e., $nP_s = 0$). However, we observed foci of low ($nP_s = 0.06 \pm 0.01$, $n = 75$) and high ($nP_s = 0.7 \pm 0.1$, $n = 58$) Ca²⁺ sparklet activity. The average nP_s value of Ca²⁺ sparklet sites under control conditions was 0.17 ± 0.05 ($n = 110$). Interestingly, nP_s values in rat arterial myocytes and tsA-201 cells (low $nP_s = 0.05 \pm 0.01$, high $nP_s = 0.6 \pm 0.01$, average $nP_s = 0.14 \pm 0.04$) expressing Cav α 1.2 and PKC α were similar ($P > 0.05$).

Application of Gö6976 (100 nM), which selectively inhibits PKC α and PKC β (Gschwendt et al., 1996), eliminated (i.e., $nP_s = 0$, number of sparklet sites = 0) Ca²⁺ sparklet activity in rat arterial myocytes (Fig. 4, C–E). This suggests that the Ca²⁺-dependent PKC α and/or PKC β isoforms underlie spontaneous persistent Ca²⁺ sparklet activity in rat arterial myocytes. To distinguish which of these two isoforms underlie spontaneous Ca²⁺ sparklet activity in these cells, we recorded Ca²⁺ sparklets before and after the application of a specific PKC β inhibitor (PKC β i, 50 nM) (Tanaka et al., 2004). In contrast to Gö6976, the averaged Ca²⁺ sparklet activity did not change after PKC β i application (control $nP_s = 0.21 \pm 0.04$, $n = 15$ vs. PKC β i $nP_s = 0.29 \pm 0.08$, $n = 15$; $P > 0.05$) (Fig. 4, D and E). In addition, we found that Ca²⁺ sparklet activity and the number of Ca²⁺ sparklet sites per cell were not changed by dialysis with a specific PKC ϵ inhibitory peptide (PKC ϵ i) (Johnson et al., 1996) (unpublished data, $n = 5$ cells, $P > 0.05$). These data indicate that neither PKC β nor PKC ϵ activity is required for spontaneous Ca²⁺ sparklet activity in arterial smooth muscle.

By excluding PKC β and PKC ϵ , our data suggest that PKC α is required for Ca²⁺ sparklet activity in arterial myocytes. To directly test this hypothesis, we examined Ca²⁺ sparklet activity in mouse wild-type (WT) and PKC α knockout (PKC α ^{-/-}) (Braz et al., 2004) arterial myocytes. Fig. 5 (A and C) shows that mouse arterial myocytes produce Ca²⁺ sparklets. Indeed, it is important to note that Ca²⁺ sparklets in WT mouse myocytes were similar ($P > 0.05$) to those in tsA-201 cells and rat arterial myocytes in all parameters examined (i.e., amplitude of the quantal events, 38 nM, and activity modalities). Control mouse-WT myocytes had silent (i.e., $nP_s = 0$) as well as low ($nP_s = 0.08 \pm 0.02$, $n = 35$) and high activity ($nP_s = 0.80 \pm 0.30$, $n = 15$) Ca²⁺ sparklet sites. Like rat arterial myocytes (Navedo et al., 2005), application of the broad-spectrum PKC activator phorbol 12, 13-dibutyrate (PDBu, 200 nM) increased Ca²⁺ influx in mouse arterial myocytes by activating new Ca²⁺ sparklet sites and by increasing the activity of previously active sites (unpublished data). Indeed, PDBu increased the number of Ca²⁺ sparklet sites in mouse arterial myocytes twofold ($n = 7$). Thus, persistent Ca²⁺ sparklets and their modulation by PKC appears to be a conserved feature of arterial myocytes.

