Biphasic Modulation of Ryanodine Receptors by Sulfhydryl Oxidation in Rat Ventricular Myocytes

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ABSTRACT To understand better the modulation of ryanodine receptors (RyRs) during oxidative stress, the effect of 4,4'-dithiodipyridine (DTDP), a cell-permeant and thiol-reactive oxidant, on global Ca$^{2+}$ signal and spontaneous Ca$^{2+}$ sparks of rat ventricular myocytes was investigated. It was shown that a brief Ca$^{2+}$ transient was elicited by DTDP, when its concentration was raised to 100 μM DTDP. In addition a dose-dependent increase of cytoplasmic free Zn$^{2+}$ concentration was induced by DTDP. An increase of the frequency of spontaneous Ca$^{2+}$ sparks appeared at 3 μM DTDP, whereas higher concentration of DTDP caused a biphasic change of the frequency in both intact and permeabilized myocytes. Consistent with the biphasic effect, caffeine-induced Ca$^{2+}$ transients were similarly affected. Because DTDP did not reduce the free Ca$^{2+}$ concentration in the sarcoplasmic reticulum lumen, it is likely that the effects of DTDP on the frequency and caffeine-induced Ca$^{2+}$ transients are due mainly to sulfhydryl oxidation-induced activation and subsequent inactivation of RyRs. Unlike the frequency, the spatio-temporal properties of Ca$^{2+}$ sparks were not influenced by DTDP. The finding that DTDP does not affect the duration of Ca$^{2+}$ sparks is inconsistent with that the DTDP-induced increase of the open time of reconstituted RyR channels. The mechanism underlying this discrepancy, especially the possible role of the interaction between arrayed RyRs in myocytes, is discussed. This study suggests that, even if oxidative stress is mild enough not to cause intracellular Ca$^{2+}$ accumulation, it may affect signaling pathways through directly modulating the RyR or its complex and in turn changing the frequency of spontaneous Ca$^{2+}$ sparks. Thus, the functional importance of moderate oxidative stress should not be overlooked.

INTRODUCTION

Normal calcium signaling is essential for regular function of cardiomyocytes. In response to oxidative stress, the intracellular Ca$^{2+}$ accumulation induced by increased formation of reactive oxygen species (ROS) would cause injury and dysfunction of cardiac myocytes (1–3).

In cardiac myocytes the sarcoplasmic reticulum (SR) is a main intracellular Ca$^{2+}$ store. Ryanodine receptors (RyRs) on the membrane of the SR play a central role in modulating Ca$^{2+}$ signaling. RyRs contain many cysteine residues in their regulatory domain and putative Ca$^{2+}$ pore region. These residues are susceptible to modification by oxidants, such as ROS and reactive disulfides. In fact, the effect of ROS on the activity of reconstituted single RyR channel has clearly been shown (4,5). In addition, ROS-induced Ca$^{2+}$ release from isolated SR vesicles was also observed (3,6). Because the effects of ROS can be prevented or reversed by thiol reducing agents, such as dithiothreitol (DTT), thiol oxidation of RyRs is suggested (4,5). 2,2'- or 4,4'-dithiodipyridine (DTDP) is a cell-permeant and thiol specific reagent, and oxidizes proteins through a thiol-disulfide exchange (7). It has been previously reported that the activation of the cardiac RyR by 2,2'- or 4,4'-DTDP in lipid bilayers, which occurs within minutes, was followed by an irreversible loss of RyR activity (8). All of these results illustrate the important role of sulfhydryl oxidation in regulating RyR gating.

It is well established that RyRs in intact muscle cells are assembled into two-dimensional arrays in the SR membrane (9,10). The interaction between isolated RyRs is modulated by various factors, including monovalent cation and functional states of RyRs (11,12). This interaction and its modulation may play an essential role in the activation and termination of RyRs gating. Recently, by analyzing the quantum nature of Ca$^{2+}$ sparks, elementary Ca$^{2+}$ release events mediated by RyRs in the array, Cheng et al. have demonstrated that the gating kinetics of multiple RyRs is reshaped by the array-based interaction between these channels (13). Moreover, it is known that the endogenous antioxidant defense system is present in various cells, including cardiomyocytes. In view of this it is likely that the effect of oxidants on in situ arrayed RyRs is different from that seen with reconstituted single RyR channel.

