Essential Role of Calmodulin in RyR Inhibition by Dantrolene

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Received January 1, 2015; accepted April 28, 2015

ABSTRACT

Dantrolene is the first line therapy of malignant hyperthermia. Animal studies suggest that dantrolene also protects against heart failure and arrhythmias caused by spontaneous Ca2+ release. Although dantrolene inhibits Ca2+ release from the sarcoplasmic reticulum of skeletal and cardiac muscle preparations, its mechanism of action has remained controversial, because dantrolene does not inhibit single ryanodine receptor (RyR) Ca2+ release channels in lipid bilayers. Here we test the hypothesis that calmodulin (CaM), a physiologic RyR binding partner that is lost during incorporation into lipid bilayers, is required for dantrolene inhibition of RyR channels. In single channel recordings (100 nM cytoplasmic [Ca2+] + 2 mM ATP), dantrolene caused inhibition of RyR1 (rabbit skeletal muscle) and RyR2 (sheep) with a maximal inhibition of Po (E_{max}) to 52 ± 4% of control only after adding physiologic [CaM] = 100 nM. Dantrolene inhibited RyR2 with an IC50 of 0.16 ± 0.03 μM. Mutant N98S-CaM facilitated dantrolene inhibition with an IC50 = 5.9 ± 0.3 nM. In mouse cardiomyocytes, dantrolene had no effect on cardiac Ca2+ release in the absence of CaM, but reduced Ca2+ wave frequency (IC50 = 0.42 ± 0.18 μM, E_{max} = 47 ± 4%) and amplitude (IC50 = 0.19 ± 0.04 μM, E_{max} = 66 ± 4%) in the presence of 100 nM CaM. We conclude that CaM is essential for dantrolene inhibition of RyR1 and RyR2. Absence explains why dantrolene inhibition of single RyR channels has not been previously observed.

Introduction

Dantrolene is a well known inhibitor of Ca^{2+} release in skeletal muscle (Hainaut and Desmedt, 1974) that has been used clinically as the treatment of malignant hyperthermia (MH). MH is a potentially fatal inherited disorder of skeletal muscle (Hainaut and Desmedt, 1974) that has been identified in the DP1 regions in RyR1 and RyR2 (Parness et al., 1995; Paul-Pletzer et al., 2002, 2005; Kobayashi et al., 2009). There has been only one direct observation of RyR inhibition by dantrolene in bilayer-based single channel recordings (Nelson et al., 1996). Studies since then find no effect of dantrolene in single channel recordings (Szentesi et al., 2001; Cherednichenko et al., 2008; Diaz-Sylvester et al., 2008; Wagner et al., 2014). Hence, it is not clear if dantrolene acts directly on the RyR or some other protein involved in excitation-contraction coupling such as the DHPR (Salata et al., 1983; Chou et al., 2014).

Dantrolene acts on skeletal and cardiac muscle by inhibiting Ca^{2+} release from the SR (Hainaut and Desmedt, 1974; Kobayashi et al., 2005; Uchinoumi et al., 2010). Assays of Ca^{2+} release in intact myocytes and cell homogenates containing SR vesicles (Fruen et al., 1997) suggest that dantrolene inhibits the SR Ca^{2+} release channel with a half-inhibiting concentration (IC50) of 0.3 μM (Kobayashi et al., 2009). Even though a dantrolene binding site has been identified in the DP1 regions in RyR1 and RyR2 (Parness and Palnitkar, 1995; Paul-Pletzer et al., 2002, 2005; Kobayashi et al., 2009), there has been only one direct observation of RyR inhibition by dantrolene in bilayer-based single channel recordings (Nelson et al., 1996). Studies since then find no effect of dantrolene in single channel recordings (Szentesi et al., 2001; Cherednichenko et al., 2008; Diaz-Sylvester et al., 2008; Wagner et al., 2014). Hence, it is not clear if dantrolene acts directly on the RyR or some other protein involved in excitation-contraction coupling such as the DHPR (Salata et al., 1983; Chou et al., 2014).