Consistent with our data from rat arterial myocytes and tsA-201 cells, we found that PKC α ^{-/-} myocytes were devoid of persistent Ca²⁺ sparklet activity under control conditions ($n = 20$ cells). Indeed, in only 1 out of 20 cells examined was a single Ca²⁺ sparklet event (amplitude = 38 nM) evident at -70 mV. Interestingly, application of 200 nM PDBu, which would activate other PKC isoforms expressed in PKC α ^{-/-} cells, had a small effect

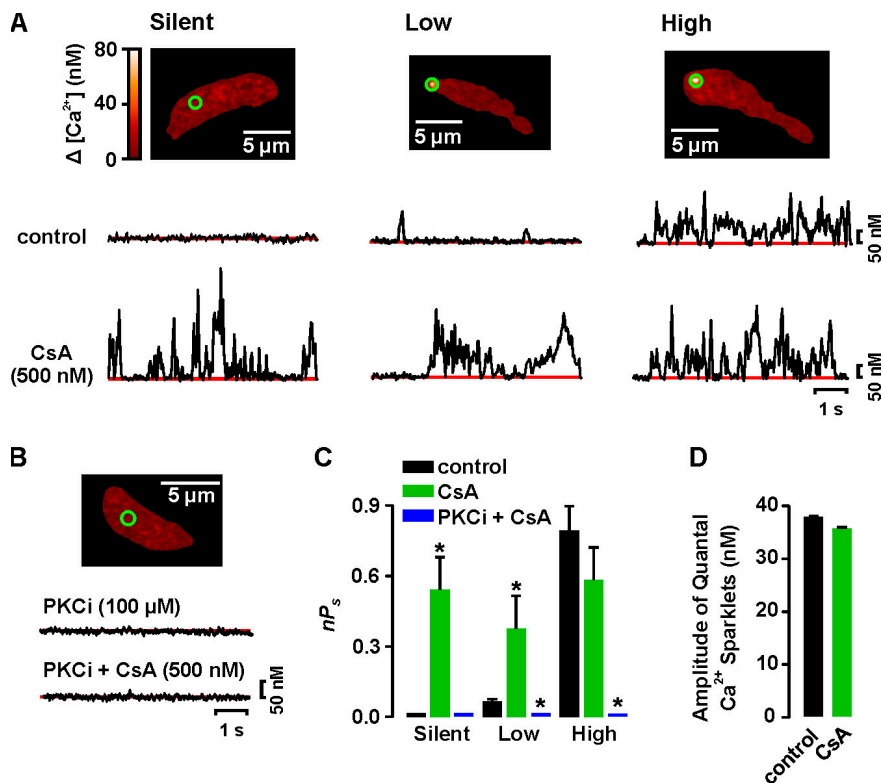


Figure 6. PP2B opposes PKC-dependent activation of Ca^{2+} sparklets. (A) Images of representative cells with silent (left), low (center), and high (right) nP_s sites. Traces below the images show the time course of $[Ca^{2+}]_i$ in the green circle before (top) and after (bottom) application of 500 nM cyclosporine A (CsA). (B) Sample image from a cell dialyzed for 10 min with an internal solution containing PKCi (100 μ M) before and after CsA. The time course of $[Ca^{2+}]_i$ in the region marked by the green circle is shown below the image. (C) Mean \pm SEM nP_s of Ca^{2+} sparklet sites in control conditions and in the presence of CsA or PKCi and CsA. (D) Mean \pm SEM amplitude of quantal Ca^{2+} sparklets before and after application of CsA. *, significantly different from control.