To understand better the modulation of RyRs during oxidative stress, it is necessary to investigate the effect of oxidants on RyRs in situ. Oxidants usually have complicated effects, including peroxidation of membrane phospholipids, oxidation of protein thiol groups, and mutagenesis of DNA (14). Comparatively, the effect of 4,4'-DTDP is more specific because its oxidizing effect is restricted to thiol groups. Therefore, the effect of 4,4'-DTDP on global and local (spontaneous Ca$^{2+}$ sparks) Ca$^{2+}$ signal in isolated rat ventricular myocytes was investigated. The frequency and spatio-temporal parameters of spontaneous Ca$^{2+}$ sparks were measured. In addition, this study was also performed on saponin...
permeabilized myocytes. The permeabilized cells probably bridge the gap between intact cell and isolated systems.

METHODS

Cell isolation

Rat ventricular myocytes were enzymatically isolated as described previously (15). Briefly, adult Sprague-Dawley rats (220–300 g) were anesthetized by sodium pentobarbital (100 mg/kg, intraperitoneal injection). The heart was quickly excised and perfused with O2 saturated Ca2+-free Tyrode’s solution at 37°C for 5 min. This solution consisted of (in mM): NaCl 150, KCl 5.4, MgCl2 1.2, CaCl2 1, HEPES 10, glucose 15, pH 7.4 (NaOH). Then, the perfusion solution was switched to Tyrode’s solution containing 1 mg/ml collagenase (type I), 0.1 mg/ml trypsin (type I), and 0.1 mM CaCl2, and the heart was perfused for another 20 min. Afterwards, the ventricle was dissected, minced, and filtered through nylon mesh. Isolated cells were resuspended in standard physiological solution containing (in mM): NaCl 120, KCl 5.4, MgCl2 1, CaCl2 1, HEPES 20, glucose 15, pH 7.4 (NaOH) at room temperature (22-24°C) for 1 h before use.

Cell permeabilization

Isolated myocytes were permeabilized with 0.01% (w/v) saponin according to a modified method described previously (16). The permeabilization solution contained (in mM): potassium aspartate 100, KCl 20, EGTA 0.1, ATP 3, MgCl2 3.81, HEPES 10, and 8% (w/v) dextran (40,000), to prevent osmotic swelling of the cells; pH 7.2 (KOH). After 1 min permeabilization, the myocytes were perfused with saponin-free internal solution composed of (in mM): potassium aspartate 100, KCl 20, ATP 3, MgCl2 3.81, EGTA 0.5, CaCl2 0.1, phosphocreatine 10, creatine phosphokinase 5 U/ml, HEPES 10, fluo-4 potassium salt 0.04, and 8% (w/v) dextran; pH 7.2 (KOH). As calculated by the computer program WinMAXC 2.5 (Stanford University, Stanford, CA), free [Ca2+]i and [Mg2+]i in this solution were 43 nM and 1 mM, respectively. The total Ca2+ and Mg2+ necessary for obtaining altered free [Ca2+]i and [ATP] were also estimated using this program.

Detection of global Ca2+ transients and spontaneous Ca2+ sparks with fluo-4

Intact myocytes were loaded with 20 μM fluo-4 AM for 30 min at room temperature. The cells were attached to a glass coverslip that formed the bottom of a chamber. The chamber was placed on the stage of an inverted microscope (Nikon Diaphot 300). Before taking records, the cells were attached to a glass coverslip that formed the bottom of a chamber. The chamber was placed on the stage of an inverted microscope (Nikon Diaphot 300). Before taking records, the cells were permeabilized with standard physiological solution for 30 min for deesterification of mag-fura-5 AM. By using polychrome system and high-speed cooled charge-coupled device (Imago QE, Till Photonics, Gräfelfing, Germany) image system, the ratio of fluorescence 340 nm/580 nm was analyzed online with Tillvision software (Till Photonics).

The data of line scan mode were processed by IDL software (Research Systems, Boulder, CO) and a modified spark detection algorithm (17). Ca2+ sparks were identified as local peak elevations of fluorescence intensity, which were >3 mean ± SD of the surrounding background levels. The parameters measured in this study included amplitude (F/F0), fullwidth at half-maximum intensity (FWHM), full duration at half-maximum intensity (FDHM). The frequency of Ca2+ sparks was represented by the number of Ca2+ sparks in one image obtained by line scan mode, event s−1 84 μm−1. Because the frequency was low, especially after DTDP exposure, the total Ca2+ sparks in successively taken 10 X-T images were measured, and an averaged frequency was estimated.