Calmodulin (CaM) is known to regulate the activity of RyR1 and RyR2 (Tripathy et al., 1995; Balshaw et al., 2001). CaM inhibits RyR2 directly by binding to residues 3583–3603 of each RyR2 subunit (Huang et al., 2013) with high affinity (K_d 20–100 nM) (Guo et al., 2011). Similarly, CaM may either increase RyR1 activity at resting cytoplasmic [Ca^{2+}] or decrease activity at higher [Ca^{2+}] (Tripathy et al., 1995).

Fruen and colleagues (Fruen et al., 1997; Zhao et al., 2001)
found that dantrolene reduces the effect of RyR1 activators (but interestingly, not in RyR2) including CaM, suggesting that CaM might augment dantrolene inhibition of RyR1. During the process of RyR2 isolation from the heart and their incorporation into artificial lipid bilayers, the RyR macromolecular complex stays mostly intact (Marks et al., 2002), except for CaM, which is reported to dissociate from the RyR complex with a time constant of less than 1 minute (Guo et al., 2011). Hence, bilayer-based channel studies would generally have been made devoid of this important regulatory molecule in the RyR complex, whereas CaM is abundant in intact cell and cell homogenates. Therefore, we hypothesize that CaM is the missing protein and that its absence in bilayer experiments provides an explanation as to why dantrolene inhibition has not been observed in single channel RyR recording experiments. We test this hypothesis by examining the effects of dantrolene, in the absence and presence of CaM, on the gating of RyR1 and RyR2 Ca$^{2+}$ release channels incorporated into artificial lipid bilayers and on the frequency and amplitude of Ca$^{2+}$ waves in permeabilized cardiomyocytes.

Materials and Methods

Chemicals. SR vesicles containing RyR1 were isolated from rabbit skeletal muscle, and RyR2 were isolated from sheep hearts (Laver et al., 1995) and incorporated in artificial bilayer membranes composed of a lipid mixture of phosphatidylethanolamine and phosphatidylcholine (8:2 wt/wt, Avanti Polar Lipids, Alabaster, AL) in n-decane (50 mg/ml, ICN Biomedicals, Irvine, CA). Experimental solutions contained (in millimolar) 150 Cs$^+$ (130 CsCH$_3$O$_3$S + 20 CsCl). All solutions were pH buffered using N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid (ICN Biomedicals) and titrated to pH 7.4 using CsOH (ICN Biomedicals). Cytoplasmic solutions were buffered to a redox potential of −232 mV with reduced glutathione disulfide (0.2 mM) and glutathione (GSH; 4 mM), and luminal solutions were buffered to a redox potential of −180 mV with reduced glutathione disulfide (3 mM) and GSH (2 mM). A Ca$^{2+}$-electrode (Radiometer, Brea, CA) was used in our experiments to determine the purity of Ca$^{2+}$$^{2-}$ buffers and Ca$^{2+}$ stock solutions as well as free [Ca$^{2+}$]$_i$$^{2-}$ > 100 nM. The cesium salts were obtained from Sigma-Aldrich (St. Louis, MO). CaCl$$_2$$ was obtained from BDH Chemicals (VWR, Radnor, PA). Calmodulin was obtained from two sources, Sigma-Aldrich (prepared from bovine testes) and Enzo Life Sciences (Farmingdale, NY; prepared from pig brain). Dantrolene (powder) was obtained from Sigma. Dantrolene was prepared as stock solutions in dimethylsulfoxide, and calmodulin was prepared at miliQ (EMD Millipore, Billericia, MA). During experiments, the concentrations of calmodulin, dantrolene, and Ca$^{2+}$ in the cytoplasmic solution were altered by a local perfusion system (O'Neil et al., 2003), which allowed exposure of a single channel to multiple bathing conditions applied in any chosen sequence with an exchange time of ~3 seconds.