on Ca^{2+} sparklet activity in these cells, only activating a few, low nP_s Ca^{2+} sparklet sites ($nP_s = 0.02 \pm 0.01$, $n = 21$; Fig. 5, B–D). In total, three Ca^{2+} sparklet sites were observed in $PKC\alpha^{-/-}$ cells in the presence of PDBu. In $PKC\alpha^{-/-}$ cells, Ca^{2+} sparklet amplitudes ranged from 34 to 41 nM. Note that this range of amplitudes is similar to the amplitude of quantal Ca^{2+} sparklets in WT mouse and rat myocytes and tsA-201 cells, indicating that the probability of coincidental openings of nearby L-type Ca^{2+} channels in $PKC\alpha^{-/-}$ cells was low. Indeed, the effects of PDBu on Ca^{2+} sparklet activity in $PKC\alpha^{-/-}$ cells ($nP_s \approx 0.02 \pm 0.01$, $n = 21$) were 15-fold smaller than in WT cells ($nP_s = 0.30 \pm 0.1$, $n = 15$) (Fig. 5 D). Taken together, these data support the hypothesis that basal $PKC\alpha$ activity is necessary for spontaneous Ca^{2+} sparklet activity. Furthermore, our data suggest that regional variations in the activity of $PKC\alpha$ underlie heterogeneous Ca^{2+} sparklet activity in arterial myocyte.

Having established that basal $PKC\alpha$ activity is necessary for spontaneous persistent Ca^{2+} sparklet activity in arterial myocytes, we investigated the role of protein phosphatases in modulating the activity of these Ca^{2+} influx events. Recent studies have suggested that the serine/threonine phosphatases PP1, PP2A, and PP2B modulate L-type Ca^{2+} channel function (Santana et al., 2002; duBell and Rogers, 2004). Thus, we tested the hypothesis that local variations in the relative activities of $PKC\alpha$ and opposing protein phosphatases determine regional variations in Ca^{2+} sparklet activity in arterial myocytes.

First, we examined the role of PP2B (calcineurin) on Ca^{2+} sparklets (Fig. 6). Inhibition of PP2B with cyclosporine A (CsA; 500 nM) increased Ca^{2+} influx by activation of previously silent sites and by increasing the activity of low activity sites (Fig. 6, A and C). CsA did not increase the activity of high nP_s sites, suggesting maximal activity at these sites (Fig. 6, A and C). CsA induced a twofold increase in the number of Ca^{2+} sparklet sites per cell ($n = 12$, $P < 0.05$). The quantal amplitude of Ca^{2+} sparklets was unchanged by CsA (Fig. 6 D). It is important to note that CsA failed to activate Ca^{2+} sparklets in cells dialyzed with the PKC inhibitor PKCi (Fig. 6 B). These results suggest that PP2B dampens Ca^{2+} sparklet activity by opposing PKC-mediated phosphorylation.

Next, we investigated the effects of the PP1 and PP2A inhibitor calyculin A (100 nM) (duBell et al., 2002) on Ca^{2+} sparklets. Like CsA, calyculin A increased Ca^{2+} influx by activating previously silent Ca^{2+} sparklet sites and by increasing the activity of low nP_s sites (Fig. 7, A and C). Again, the activity of high nP_s sites did not change upon application of calyculin A, suggesting maximal activity at these sites. Calyculin A induced a 2.3-fold increase in the number of Ca^{2+} sparklets sites per cell ($n = 7$, $P < 0.05$). As with CsA, calyculin A did not activate Ca^{2+} sparklets in cells dialyzed with PKCi (Fig. 7 B). These results suggest that PP1 and/or PP2A modulate Ca^{2+} sparklets by opposing PKC.

We used different concentration of okadaic acid (OA) to determine the relative contribution of PP1 and PP2A to Ca^{2+} sparklet activity (Fig. 8). At concentrations