Detection of intracellular Zn2+ with mag-fura-5

Mag-fura-5, a fluorescent dye sensitive to many divalent cations, was used to measure intracellular free Zn2+ concentration ([Zn2+]i), because it has high affinity for Zn2+ (KdZn = 28 nM) (18,19). Intact myocytes were loaded with 5 μM mag-fura-5 AM for 30 min at room temperature and then washed with standard physiological solution for 30 min for deesterification of mag-fura-5 AM. By using polychrome system and high-speed cooled charge-coupled device (Imago QE, Till Photonics, Gräfelfing, Germany) image system, the ratio of fluorescence 340 nm/580 nm was analyzed online with Tillvision software (Till Photonics).

Detection of Ca2+ in SR lumen with mag-fluo-4

Mag-fluo-4, a Ca2+ indicator with low affinity (Kd = 22 μM), was used for detecting free [Ca2+]i in the SR lumen ([Ca2+]SR). For this purpose, isolated myocytes were loaded with 10 μM mag-fluo-4 AM for 2 h, and then incubated in physiological solution for another 2 h at 37°C to allow deesterification and outward leak of this dye from cytoplasm (20,21). Mag-fluo-4 was excited at 488 nm, and the fluorescence measured at >522 nm. A striation pattern of the fluorescence appeared in the dye-loaded resting myocyte, indicating the localization of the dye in the junctional SR. This localization is supported by the finding that the fluorescence decreased when 10 mM caffeine was briefly applied, which caused translocation of Ca2+ from the SR to the cytoplasm.

Statistical analysis

Results are expressed as mean ± SE for the indicated number of the myocytes isolated from at least three animals. Statistical significance was determined by Student’s t-test. P<0.05 was considered to be statistically significant.

RESULTS

Effect of DTDP on [Ca2+]i in intact myocytes

As illustrated in Fig. 1A, low concentration of DTDP such as 3 μM did not cause any change of the fluorescence within 5 min, whereas a slow rise of fluorescence was seen with 10 μM DTDP. The rate of rise and amplitude of the fluorescence signal increased with increasing DTDP concentration. More interestingly, a complicated change of the fluorescence was consistently observed, when the myocytes were exposed to 100 μM DTDP. A rapid increase of the fluorescence, referred to as phase I, appeared within ~0.5 min. After the increased levels during phase I returned to the baseline, the fluorescence started to increase again, which is referred to as phase II. The phase I but not the phase II was blocked by 10 μM ryanodine (data not shown).
In the absence of external Ca\(^{2+}\), 100 \(\mu M\) DTDP caused a similar change of the fluorescence (Fig. 1B), indicating that these effects of DTDP are independent of Ca\(^{2+}\) entry. In addition, in the absence of external Ca\(^{2+}\) the phase I but not the phase II was abolished, if Ca\(^{2+}\) in the SR was depleted by caffeine combined with thapsigargin, an inhibitor of Ca\(^{2+}\)-ATPase (Fig. 1C). This illustrates that the increase of Ca\(^{2+}\) during phase I is due mainly to release from the SR, although other sources, e.g., mitochondria etc., cannot be ruled out.

**Effect of DTDP on \([Zn^{2+}]_i\) in intact myocytes**

The results described above clearly show that neither extracellular Ca\(^{2+}\) entry nor Ca\(^{2+}\) release from the SR contribute to phase II. In addition, washing out DTDP could not reverse the fluorescence changes that occurred during phase II. However, washing with 1 mM DTT significantly reduced the changes during phase II (Fig. 2A), indicating that oxidation of some residues might be involved. As a result of the oxidation, an increase of cytoplasmic free Zn\(^{2+}\) concentration ([Zn\(^{2+}\)_i]) might be induced. In fact, it has previously been shown that oxidants, such as hypochlorous acid and selenite, could increase [Zn\(^{2+}\)_i] in rabbit ventricular myocytes (22).

