Contractile Properties and Ca\(^{2+}\) Release Activity of the Sarcoplasmic Reticulum in Dilated Cardiomyopathy

A. D’Agnolo, MD; G.B. Luciani, MD; A. Mazzucco, MD; V. Gallucci, MD; and G. Salviati, MD

Background. We performed a comparative study on Ca\(^{2+}\) release activity of the sarcoplasmic reticulum and calcium sensitivity of contractile apparatus of chemically skinned myocardial fibers obtained from four nonfailing human hearts and 13 excised hearts from patients with idiopathic dilated cardiomyopathy.

Methods and Results. Ca\(^{2+}\) sensitivity of contractile apparatus was studied by following the isometric tension developed by chemically skinned myocardial fibers challenged with solutions of decreasing pCa. Ca\(^{2+}\) release from sarcoplasmic reticulum was monitored indirectly by measurement of the isometric tension developed by skinned fibers challenged with caffeine. We observed no significant difference of Ca\(^{2+}\) sensitivity and cooperativity between normal myocardium (pCa\(_{50}=6.00\pm0.05\); Hill coefficient, n\(_{Hill}=2.07\pm0.10\)) and dilated cardiomyopathy (pCa\(_{50}=6.03\pm0.07\); n\(_{Hill}=2.72\pm0.30\)) when the fibers were stretched to 130% of the resting length. We also found that both in normal myocardium and dilated cardiomyopathy, stretching to 150% of the resting length increased the Ca\(^{2+}\) sensitivity of the contractile system; pCa\(_{50}=6.21\pm0.01\) and 6.13\pm0.04 in normal and dilated cardiomyopathy, respectively, whereas in dilated cardiomyopathy there was a decrease of Hill coefficient with stretching that was not observed in the control group. The caffeine threshold in idiopathic dilated cardiomyopathy was markedly increased compared with the control group, 1.94\pm0.27 mmol/l and 0.29\pm0.04 mmol/l cafeine, respectively, whereas there were no significant differences in the extent and rate of caffeine-induced Ca\(^{2+}\) release.

Conclusions. These results indicate that in idiopathic dilated cardiomyopathy there is no alteration of contractile and regulatory proteins; on the contrary, the gating mechanism of the Ca\(^{2+}\) release channel of sarcoplasmic reticulum is abnormal, suggesting a possible involvement of the excitation–contraction coupling in the pathogenesis of this disease. It should also be taken into account, however, that the increased caffeine threshold in dilated cardiomyopathy would be a result of the enhanced resistance to the skinning procedure secondary to the modification of lipid species and/or content in sarcoplasmic reticulum membrane. (Circulation 1992;85:518–525)

Dilated cardiomyopathy (DCM) is defined as a heart muscle disease of unknown origin characterized by a deficient contractility of ventricular myocardium with dilation of the ventricles and inadequate biventricular hypertrophy, reduced cardiac output and ejection fraction, and increased systemic vascular resistance.\(^1\)

Several theories have been proposed to explain the etiology of the disease. Potential roles for viral infection, in particular for coxsackievirus B,\(^2\) autoimmune damage of myocardium,\(^3–5\) and myocardial damage by ethanol\(^6\) have been suggested. In addition, a selective downregulation of \(\beta_{1}\)-receptor subpopulation\(^7\) and an increase of \(G_{i}\) protein have been reported.\(^8,9\) In summary, it seems likely that DCM is an end-stage condition resulting from various diseases of the myocardium.

The pathogenesis of the contractile failure in DCM is also unknown. Possible mechanisms could involve the contractile and regulatory proteins, the mechanism of excitation–contraction coupling, or both. In this study, we measured the pCa/tension

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\(^2\) Supported by institutional funds from the National Research Council Target project FATMA and by grants from the Ministero della Pubblica Istruzione.

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\(^4\) Received April 23, 1991; revision accepted September 23, 1991.
relation and Ca\(^{2+}\) release from sarcoplasmic reticulum of chemically skinned cardiac fibers from patients with idiopathic DCM. We show here that in DCM there is a reduction of the release of Ca\(^{2+}\) from sarcoplasmic reticulum, whereas there are no alterations of Ca\(^{2+}\) sensitivity of contractile proteins.