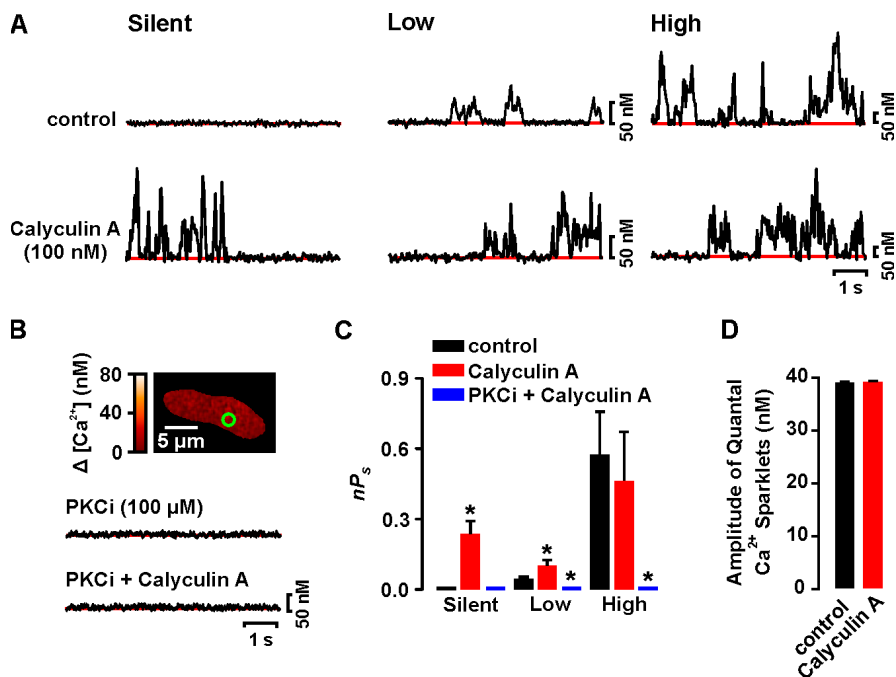


Figure 7. Calyculin A increases Ca^{2+} sparklet activity. (A) Time course of $[Ca^{2+}]_i$ in silent (left), low (center), and high (left) nP_s sites from a representative cell before (top) and after (bottom) application of 100 nM calyculin A. (B) Sample image from a typical cell dialyzed for 10 min with an internal solution containing PKCi (100 μ M). The traces below show the time course of $[Ca^{2+}]_i$ in the region of the cell marked by the green circle before (top trace) and after (bottom trace) calyculin A. (C) Mean \pm SEM nP_s before and after application of calyculin A or PKCi plus calyculin A. (D) Mean \pm SEM amplitude of quantal Ca^{2+} sparklets before and after application of calyculin A. *, significantly different from control.

of between 1 and 10 nM, OA inhibits PP2A, while at concentrations >100 nM, it blocks PP2A and PP1 (Bialojan and Takai, 1988; duBell and Rogers, 2004). Application of 1 nM OA increased Ca^{2+} sparklet activity nearly twofold (Fig. 8 A). It is important to note that the increase in Ca^{2+} sparklet activity produced by 1 nM OA was not significantly different from that observed with calyculin A ($P > 0.05$). At 1 nM, OA induced a 2.5-fold increase in the number of Ca^{2+} sparklets sites per cell ($n = 9$, $P < 0.05$). These data suggest that calyculin A increased Ca^{2+} sparklet activity by inhibiting PP2A. Consistent with this hypothesis, increasing OA in the bath from 1 to 100 nM, thus also inhibiting PP1, did not elicit any further increase in Ca^{2+} sparklet activity ($nP_s = 0.37 \pm 0.1$, $n = 31$, $P > 0.05$) or sparklet density ($n = 9$, $P > 0.05$). As with CsA and calyculin A, OA did not activate Ca^{2+} sparklets in cells dialyzed with PKCi and did not change the quantal level of the Ca^{2+} sparklets (Fig. 8). From these data we conclude that PP2A, PP2B, and PKC α form a signaling module that tunes local Ca^{2+} influx via L-type Ca^{2+} channels in arterial smooth muscle cells.

DISCUSSION

We used TIRF microscopy to investigate with high resolution the spatiotemporal organization of functional Ca^{2+} channels. Using this approach, we provide the first direct demonstration that persistent Ca^{2+} sparklet activity is a fundamental property of L-type Ca^{2+} channels when associated with PKC. Furthermore, we describe a novel mechanism for the local, dynamic control of steady-state Ca^{2+} influx via L-type Ca^{2+} channels. We

found that PKC α and the phosphatases PP2A and PP2B (calcineurin) have opposing effects on persistent Ca^{2+} sparklet activity. Our results also suggest that PP2A and PP2B oppose the actions of PKC α on L-type Ca^{2+} channels. Based on these findings we propose that the degree of steady-state Ca^{2+} influx in various regions of the cell via L-type Ca^{2+} channels is determined by a local balance between PKC α , PP2A, and PP2B activities.