Data Acquisition and Analysis. Experiments were carried out at room temperature (23 ± 2°C). Electric potentials are expressed using standard physiologic convention (i.e., cytoplasm relative to SR lumen at virtual ground). Control of the bilayer potential and recording of unitary currents were done using an Axopatch 200B amplifier (Axon Instruments/Molecular Devices, Sunnyvale, CA). The current signal was digitized at 5 kHz and low pass-filtered at 1 kHz. Single channel dwell-time histograms of open and closed time, open probability, and mean open time and mean closed time, were measured using a threshold discriminator at 50% of channel amplitude (Channel13 software; N. W. Laver, nie@nicelaver.com). Individual readings were derived from 45-120 seconds of RyR2 recording. Hill equations were fitted to the dose-response data by the method of least squares. Average data are given as mean ± S.E.M. The statistical significance of differences was tested using Student's t test.

Ca$^{2+}$ Waves Experiments in Ventricular Myocytes. Single ventricular myocytes from 12- to 16-week-old C57Bl/6 mice were isolated using an enzymatic digestion method as previously described (Knollmann et al., 2006). Myocytes were first exposed to a Ca$^{2+}$-free relaxing solution and then permeabilized with suxmon (40 µg/ml) for 60 seconds and placed in internal solution composed (in millimolar) of 120 K-aspartate, 15 KCl, 5 KH$_2$PO$_4$, 0.75 MgCl$_2$, 0.4% dextran (40,000), 10 HEPES, 5 Mg$_2$ATP, 10 glutathione (reduced), 0.025 Fluo-4, and 10 phosphocreatine (di-Na). These solutions also contained 10 U/ml creatine phosphokinase (Hwang et al., 2014) and had free [Ca$^{2+}$]$^{2-}$ = 120 nM. To allow complete removal of CaM binding to RyR2 in permeabilized myocytes (Yang et al., 2014), all Ca$^{2+}$ wave recordings were done after 30-minute incubation with either dantrolene alone or dantrolene + CaM. Free [CaM] was kept at the physiologic concentration of 100 nM. Ca$^{2+}$ waves in myocytes were imaged with a confocal microscope (LSM 510; Zeiss, Thornwood, NY) in line scan mode. Ca$^{2+}$ wave analysis was performed as described (Hwang et al., 2014). Given the variability between different experimental days, the Ca$^{2+}$ wave frequency and amplitude data were normalized to the mean of vehicle group obtained on the same day.

Results

**Essential Role of CaM on RyR1 and RyR2 Inhibition by Dantrolene.** To investigate if CaM binding to RyR1 and RyR2 is a prerequisite for their inhibition by dantrolene, RyR activity was measured in the presence of 100 nM cytoplasmic Ca$^{2+}$ (+2 mM ATP) for periods of 1 minute (vehicle) and then during 1-minute exposure to added dantrolene and then again after dantrolene washout. This sequence was repeated in the absence and presence of exogenous 100 nM CaM as shown in Fig. 1A for rabbit RyR1 and Fig. 1B for sheep RyR2. In the absence of CaM, dantrolene had no observable effect on the channel open probability ($P_o$) of either RyR1 or RyR2. However, when CaM was present in the experimental solutions, dantrolene reduced the $P_o$ of both RyR isoforms. This effect of dantrolene on RyR1 and RyR2 was reversible on washout and inhibition could be seen in multiple applications (Fig. 1C). The data summary from application-washout experiments in Fig. 1D shows that dantrolene at both 10 and 50 µM significantly reduced the open probability of RyR1 to 50% and RyR2 to 45% of control (i.e., vehicle alone), respectively. When CaM was subsequently washed out by perfusion with CaM-free solutions for 1 minute, dantrolene inhibition was abolished (RyR2 open probability 95 ± 9% of vehicle, $P = 0.24$). Therefore, dantrolene inhibition of RyR1 and RyR2 requires the presence of CaM. The concentration-dependence of dantrolene inhibition of RyR2 is shown in Fig. 1E. In the presence of 100 nM CaM (●), inhibition exhibited a sigmoidal dependence on log-concentration with an IC$_{50}$ of 0.16 ± 0.03 µM, a Hill coefficient of ~1 and with a saturating RyR2 open probability ($P_{max}$) of 52 ± 4% compared with the absence of dantrolene. Reducing the CaM concentration to 10 nM approximately halved the magnitude of dantrolene inhibition (Fig. 1E, ○, $P_{max}$ = 80 ± 5%).