**Methods**

**Tissue Procurements**

Samples of the left ventricular myocardium were obtained from the explanted hearts of 13 patients, all male, aged 1–59 years (mean, 36.9±5.76), with class IV idiopathic DCM undergong orthotopic cardiac transplantation. The diagnosis of DCM was made according to established criteria.\(^1\) All patients had undergone complete hemodynamic evaluation. They all had normal coronary arteries and no evidence of valvular or congenital heart disease, and the average cardiac catheterization values obtained before transplantation were: right atrial mean pressure, 10.71±3.05 mm Hg; pulmonary wedge mean pressure, 30.33±4.37 mm Hg; and cardiac index, 2.03±0.25 l/min/m\(^2\). All the patients were on therapy with various combinations of digitalis, diuretics, vasodilators, and antiarrhythmics.

As controls, samples of the right ventricle were obtained from four patients with coronary heart disease undergoing coronary artery bypass graft (CABG) surgery. All patients suffered from angina pectoris with no precedent myocardial infarction. Coronary angiography demonstrated involvement of the left coronary artery; the right coronary artery was not involved, and the cardiac catheterization values were in the normal range. Preliminary experiments showed that in rabbits, right and left ventricular myocardium were comparable with regard to the function of both sarcoplasmic reticulum and contractile and regulatory proteins. Thus, it is possible to compare results obtained from samples of left ventricular myocardium affected by DCM with right normal ventricular myocardium. Biopsies of normal myocardium were taken during the CABG surgery before the aorta was cross-clamped on the right ventricle and were immediately immersed in ice-cold 0.9\% NaCl solution. Hearts affected with DCM were excised after infusion of cold cardioplegia solution into the roots of the cross-clamped aortas and immediately immersed in ice-cold 0.9\% NaCl solution. The dimensions of each biopsy were about 2x5 mm.

**Preparation of Skinned Fibers**

Chemically skinned fibers were prepared as previously reported.\(^10\) Briefly, immediately after collection, each biopsy was attached to a wooden stick at the resting length and chemically skinned by incubation at 0–4°C for 24 hours in 10 ml of skinnning solution (170 mmol/l K\(^+\), 2.5 mmol/l Mg\(^{2+}\), 2.5 mmol/l ATP, 5 mmol/l EGTA, 10 mmol/l imidazole, pH 7.0; all cations were salts of propionate). The skinnning solution was changed at hours 1, 4, 6, 16, and 23. The biopsies were then stored at −20°C in the same solution supplemented with 50% glycerol until used (1–2 weeks).

**Calcium Sensitivity of Contractile Apparatus**

Single bundles were isolated with the help of a dissecting microscope, transferred into a chamber containing 1.0 ml of relaxing solution (170 mmol/l K\(^+\), 2.5 mmol/l Mg\(^{2+}\), 5 mmol/l ATP, 5 mmol/l EGTA, 10 mmol/l imidazole, pH 7.0), and inserted between two clamps, one of which was attached to a tension transducer. The fibers were stretched to 130\% of the resting length. All the experiments were carried out at room temperature (20–23°C).

Calcium sensitivity was measured by following the isometric tension developed by the fibers rinsed in solutions of decreasing pCa (170 mmol/l K\(^+\), 2.5 mmol/l Mg\(^{2+}\), 5 mmol/l ATP, 5 mmol/l EGTA, 10 mmol/l imidazole, pH 7.0, and Ca\(^{2+}\) from 0.8 mmol/l [pCa 7.0] to 4.76 mmol/l [pCa 5.0]). Free Ca\(^{2+}\) and Mg\(^{2+}\) concentrations were calculated according to the method of Orentlicher et al.\(^11\) After tension attained plateau, fibers were released by the addition of 10 mmol/l EGTA and then incubated with the solutions of successively decreasing pCa. The experiments were carried out on fibers stretched to 130\% of the resting length. In seven biopsies of DCM and four normal biopsies, we repeated the experiments with the fibers stretched to 150\% of the resting length. The isometric tension, P\(_0\), developed at a given pCa, was normalized to the maximum tension, P\(_{iso}\), developed at pCa 5.0.