Our data, in conjunction with earlier studies (Nelson et al., 1995), suggest that smooth muscle cells are capable of generating multiple types of local Ca^{2+} signals. For example, simultaneous activation of a small cluster of ryanodine-sensitive Ca^{2+} channels in the sarcoplasmic reticulum of these cells produces Ca^{2+} sparks (Nelson et al., 1995). Because Ca^{2+} sparks are produced by the release of Ca^{2+} from intracellular stores, they are insensitive to dihydropyridines or the removal of external Ca^{2+} , their amplitude is independent of changes in membrane voltage, and they are abolished by depleting the SR of Ca^{2+} with thapsigargin (Cannell et al., 1995; López-López et al., 1995; Nelson et al., 1995). In sharp contrast to Ca^{2+} sparks, the Ca^{2+} sparklets described here meet all the criteria for a Ca^{2+} influx event via sarcolemmal Ca^{2+} channels. First, Ca^{2+} sparklets are insensitive to thapsigargin. Second, Ca^{2+} sparklets are rapidly eliminated by perfusion with a Ca^{2+} -free solution. Third, the amplitude of quantal Ca^{2+} sparklets decreased with membrane depolarization (i.e., decreased driving force). Fourth, a dihydropyridine antagonist and agonist inhibited and activated Ca^{2+} sparklets, respectively. Taken together, these data provide compelling support to the hypothesis that Ca^{2+} sparklets are produced by Ca^{2+} influx via sarcolemma Ca^{2+} channels.

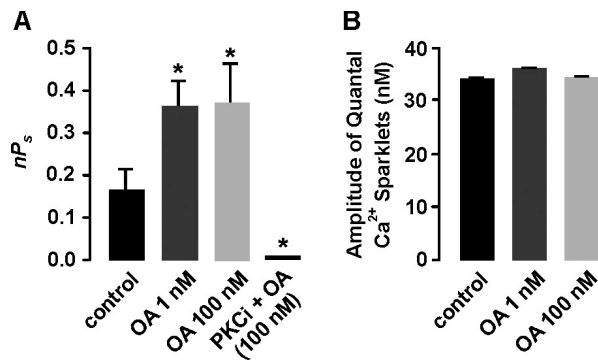


Figure 8. PP2A inhibits Ca^{2+} sparklets by opposing PKC. (A) Mean \pm SEM nP_s before and after application of 1 or 100 nM okadaic acid (OA) in the absence or presence of PKCi in the patch pipette. (B) Mean \pm SEM amplitude of quantal Ca^{2+} sparklets before and after application of 1 nM or 100 nM OA. *, significantly different from control.

Our data unequivocally demonstrate that L-type $Cav\alpha 1.2$ channels have the ability to produce persistent Ca^{2+} sparklets even at hyperpolarized potentials (-70 mV). Interestingly, L-type $Cav\alpha 1.2$ channels require PKC α activity to operate in a persistent gating mode; in the absence of PKC α activity, Ca^{2+} sparklet activity in tsA-201 cells and arterial myocytes at -70 mV was very low. An important finding in our study is that heterologous expression of PKC α and $Cav\alpha 1.2$ channels was sufficient to reproduce all the basic features (i.e., amplitude of quantal event, voltage dependencies, gating modalities, and pharmacology) of persistent Ca^{2+} sparklet activity in arterial myocytes. These results strongly support the hypothesis that, as in heart (Wang et al., 2001), L-type Ca^{2+} channels underlie Ca^{2+} sparklets in arterial smooth muscle.