**Effect of Dantrolene on RyR Dwell Times.** To gain more insight into the mechanism of dantrolene inhibition, we compiled dwell-time histograms of channel open and closed events of sheep RyR2 at four cytoplasmic [Ca$^{2+}$], ranging from 0.1 µM (end diastolic) to 100 µM (systolic) (Fig. 2, A and B; Supplemental Fig. 1). Histograms are displayed using the log-bin method of Sigworth and Sine (1987), where individual
exponential components appear as peaks centered on their time constant value. In the absence of dantrolene, open and closed dwell times in 1 μM cytoplasmic Ca\(^{2+}\) exhibited peaked distributions that were fitted by two exponential components (see Supplemental Table 1). Addition of dantrolene (10 μM) shifted the peak of the open distributions to shorter times and closed distribution to longer times. Dantrolene had a similar effect in 0.1 μM cytoplasmic Ca\(^{2+}\) but had no effect at 100 μM cytoplasmic Ca\(^{2+}\) (Supplemental Fig. 1). It was not possible to resolve significant differences in the parameters of the exponential fits except for the slow time-constant of the closed times at 0.1 μM cytoplasmic Ca\(^{2+}\) (T2; Supplemental Table 1). However, it was possible to resolve relative changes in the RyR2 mean open and closed durations (Fig. 2C). In 0.1 μM cytoplasmic Ca\(^{2+}\), dantrolene reduced RyR2 \(P_o\) via a decrease in mean channel open duration and an increase in mean closed duration. At 1 μM cytoplasmic Ca\(^{2+}\), the effect of dantrolene was diminished and there was no significant inhibition occurring at higher [Ca\(^{2+}\)]. The effect of dantrolene was to shift the Ca\(^{2+}\)-activation response of RyR2 to higher [Ca\(^{2+}\)].

**Effect of Dantrolene on Ca\(^{2+}\) Waves in Mouse Cardiomyocytes.** The amplitude and frequency of spontaneous Ca\(^{2+}\) waves, two parameters that have been implicated as independent predictors of arrhythmogenicity (Galimberti and Knollmann, 2011), were measured in mouse ventricular myocytes. Examples of the effect of 30-minute exposure to dantrolene (3, 10, 50 μM) on Ca\(^{2+}\) waves recorded in the presence or absence of CaM are presented in Fig. 3A. Dantrolene reduced Ca\(^{2+}\) wave amplitude and frequency in the presence of CaM but had no effect in the absence of CaM (Fig. 3A). This finding is consistent with the single channel experiments with the concentration dependence of these effects (Fig. 3, B and C) exhibiting remarkably similar IC\(_{50}\) and \(E_{max}\) values to that measured in the single channel experiments (Fig. 1).

**Dantrolene Inhibition Can Be Mediated by CaM Mutants.** Because both dantrolene and CaM are RyR2 inhibitors, we investigated the possibility that dantrolene acts by amplifying CaM inhibitory action on RyR2. To test this possibility we measured dantrolene inhibition of sheep RyR2 in the presence of 10 nM (C); mean ± S.E.M., n = 3 to 4) and 100 nM CaM (D); mean ± S.E.M., n = 7 to 20). The luminal [Ca\(^{2+}\)] is 0.1 mM and cytoplasmic [Ca\(^{2+}\)] is 100 nM. The solid curve shows the Hill fit to the data using the equation: where IC\(_{50}\) = 0.16 ± 0.03 μM, \(H = 1.3 ± 0.3\), and \(E_{max} = 52 ± 4\%\). The dashed curve uses the same parameter values, except \(E_{max} = 80 ± 5\%\).
We also investigated dantrolene inhibition in the presence of N98S-CaM that is a CaM mutant that has no inhibitory effect on RyR2 in the absence of dantrolene (Fig. 4C). The advantage of this CaM mutant is that we can examine the effect of varying its concentration on facilitating dantrolene inhibition without the confounding effect of CaM inhibition. In the absence of CaM, dantrolene (10 μM) had no effect on RyR2 Po. Figure 4C shows that addition of only 6 nM N98S-CaM was sufficient to facilitate significant dantrolene inhibition of RyR2. The N98S-CaM facilitation of dantrolene inhibition had a sigmoidal dependence on log-concentration with an IC50 of 5.9 ± 0.3 nM, a Hill coefficient of 5 ± 2.6, and an Emax of 53 ± 4%.