To explore if the DTDP-induced phase II really is due to an increase of [Zn\(^{2+}\)_i], the effect of DTDP in the presence of TPEN, a cell-permeable Zn\(^{2+}\) chelator was studied. In control experiments, exposure of the myocytes to 1 \(\mu M\) Zn\(^{2+}\)-pyrithione (ZnPyr), a specific Zn\(^{2+}\) ionophore, caused an increase of the fluo-4 fluorescence (Fig. 2B). This signal was reversed by 10 \(\mu M\) TPEN. Interestingly, 10 \(\mu M\) TPEN could significantly reverse the DTDP-induced phase II changes (Fig. 2C). If the myocytes were pretreated with 10 \(\mu M\) TPEN, the DTDP-induced phase I was not affected, but the phase II was clearly depressed (Fig. 2D).

The effects of DTDP and TPEN on the fluo-4 fluorescence during phase II were summarized in Fig. 2E. As control, the ratio of F/F\(_0\) was 1. TPEN (10 \(\mu M\)) did not change this ratio. Perfusing the myocytes with 100 \(\mu M\) DTDP for 4 min significantly increased the fluorescence signal and most (~75%) of this increase could be reversed by 10 \(\mu M\) TPEN.

Moreover, mag-fura-5, a Zn\(^{2+}\) indicator, was used to further test if DTDP increases [Zn\(^{2+}\)_i]. It was apparently observed that only phase II occurred when the myocytes were exposed to 100 \(\mu M\) DTDP (Fig. 2F). Similar results were observed in the other five myocytes.

Taken together, these observations clearly indicate that oxidation-induced increase of [Zn\(^{2+}\)_i] contributes largely to phase II.

**Effect of DTDP on spontaneous Ca\(^{2+}\) sparks and caffeine-induced Ca\(^{2+}\) transients in intact myocytes**

Because the DTDP-induced phase I is due mainly to Ca\(^{2+}\) release from the SR, the effects of DTDP on the local (spontaneous Ca\(^{2+}\) sparks) and global (caffeine-induced Ca\(^{2+}\) transients (CaTs)) Ca\(^{2+}\) release of RyRs in intact myocytes were examined in the following experiments.

First, the effect of DTDP on the spontaneous Ca\(^{2+}\) sparks in intact myocytes was investigated. Representative confocal images of Ca\(^{2+}\) sparks obtained by line scanning mode are shown in Fig. 3. One spontaneous Ca\(^{2+}\) spark was observed in this myocyte before DTDP exposure (Fig. 3A). At 0.5 min of 30 \(\mu M\) DTDP perfusion the number of Ca\(^{2+}\) sparks increased to 9 (Fig. 3B). With prolonged treatment of DTDP, an evident increase of the basal fluorescence appeared and no Ca\(^{2+}\) sparks were detected (Fig. 3C). According to the results illustrated in Fig. 2, the enhancement of the basal fluorescence is due to an increase of [Zn\(^{2+}\)_i]. It has been shown that Zn\(^{2+}\) ions have depression effect on the activities of RyRs (23). Moreover, the increase of the basal fluorescence may interfere with the identification of Ca\(^{2+}\) sparks.
It would be interesting to explore if the DTDP-induced increase of both basal fluorescence and \([\text{Zn}^{2+}]_i\) is responsible for the vanishing of Ca\(^{2+}\) sparks. Fig. 3, D–F, obtained from another myocyte clearly shows that in the presence of 10 \(\mu\)M TPEN 30 \(\mu\)M DTDP still caused biphasic effect on Ca\(^{2+}\) sparks. In this myocyte 2 spontaneous Ca\(^{2+}\) sparks were found before DTDP exposure (Fig. 3 D). At 0.5 min of 30 \(\mu\)M DTDP perfusion the number of Ca\(^{2+}\) sparks increased to 5 (Fig. 3 E). With prolonged treatment of DTDP the basal fluorescence did not change, but Ca\(^{2+}\) sparks vanished (Fig. 3 F). Therefore, the decrease of Ca\(^{2+}\) sparks induced by DTDP cannot be ascribed to the increase of basal fluorescence and \([\text{Zn}^{2+}]_i\). This conclusion is further confirmed by the results of permeabilized myocytes (see the next section).