**Ca\(^{2+}\) Release From Sarcoplasmic Reticulum**

Ca\(^{2+}\) release was monitored indirectly by following tension development.\(^12\) Fibers were allowed to accumulate Ca\(^{2+}\) into the sarcoplasmic reticulum by incubating in a pCa 7.0 solution (170 mmol/l K\(^+\), 2.5 mmol/l Mg\(^{2+}\), 5 mmol/l ATP, 5 mmol/l EGTA, 10 mmol/l imidazole, pH 7.0, 0.8 mmol/l Ca\(^{2+}\)). After loading, fibers were rinsed in wash solution (same as relaxing solution but without EGTA; free Mg\(^{2+}\) was 0.1 mmol/l\(^{13,15}\)) and challenged with 20 mmol/l caffeine (an agonist of channels of sarcoplasmic reticulum\(^13,15\)). At this concentration of caffeine, all the sarcoplasmic reticulum Ca\(^{2+}\) is released.\(^18\) After bundles were rinsed in relaxing solution, the sarcoplasmic reticulum was reloaded with Ca\(^{2+}\) and then challenged with stepwise-increasing concentrations of caffeine (from 0.1 mmol/l caffeine) until development of tension was measured (caffeine threshold). Caffeine threshold concentration was defined as the lowest caffeine concentration that induced a tension, P\(_t\), greater than 10\% of P\(_{iso}\) (P\(_t\)=tension in the presence of 20 mmol/l caffeine). We also measured the rate of rise in tension at caffeine threshold concentration and at 20 mmol/l caffeine concentration. At the end of the experiment, fibers were rinsed in a pCa 5.0 solution (170 mmol/l K\(^+\), 2.5 mmol/l Mg\(^{2+}\), 5 mmol/l ATP, 10 mmol/l imidazole, pH 7.0, 4.78
mmol/l Ca\textsuperscript{2+}) to record the maximum Ca\textsuperscript{2+}-activated tension and maximum rate of rise in tension.

**Calcium Uptake**

Calcium uptake of two normal and two DCM biopsies was measured by incubating skinned fibers for 10 minutes in 2.0 ml of a solution containing \textsuperscript{45}Ca\textsuperscript{2+} buffered at pCa 7.0 (specific activity, 50×10\textsuperscript{2} cpm/nmol Ca\textsuperscript{2+}). After incubation, the fibers were rinsed for 5 minutes in a modified wash solution containing 0.25 mmol/l EGTA, 10 mmol/l MgCl\textsubscript{2}, and 50 μmol/l ruthenium red, and the fibers were then solubilized by incubating overnight in 100 μl of 1 N NaOH. Radioactivity was measured by liquid scintillation spectrometry.

Protein concentration was determined with the method of Lowry et al,19 using bovine serum albumin as standard.

**Statistical Analysis**

Results were expressed as mean±SEM. Statistical significance was performed by Student's t test. A value of p<0.05 was considered significant.

The Hill coefficient (n) and K (the Ca\textsuperscript{2+} concentration giving half-maximal activation) were calculated by fitting the data to the equation

\[ Y = \frac{\text{Ca}^{2+}}{(\text{Ca}^{2+})^n + K^n} \]

by the least-squares method, where Y is the normalized P/P\textsubscript{0} tension. The Hill coefficient describes the degree of cooperativity of many steps between calcium binding to troponin C and tension development. Therefore, it does not represent a quantitative measurement of the number of calcium binding sites to the myofilaments. Maximal tension was the tension developed at pCa 5.0 normalized to the cross-sectional area of myocardial fibers (mN/mm\textsuperscript{2}).

**Materials**

Caffeine, ruthenium red, ATP, imidazole, and EGTA were purchased from Sigma Chemical Co., St. Louis, Mo. All chemicals were analytical grade.