Discussion

Our study presents the first demonstration of dantrolene inhibition of mammalian RyR1 and RyR2 from recordings of single RyR and permeabilized cardiomyocytes. The finding that a physiologic concentration of CaM is required for dantrolene inhibition of these RyRs provides an answer to the long-standing question of why dantrolene, an inhibitor of SR Ca2+ release, had no effect on the activity of mammalian RyR1 and RyR2 in previous single channel studies (Szentesi et al., 2001; Diaz-Sylvester et al., 2008; Wagner et al., 2014). Because CaM readily dissociates from the RyR complex (Guo et al., 2011), CaM would have been absent during those experiments. IC50 for CaM facilitation of dantrolene inhibition appears to be ~10 nM for wt-CaM (Fig. 1E) and 5.9 nM for N98S-CaM (Fig. 4C). These values are ~2-fold lower than the binding affinities for these CaMs on RyR2 (Guo et al., 2011; Hwang et al., 2014).

[3H]ryanodine binding assays have demonstrated a reduction of CaM activation of purified pig RyR1 by dantrolene (Fruen et al., 1997). However, that finding was contradicted by a single channel study (Cherednichenko et al., 2008) that, using similar experimental conditions (100 nM cytoplasmic Ca2+ and 35°C), reported no inhibition by dantrolene (20 μM) of purified rabbit RyR1 channels in bilayers in the presence of exogenous FKBP12 and CaM. Together with the findings reported here, these results suggest that the inhibitory effect of dantrolene on RyR not only requires CaM but also other RyR-associated proteins that are present in native preparations but presumably absent in some purified RyR preparations.

The maximum RyR2 inhibition (Emax = 52%) and IC50 (0.16 ± 0.03 μM; Fig. 1D) are in close agreement with dantrolene inhibition of Ca2+ wave frequency and amplitude in saponin permeabilized cardiomyocytes (Fig. 3) and inhibition of Ca2+ release in SR vesicles from failing dog heart [IC50 = 0.3 ± 0.07 μM (Kobayashi et al., 2009)] and activity of purified RyR1 in [3H]ryanodine binding assays [0.15 ± 0.02 μM (Fruen et al., 1997)]. The dantrolene IC50 reported here coincides with the binding affinity of dantrolene to skeletal muscle SR vesicles [0.277 ± 0.025 μM (Parness and Palnitkar, 1995)] and its IC50 (0.3 ± 0.11 μM) for inhibiting the unzipping of the central and...
N-terminal domains of RyR2 (Kobayashi et al., 2009). The potency of dantrolene in our study is also consistent with the inhibitory action of 1 μM dantrolene on Ca\(^{2+}\) spark frequency in isoproterenol-stimulated cardiomyocytes from R2474S knock-in mice (Kobayashi et al., 2010), increase survival after ventricular fibrillation (Zamiri et al., 2014). Because dantrolene is not an effective RyR inhibitor at high cytoplasmic [Ca\(^{2+}\)]\(_{i}\) yet does not inhibit the amplitude of Ca\(^{2+}\) transients (Maxwell et al., 2012; Zamiri et al., 2014). Thus dantrolene is a diastolic inhibitor of Ca\(^{2+}\) release in failing heart, which has the beneficial actions of increasing diastolic Ca\(^{2+}\) loading of the SR (Maxwell et al., 2012) and reducing diastolic SR Ca\(^{2+}\) leak after ventricular fibrillation (Zamiri et al., 2014). Because dantrolene is not an effective RyR inhibitor at high cytoplasmic [Ca\(^{2+}\)]\(_{i}\), it is not surprising that other dantrolene mechanisms may be more important for suppressing Ca\(^{2+}\) release during skeletal muscle twitches (Flewelling et al., 1983) or suppressing MH episodes.