The statistical data are represented in Fig. 4. Fig. 4 A shows that, the frequency of spontaneous Ca\(^{2+}\) sparks in control myocytes was 1.78 ± 0.23 event s\(^{-1}\) 84 \(\mu\)m\(^{-1}\) (n = 22), and it was relatively stable within 5 min. Exposure to 3 \(\mu\)M DTDP slightly but significantly increased the frequency, and this increase was sustained within at least 5 min. When DTDP was raised to 30 \(\mu\)M, a biphasic change of the frequency occurred. At 0.5 min, the frequency increased to 270 ± 35% of the control. Then, the frequency gradually decreased. After 4 min, almost no spontaneous Ca\(^{2+}\) sparks could be detected (Fig. 4 A). Fig. 4 A also shows that the DTDP-induced biphasic change of the frequency was not affected by 10 \(\mu\)M TPEN. Although the frequency was affected, all of the spatio-temporal parameters of the spontaneous Ca\(^{2+}\) sparks, including \(F/F_0\), FWHM and FDHM, were not changed by 3 \(\mu\)M or 30 \(\mu\)M DTDP (Fig. 4 C). Unlike that in single reconstructed RyRs where DTDP induced an increase of the mean opening time (8), no alteration of temporal parameters, especially FDHM were seen in these experiments.

The depression of spontaneous Ca\(^{2+}\) sparks induced by 30 \(\mu\)M DTDP for 5 min could not be restored by washing out DTDP for 10 min (data not shown). To see whether the thiol oxidation of RyRs is involved in this depression, the effect of DTT was examined. One mM DTT alone had no obvious effect on the frequency, indicating that most of the RyRs in resting myocytes probably exist in a reduced form.
However, spontaneous Ca\textsuperscript{2+} sparks depressed by DTDP could be fully recovered by 1 mM DTT (Fig. 4 B).

Second, the effect of DTDP on the CaTs was investigated. In intact myocytes CaTs could be evoked by repetitive exposures to 10 mM caffeine at an interval of 10 min without evident decline in their amplitude, as shown in Fig. 5 A (a). This figure also indicates that the amplitude of CaTs was not affected by 10 μM TPEN. With 3 μM DTDP for 5 min, a slight increase of CaTs was observed (Fig. 5 A (d) and B). Prolonging 3 μM DTDP exposure to 10 min reversed the increase to an evident decrease, 68 ± 3% of the control (Fig. 5 A (c) and B). The biphasic effect of DTDP on CaTs was more clearly seen with 30 μM DTDP. In comparison with the control, a significantly increased CaT was evoked by caffeine combined with 30 μM DTDP (Fig. 5 A (d) and B). However, if the myocytes had been treated with 30 μM DTDP for 5 min before caffeine exposure, the CaT was greatly decreased to 34 ± 2% of the control (Fig. 5 A (e) and B). The DTDP-induced decrease of CaTs could not be prevented by TPEN (Fig. 5 A (f)), indicating that the increase of [Zn\textsuperscript{2+}]\textsubscript{i} cannot account for the DTDP-induced depression of CaTs. However, as shown in Fig. 5 A (g), 1 mM DTT could antagonize the DTDP-induced depression of CaTs. The summarized results of the effect of DTDP on CaTs are represented in Fig. 5 B.

The biphasic effects of DTDP on the spontaneous Ca\textsuperscript{2+} sparks and global CaTs suggest that thiol oxidation might biphasically modulate Ca\textsuperscript{2+} release of RyRs in situ.

**Effect of DTDP on spontaneous Ca\textsuperscript{2+} sparks in permeabilized myocytes**

DTDP may change the activity of RyRs through altering [Ca\textsuperscript{2+}]\textsubscript{i} in intact myocytes. To examine the probable role of DTDP-altered [Ca\textsuperscript{2+}]\textsubscript{i} in the effects of DTDP, the following experiments were performed on saponin permeabilized myocytes. In these experiments 0.5 mM EGTA was used to buffer [Ca\textsuperscript{2+}]\textsubscript{i} usually to 43 nM. As shown in Fig. 6 A, under this condition 100 μM DTDP did not change fluo-4 fluorescence in permeabilized myocyte in the presence or absence of 10 μM TPEN, demonstrating proper buffering of [Ca\textsuperscript{2+}]\textsubscript{i} and [Zn\textsuperscript{2+}]\textsubscript{i}.