**Results**

**Calcium Sensitivity and Cooperativity of Contractile Apparatus**

To analyze the Ca\textsuperscript{2+} affinity of regulatory proteins in DCM, we studied the dependence of isometric tension of human cardiac skinned fibers on free Ca\textsuperscript{2+} concentration. Figure 1 shows a typical experiment of isometric pCa/tension measurement. Incubation of cardiac fibers with increasing concentrations of Ca\textsuperscript{2+} caused an increase in the isometric tension.

Figure 2 shows that the averaged curves representative of the isometric pCa/tension relations of ventricular skinned fibers obtained from 13 patients with DCM and from 4 patients with normal myocardium are sigmoidal. The sigmoidal morphology of the curves stands for the existence of cooperativity between the regulatory mechanisms. Ca\textsuperscript{2+} sensitivity was expressed as pC\textsubscript{50} that is, the Ca\textsuperscript{2+} concentration (expressed as −log([Ca\textsuperscript{2+}])) that gives half maximal tension. Ca\textsuperscript{2+} sensitivity in DCM (pC\textsubscript{50}=6.03±0.07) was similar to that obtained in control myocardium (pC\textsubscript{50}=6.00±0.05) (Table 1), with no statistically significant difference. The cooperativity indexes (Hill coefficient, n) of normal myocardium and DCM were also similar; n\textsubscript{Hill} was 2.07±0.13 in normal myocardium and 2.72±0.30 in DCM, with no statistical difference (Table 1). Maximal tension developed at pCa 5.0 was similar in DCM and in control myocardium (46.9±7.0 and 40.2±3.8 mN/mm\textsuperscript{2}, respectively). Also, the rate of rise of isometric tension developed at pCa 5.0 was similar.
(1.32±0.15 and 1.44±0.16 mN/mm²/sec in DCM and normal heart, respectively). No significant differences of maximal tension developed by DCM fibers were found when measurements were carried out in the presence of 15 mmol/l total EGTA and ATP regenerating system.

It is known that stretching of normal myocardial fibers causes an increase of Ca²⁺ sensitivity²⁰ (Stirling’s law). Thus, we checked whether there was an alteration in the response to the stretching in DCM. As shown in Table 2 and in Figure 3, in both normal human myocardium and DCM, Ca²⁺ sensitivity of fibers stretched to 150% of the resting length (pCa5₀=6.21±0.01 and 6.13±0.04 in normal myocardium and DCM, respectively) was significantly higher than that measured at 130% of the resting length (pCa5₀=6.02±0.07 and 6.01±0.03 in normal myocardium and DCM, respectively). The shift in pCa was 0.24±0.03 in DCM and 0.38±0.16 in the normal group, with no significant differences between the two groups. The cooperativity index did not vary with stretching in normal human myocardium (the values were 2.07±0.10 and 1.95±0.02 at 130% and 150% of the resting length, respectively), whereas it decreased significantly (p<0.01) in DCM (3.42±0.62 and 2.73±0.20 at 130% and 150% of the resting length, respectively) (Figure 3 and Table 2).

Caffeine Sensitivity of Sarcoplasmic Reticulum

Sarcoplasmic reticulum Ca²⁺ release activity was studied by following the effects of caffeine. Caffeine is a well-known activator of the Ca²⁺ release channel of terminal cisternae of the sarcoplasmic reticulum.¹⁶,¹⁷,²¹-²⁵ It is also known that caffeine activates the Ca²⁺-induced Ca²⁺ release, the physiological process involved in excitation–contraction coupling in the heart.¹⁸,²¹,²²,²⁶ The release of Ca²⁺ is usually transient and is dependent on the Ca²⁺ load in the sarcoplasmic reticulum and on the concentrations of Mg²⁺, Ca²⁺, and ATP in the medium.¹⁵,¹⁶,²⁶ The effects of caffeine on Ca²⁺ release from the sarcoplasmic reticulum of skinned fibers of four normal and 13 DCM hearts is summarized in Figures 4 and 5 and Table 3. After the sarcoplasmic reticulum is loaded with Ca²⁺, the addition of caffeine at concentrations higher than a definite threshold causes a tension development (Figure 4).