Single channel recordings of dantrolene inhibition provide a unique opportunity to probe the mechanism of dantrolene inhibition. RyR2 dwell-time distributions (Fig. 2, A and B) indicate that dantrolene decreases the duration of channel openings and increases the duration of closures, characteristics typical of an allosteric inhibitor rather than a channel blocker like the local anesthetics that cause distinct blocking events in single channel recordings that introduce new exponential components in closed time distributions (Tinker and Williams, 1993; Xu et al., 1993; Taushima et al., 2002). Like CaM, dantrolene inhibits RyR2 by destabilizing their open state and stabilizing their closed state. By using a CaM mutation that causes CaM to activate RyR2, we show that dantrolene does not merely increase the efficacy of CaM, but is an inhibitor in its own right (Fig. 4). The RyR has a homotetrameric structure that includes four dantrolene binding sites and at least four CaM binding sites. The dantrolene dose-response (Fig. 1D) exhibited a Hill coefficient of \(\sim 1\), consistent with values obtained from \(^{3}H\)ryanodine binding assays (Fruen et al., 1997). Such a value indicates that the binding of only one dantrolene molecule is sufficient to cause inhibition of RyR2 activity. Interestingly, the dose-response of N98S-CaM facilitation of dantrolene inhibition (Fig. 4C) had a much higher Hill coefficient, consistent with a requirement for multiple CaM molecules on RyR2. The mechanism by which CaM facilitates dantrolene inhibition remains unclear. It is unlikely that dantrolene acts by binding to a site on CaM, because that would not explain the different Hill coefficients for the dantrolene and N98S-CaM dose responses. Also, given the redox buffering of our experimental solutions (4 mM GSH in bilayer experiments and 10 mM GSH in myocyte experiments) it is unlikely that the reducing
properties of dantrolene underlie its inhibition. However, one possibility is that CaM puts the RyR into a conformation that gives dantrolene access to its binding site on the RyR. Two studies have demonstrated that dantrolene has restricted access to its binding site that is regulated by RyR conformation and on the presence of RyR ligands such as Ca²⁺ and ATP (Paul-Pletzer et al., 2001, 2005). An alternative possibility is that CaM is a part of the signaling pathway that transduces dantrolene binding into RyR inhibition. Several studies present evidence that dantrolene modulates interdomain interactions in RyR1 (Kobayashi et al., 2005) and RyR2 (Kobayashi et al., 2009; Uchinoumi et al., 2010; Suetomi et al., 2011; Maxwell et al., 2012) between the N-terminal (1–619 aa), central (2000–2500 aa), and C-terminal domains (3900–end). Our data are consistent with both these possibilities.

In conclusion, we show that CaM binding to the RyR is required to produce dantrolene inhibition in both RyR1 and RyR2. It is likely that other, as yet undefined, factors play a similar role in facilitating dantrolene inhibition.

Acknowledgments

The authors thank Paul Johnson for assisting with the experiments.

Author Contributions

Participated in research design: Oo, Imitaz, Knollmann, Laver.
Conducted experiments: Oo, Gomez-Hurtado, Walweel.
Contributed to new reagents or analytic tools: Knollmann, Laver.
Performed data analysis: Oo, Gomez-Hurtado, Knollmann, Laver.
Wrote or contributed to the writing of the manuscript: Oo, Gomez-Hurtado, van Helden, Knollmann, Laver.

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Fig. 4. Dantrolene inhibition of RyR in the presence of wt-CaM and mutant CaM. (A) Relative effect of wt-CaM (100 nM) on the open probability of RyR1 and RyR2 and N541-CaM on RyR2. (B) Relative effect of 10 μM dantrolene (dan) on RyR2 P₀ in the presence of wt- and mutant-CaM. Mean values are indicated by the horizontal bars and S.E.M. by the vertical bars. P values indicate significant differences between wt- and mutant-CaM. (C) Facilitation of dantrolene (10 μM) inhibition by N98S-CaM (•). In the absence of dantrolene (○), N98S-CaM has no inhibiting action on RyR2. Asterisks indicate significantly different than 100% (*P < 0.05; **P < 0.01). The solid curve shows the fit of the Hill equation (see legend to Fig. 1) to the data.