The control frequency of spontaneous Ca\textsuperscript{2+} sparks in permeabilized myocytes was 3.56 ± 0.22 event s\textsuperscript{-1} 84 μm\textsuperscript{-1} (n = 37). It did not change within 10 min, showing that the activity of RyRs is well maintained after permeabilization (Fig. 6 B). It was observed that 100 μM DTDP caused a biphasic change of the frequency. At 0.5 min, the frequency was raised to 243 ± 24% of the control, and then decreased gradually. Five minutes later, spontaneous Ca\textsuperscript{2+} sparks were rarely detectable (Fig. 6 B). This time course was basically similar to that seen in intact cells. Biphasic change of the frequency also appeared after 10 μM DTDP exposure, but 10 μM DTDP could not completely abolish Ca\textsuperscript{2+} sparks even 10 min later (Fig. 6 B). With 3 μM DTDP, only a transient increase of the frequency was observed.

Similar to intact cells, all of the spatio-temporal parameters of spontaneous Ca\textsuperscript{2+} sparks in permeabilized myocytes were not affected by DTDP (data not shown). One mM DTT alone had no evident effect on the frequency, but DTT could reverse the depression effects of DTDP (Fig. 6 C). The antagonizing effect of DTT further suggests the involvement of the thiol oxidation of RyRs.

Ca\textsuperscript{2+} accumulation and ATP depletion have been shown in the cells subjected to oxidative stress (1–3,24,25). Both Ca\textsuperscript{2+} and ATP are known to modulate RyRs. Thus, the effect of DTDP was investigated at different Ca\textsuperscript{2+} or ATP concentrations. It was found that the biphasic effect of 100 μM DTDP on the frequency was not altered by changing [Ca\textsuperscript{2+}] (from 43 to 115 nM) and [ATP] (from 0.3 to 3 mM) (data not shown). It should be emphasized that, compared with that at 43 nM [Ca\textsuperscript{2+}], an increase of Ca\textsuperscript{2+} sparks frequency could still be observed at 115 nM [Ca\textsuperscript{2+}], although the basal fluorescence at 115 nM [Ca\textsuperscript{2+}] was increased to an extent similar to that with 30 μM DTDP (Fig. 3 C).

**Effect of DTDP on Ca\textsuperscript{2+} content in the SR lumen**

It is well established that the gating of RyRs is modulated by Ca\textsuperscript{2+} content in the SR lumen (26,27). To observe if the effect of DTDP on the gating of RyRs is mediated by altering Ca\textsuperscript{2+} content in the SR lumen, mag-fluo-4 was used to directly measure the [Ca\textsuperscript{2+}]\textsubscript{SR}. Because the Ca\textsuperscript{2+} affinity of mag-fluo-4 (K\textsubscript{s} = 22 μM) is evidently lower than the [Ca\textsuperscript{2+}]\textsubscript{SR} (~1 mM), this dye may not be proper for measuring [Ca\textsuperscript{2+}]\textsubscript{SR}. However, many previous studies have used this dye to successively estimate [Ca\textsuperscript{2+}]\textsubscript{SR} or free [Ca\textsuperscript{2+}] in the endoplasmic reticulum under various conditions (20,28–30).
The localization of mag-fluo-4 is illustrated in Fig. 7A. In the resting myocytes the fluorescence of this dye showed a striation pattern, and this pattern disappeared following caffeine exposure, indicating that this dye was localized in the SR lumen. It was found that caffeine caused an evident reduction of the fluorescence, illustrating a caffeine-induced decrease of the \([\text{Ca}^{2+}]_{\text{SR}}\). In the presence of 30 \(\mu\)M DTDP the caffeine-induced response became smaller (Fig. 7A). Fig. 7B (a)
shows that the DTDP-induced decrease of the caffeine response could be antagonized by 1 mM DTT. More interestingly, it was consistently observed that DTDP increased but not decreased the fluorescence, and this increase could be reversed by DTT (Fig. 7B(b)). It is clearly indicated that the \([\text{Ca}^{2+}]_{\text{SR}}\) is not reduced by DTDP. So, the depression and final disappearance of spontaneous \(\text{Ca}^{2+}\) sparks caused by DTDP is likely to be due to sulfhydryl oxidation-induced inactivation of RyRs.