The data presented here, and in our previous study (Navedo et al., 2005), indicate that Ca^{2+} sparklets in arterial smooth muscle are produced by the opening of a single or a cluster of L-type Ca^{2+} channels. Accordingly, simultaneous recordings of single Ca^{2+} channel currents and Ca^{2+} sparklets under conditions similar to those used in the current study (i.e., -70 mV and 20 mM external Ca^{2+}) demonstrated that an opening of a single Ca^{2+} channel (0.5 pA) could produce a Ca^{2+} sparklet of an amplitude of ≈ 37 nM (Navedo et al., 2005). Because Ca^{2+} sparklet amplitude is variable (between 38 and 300 nM), these data strongly suggest that Ca^{2+} sparklets could be produced by the opening of a single or a cluster of L-type Ca^{2+} channels. Consistent with this, the all-points (see Fig. S1, available at <http://www.jgp.org/cgi/content/full/jgp.200609519/DC1>) and event histograms (e.g., Fig. 1 C, Fig. 2 D, and Fig. 3 B) presented here clearly show well-defined peaks with a quantal unit of Ca^{2+} influx of ~ 34 – 37 nM. Together, these data provide compelling support to the view that openings of a single or a cluster of L-type Ca^{2+} channels underlie Ca^{2+} sparklets in arterial smooth muscle.

Although Ca^{2+} sparklets in cardiac and smooth muscle are produced by the same molecular entity (hence the same name), there are important differences between Ca^{2+} sparklets in these cells. For example, as noted above, Ca^{2+} sparklets in arterial myocytes are produced by the opening of a single or a cluster of L-type $Cav\alpha 1.2$ channels. In contrast, Ca^{2+} sparklets in ventricular myocytes are produced by the opening of a single L-type Ca^{2+} channel (Wang et al., 2001). Furthermore, unlike smooth muscle, persistent Ca^{2+} sparklet activity has not been observed in ventricular myocytes. This is interesting because PKC α is expressed in ventricular myocytes (Braz et al., 2004) and $Cav\alpha 1.2$ is the predominant $Cav\alpha$ isoform in these cells. Although the reasons for these differences are presently unclear, it is intriguing to speculate that basal PKC α activity in ventricular myocytes is lower than in arterial myocytes, thus decreasing the probability of persistent Ca^{2+} sparklet activity in these cells. This hypothesis is supported by our data suggesting that Ca^{2+} sparklet activity in tsA-201 expressing $Cav\alpha 1.2$ only and PKC $\alpha^{-/-}$ myocytes, as in ventricular myocytes, is low and mostly resulting from the activation of single $Cav\alpha 1.2$ channels. Future studies should examine the mechanisms modulating Ca^{2+} sparklet activity in heart.

A recent study (Yang et al., 2005) provides insight into the molecular mechanisms underlying PKC-induced modulation of Ca^{2+} sparklet activity. Yang and coworkers found that PKC could directly phosphorylate serine 1928 of $Cav\alpha 1.2$ channels. Thus, it is intriguing to speculate that direct phosphorylation of this L-type Ca^{2+} channel subunit by PKC may play a critical role in the induction of persistent Ca^{2+} sparklet activity. Consistent with this, our data indicate that coexpression of $Cav\alpha 1.2$ and PKC α is sufficient to produce persistent Ca^{2+} sparklet activity in tsA-201 cells. Future experiments should examine the molecular mechanisms by which PKC α promotes persistent L-type Ca^{2+} channel gating.

A particularly interesting finding in this study is that inhibition of PP2A or PP2B increases Ca^{2+} influx by recruiting new Ca^{2+} sparklet sites and increasing the activity of low nP_s sites. Inhibition of these phosphatases did not increase the activity of high nP_s sites, indicating that these sites were maximally activated under control conditions. These findings suggest that in silent Ca^{2+} sparklet sites, PP2A and/or PP2B activity is sufficiently high to exceed PKC α , thus favoring dephosphorylation of Ca^{2+} channels. Accordingly, in low nP_s sites, the balance between PKC α and PP2A/PP2B favors submaximal PKC α -dependent phosphorylation, which induces low Ca^{2+} sparklet activity. In high nP_s sites, however, PKC α activity exceeds PP2A/PP2B activity, thus favoring maximal phosphorylation of Ca^{2+} channels. Because silent, low, and high activity Ca^{2+} sparklet sites coexist in the same cell, these findings support a model in which Ca^{2+} influx is determined locally by the relative balance

between PKC α -dependent phosphorylation and opposing phosphatases. Such a situation allows for dynamic, local modulation of Ca²⁺ channel gating modalities.