To clarify if DTDP-induced rise of the fluorescence in intact myocytes really represents an increase of the \([\text{Ca}^{2+}]_{\text{SR}}\), we performed the experiments on permeabilized myocytes using mag-fluo-4. It was seen that DTDP evidently increased mag-fluo-4 fluorescence in permeabilized myocytes (Fig. 7C). This increase was not affected by the presence of thapsigargin and TPEN, indicating that the change of mag-fluo-4 fluorescence does not result from \(\text{Ca}^{2+}\) uptake via \(\text{Ca}^{2+}\)-ATPase or \(\text{Zn}^{2+}\) increase in the SR lumen and DTDP may induce an increase of \([\text{Ca}^{2+}]_{\text{SR}}\). Because mag-fluo-4 is not an ideal dye for this purpose, it is desirable to use the dye with lower \(\text{Ca}^{2+}\) affinity to repeat these observations in further study.

In addition, it was noted that, DTDP-induced increase of the frequency of \(\text{Ca}^{2+}\) sparks appeared evidently earlier than DTDP-induced increase of mag-fluo-4 fluorescence. Thus, the increase of \([\text{Ca}^{2+}]_{\text{SR}}\) is not the main factor for the activation of RyRs, although it may play some roles.

**DISCUSSION**

**Modulation of RyRs by sulfhydryl oxidation in cardiac myocytes**

Consistent with previous results obtained on reconstituted single RyR channels (8), the results of this study show that the activity of RyRs in situ is also affected by DTDP in a
dose-dependent manner. An increase of the frequency of spontaneous Ca\(^{2+}\) sparks was seen at 3 \(\mu\)M DTDP, whereas higher concentrations of DTDP caused a biphasic change of the frequency and finally disappearance of spontaneous Ca\(^{2+}\) sparks in both intact and permeabilized myocytes, in which cytoplasmic Ca\(^{2+}\) was buffered with EGTA. Moreover, the amplitude of CaTs was similarly altered by DTDP.

In addition, this study showed that \([\text{Ca}^{2+}]_{\text{SR}}\) was not decreased, but increased by DTDP and this increase was independent of Ca\(^{2+}\) uptake via Ca\(^{2+}\)-ATPase in the SR membrane. The exact origin of this increase remains unclear. As one possibility, we suppose that it results from DTDP oxidation-induced dissociation of Ca\(^{2+}\) ions from Ca\(^{2+}\)-binding proteins in the SR lumen, e.g., calsequestrin. Calsequestrin is known as the most abundant Ca\(^{2+}\)-binding protein in the SR. Its thioredoxin structure suggests that calsequestrin may be sensitive to redox (31,32). Actually, it has been reported that oxidation of calbindin D\(_{28k}\), a Ca\(^{2+}\)-binding protein, decreases its Ca\(^{2+}\)-binding affinity (33). Because DTDP-induced increase of the free \([\text{Ca}^{2+}]\) in the SR lumen was later than DTDP-induced increase of Ca\(^{2+}\) sparks frequency, it is unlikely to be the main factor for the activation of RyRs, although it may play some roles.

Taken together these results indicate that the effects of DTDP on the frequency of spontaneous Ca\(^{2+}\) sparks and the amplitude of CaTs are due mainly to sulfhydryl oxidation-induced activation and subsequent inactivation of RyRs. In fact, it has previously been proposed that, there are at least two classes of sulfhydryls in RyR (8). Oxidation of one class of sulfhydryls causes activation, and inactivation would occur due to further oxidation of the other class.

However, compared with previous studies, an apparent disparity is noted. Single channel studies have evidently shown DTDP-induced increase of mean open time of RyR (8). Accordingly, it should be expected that the temporal parameters of spontaneous Ca\(^{2+}\) sparks would be prolonged by DTDP. In contrast, in this study we found no change of the temporal parameters. This discrepancy needs to be explained.

Recent evidence shows that isolated RyRs interact with each other (11,12,34). This interaction can be modulated by several factors, e.g., the concentration of monovalent cations (12), the functional states of RyRs (11). A possible explanation of the inconsistency between the behavior of isolated RyRs and that of RyRs in intact cells could be that the interaction between arrayed RyRs prevents the change of the number of RyRs forming single spontaneous Ca\(^{2+}\) sparks. Because the average release duration of simulated Ca\(^{2+}\) sparks is inversely related to the number of RyRs of which Ca\(^{2+}\) sparks consist (13), the temporal parameters of Ca\(^{2+}\) sparks would not be altered, if the number of RyRs does not change. This study did not find any change of \(F/F_0\) in the presence of DTDP, indicating that the mean number of RyRs forming single spontaneous Ca\(^{2+}\) sparks is not affected by DTDP. At the moment, this assumption is speculative. Direct evidence for the role of the interaction between arrayed RyRs for Ca\(^{2+}\) movements in the living cell obviously is needed.

In addition to the interaction between arrayed RyRs, under in situ condition numerous proteins physically associating with RyR are involved in modulating the gating of RyRs (35). Thus, the possibility that the discrepancy between results obtained from intact cells and those obtained from isolated receptors could be due to the absence of some RyR-associated proteins in the isolated system cannot be excluded.
Regardless of the exact reason for this inconsistency, this study clearly indicates that extrapolating in vitro result to in vivo events is not straightforward.

**Functional consequence of DTDP-induced change of the frequency of spontaneous Ca\(^{2+}\) sparks**

Ca\(^{2+}\) sparks are fundamental SR Ca\(^{2+}\) release events, representing the nearly synchronous activation of a cluster of RyRs at a single junction. In smooth muscle the roles of spontaneous Ca\(^{2+}\) sparks in positive- and negative-feedback regulation have been proposed (36). Although spontaneous Ca\(^{2+}\) sparks are generally thought to play an important role in modulating Ca\(^{2+}\) homeostasis in cardiac myocytes (37), it is unclear what the functional consequence of the change of the frequency of spontaneous Ca\(^{2+}\) sparks is.

The change in the frequency might alter local Ca\(^{2+}\)-mediated signaling pathways. Actually, it has recently been shown that Ca\(^{2+}\) sparks are able to elicit miniature mitochondria matrix Ca\(^{2+}\) signal called Ca\(^{2+}\) sparks, which are restricted to single mitochondria and last <500 ms (38). As a result, it would affect the metabolism and ATP production in mitochondria. Therefore, the functional consequence of the change of the frequency of spontaneous Ca\(^{2+}\) sparks should not be overlooked, even if the change of the frequency is not accompanied by apparent intracellular Ca\(^{2+}\) accumulation.

Compared with the functional consequences of low concentrations of DTDP those of high concentrations of DTDP are more obvious because they lead to the failure of the RyRs to release Ca\(^{2+}\). Although it would cause dysfunction of excitation-contraction coupling, this failure of RyRs to release Ca\(^{2+}\) may reduce intracellular Ca\(^{2+}\) accumulation and in turn cellular injury. It probably represents a novel mechanism to protect cardiomyocytes from the injury induced by oxidative stress.

**DTDP-induced increase of [Zn\(^{2+}\)], and its role in DTDP-induced modulation of RyR**

It is known that oxidants, such as hypochlorous acid and selenite, increased [Zn\(^{2+}\)], in rabbit ventricular myocytes (22). More recently, DTDP-induced increase of [Zn\(^{2+}\)] has been reported in neurons (39,40). In this study we provided clear evidence for DTDP-induced increase of [Zn\(^{2+}\)], in rat ventricular myocytes.

We have shown that ryanodine binding to the SR vesicle of calf cardiac muscle was inhibited by Zn\(^{2+}\) (23). Therefore, we were interested in examining if the sulfhydryl oxidation-induced increase of [Zn\(^{2+}\)], influences the effect of DTDP on Ca\(^{2+}\) signaling. It was observed that the effect of DTDP on the frequency of spontaneous Ca\(^{2+}\) sparks was not altered by TPEN (Fig. 4A), representing that the depression effect of DTDP on spontaneous Ca\(^{2+}\) sparks cannot be ascribed to the increase of [Zn\(^{2+}\)]. In addition, it was found that 3 \(\mu\)M DTDP did not increase [Zn\(^{2+}\)] (Fig. 1A), but persistently activated RyRs (Fig. 4A), indicating that the increase of [Zn\(^{2+}\)] also is not the prerequisite for DTDP-induced activation of RyRs.

Although DTDP-induced increase of [Zn\(^{2+}\)] does not influence the effect of DTDP on spontaneous Ca\(^{2+}\) sparks, some of the physiological and pathological processes may be affected by increased [Zn\(^{2+}\)] (41). As an example, it has been shown that Zn\(^{2+}\) regulates K\(_{ATP}\) channels from both intracellular and extracellular sides (42). This regulation is thought to protect the cardiomyocytes from the injury induced by ischemia and reperfusion. Thus, it is reasonable to propose that fair increase of [Zn\(^{2+}\)] induced by moderate oxidative stress may provide another novel mechanism to protect cardiac myocytes from ischemia-reperfusion insult.

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