This model implies that when prevailing conditions favor dephosphorylation, Ca²⁺ influx is most likely dominated by random, sporadic openings of solitary L-type Ca²⁺ channels. A physiological stimulus that increases PKC α activity, or decreases the activity of PP2A and/or PP2B, would promote Ca²⁺ influx via persistent Ca²⁺ sparklet sites. In this case, Ca²⁺ influx would be determined by persistent Ca²⁺ channel activity in addition to rare, stochastic openings of these channels. Accordingly, vasoactive agents that activate PKC (e.g., angiotensin II and UTP) would increase Ca²⁺ influx and thereby constrict arterial smooth muscle, at least in part, by increasing persistent Ca²⁺ sparklet activity.

An issue that was not addressed by Navedo et al. (2005) is the mechanism by which Bay-K 8644 increases persistent Ca²⁺ sparklet activity in arterial myocytes. To our knowledge, Bay-K 8644 does not directly activate PKC α . Thus, how does Bay-K 8644 increase persistent Ca²⁺ sparklet activity in these cells? The experiments in this study provide insight into this issue. Note that Bay-K 8644 induced high nP_s , persistent Ca²⁺ sparklet activity in tsA-201 cells expressing Cav α 1.2 and PKC α ; persistent Ca²⁺ sparklet activity was never observed after Bay-K 8644 in tsA-201 cells expressing Cav α 1.2 only. Based on these data, we propose a positive feedback model to explain this apparent discrepancy between the effects of Bay-K 8644 on Ca²⁺ sparklets in tsA-201 cells and arterial myocytes. In this model, application of Bay-K 8644 (or any other Ca²⁺ channel opener) to arterial myocytes causes an increase in Ca²⁺ influx (i.e., by increasing the mean open time and/or open probability of L-type Ca²⁺ channels) that would activate nearby Ca²⁺-sensitive PKC α . Once activated, this kinase can induce L-type Ca²⁺ channels to operate in a persistent gating mode, thus causing further Ca²⁺ influx, which could presumably maintain PKC α activity and hence persistent Ca²⁺ sparklet activity. Because PKC expression in arterial myocytes is punctate (Maasch et al., 2000; Navedo et al., 2005), PKC α and persistent Ca²⁺ sparklet activity does not propagate throughout the cell. As noted above, protein phosphatases provide additional negative control to this system. An interesting implication of this model is that while Ca²⁺ sparklets could be evoked by voltage or pharmacological means wherever Cav α 1.2 channels are expressed, persistent Ca²⁺ channel activity would only occur in regions of the cell membrane where both Cav α 1.2 and PKC α are expressed. Future experiments should investigate the spatial and temporal relationship between Ca²⁺ sparklets and PKC α in arterial myocytes.

To conclude, we demonstrated that persistent Ca²⁺ sparklet activity is a fundamental feature of L-type Cav α 1.2 channels in association with PKC α . Because PKC and Cav α 1.2 are ubiquitously expressed, our find-

ings support the concept that Ca²⁺ influx via persistent L-type Ca²⁺ channels may represent a general mechanism for the control of steady-state Ca²⁺ influx in excitable cells.

Furthermore, our observations support the concept that PKC α , PP2A, and PP2B form a signaling module that tunes the activity of Ca²⁺ channels, allowing for local, dynamic regulation of steady-state Ca²⁺ influx in cerebral arterial smooth muscle and perhaps excitable cells in general.

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