The caffeine threshold was higher in DCM than in the normal group. The mean caffeine threshold was 0.29±0.03 mmol/l in normal human ventricle and 1.94±0.27 mmol/l in DCM (p<0.01) (Table 3). Fig-

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**Table 1. Ca²⁺ Sensitivity of Normal and Dilated Cardiomyopathy Fibers Stretched to 130% of the Resting Length**

<table>
<thead>
<tr>
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<th>Normal human myocardium</th>
<th>Dilated cardiomyopathy</th>
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<tbody>
<tr>
<td>pCa5₀</td>
<td>6.12±0.07</td>
<td>6.21±0.01*</td>
</tr>
<tr>
<td>n</td>
<td>4.20±0.10</td>
<td>1.95±0.02</td>
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Data represent mean±SEM of 16 bundles from four normal ventricles and 47 bundles from 13 ventricles with dilated cardiomyopathy. Ca²⁺ sensitivity is expressed as pCa5₀, i.e., the Ca²⁺ concentration (expressed as −log[Ca²⁺]) at which half-maximal tension is achieved. The degree of positive cooperativity is expressed by the Hill coefficient, n. *p<0.01; †p<0.005.

**Table 2. Effect of Stretching on Ca²⁺ Sensitivity of Normal and Dilated Cardiomyopathic Fibers**

<table>
<thead>
<tr>
<th></th>
<th>Normal</th>
<th>Dilated cardiomyopathy</th>
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<tbody>
<tr>
<td>pCa5₀</td>
<td>6.01±0.03</td>
<td>6.13±0.04†</td>
</tr>
<tr>
<td>n</td>
<td>2.07±0.10</td>
<td>1.95±0.02</td>
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Data represent mean±SEM of 16 fibers from four normal right ventricles and 27 fibers from seven ventricles with dilated cardiomyopathy. Ca²⁺ sensitivity is expressed as pCa5₀, i.e., the Ca²⁺ concentration (expressed as −log[Ca²⁺]) at which half-maximal tension is achieved. The degree of positive cooperativity is expressed by the Hill coefficient, n. *p<0.01; †p<0.005.
Figure 4. Graphs show caffeine sensitivity of sarcoplasmic reticulum. Fibers were incubated at pCa 7.0 for 30 seconds, rinsed in wash (W) solution, and then challenged with 20 mmol/l caffeine to measure maximum tension (P0). Thereafter, fibers were incubated again at pCa 7.0 for 30 seconds, rinsed in wash solution, and challenged with increasing concentrations of caffeine until tension was recorded. Caffeine threshold was defined as the lowest concentration of caffeine that induced tension development, P, larger than 10% of P0. At the end of the experiment, fibers were rinsed in a pCa 5.0 solution, and tension and rate of rise in tension were measured again. Panel A: normal; panel B: dilated cardiomyopathy. R, relaxing solution.

Figure 5. Distribution of caffeine threshold concentrations in 15 bundles from four normal ventricles and 42 bundles from 13 dilated cardiomyopathy ventricles. Caffeine threshold was measured as described in the legend of Figure 4. □, Normal group; ■, dilated cardiomyopathy.

Table 3. Caffeine Sensitivity of Sarcoplasmic Reticulum in Normal and Dilated Cardiomyopathy

<table>
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<tr>
<th></th>
<th>Normal human right ventricle</th>
<th>Dilated cardiomyopathy</th>
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<tbody>
<tr>
<td>Caffeine threshold mmol/l</td>
<td>0.29±0.03</td>
<td>1.94±0.27*</td>
</tr>
<tr>
<td>P/P0</td>
<td>0.20±0.02</td>
<td>0.18±0.01</td>
</tr>
<tr>
<td>P0/X0</td>
<td>0.71±0.06</td>
<td>0.65±0.07</td>
</tr>
<tr>
<td>V/V0</td>
<td>At threshold 0.08±0.03</td>
<td>0.06±0.01</td>
</tr>
<tr>
<td></td>
<td>At 20 mmol/l caffeine 0.36±0.07</td>
<td>0.32±0.04</td>
</tr>
</tbody>
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Data represent mean±SEM of 15 fibers from four normal right ventricles and 42 fibers from 13 ventricles with dilated cardiomyopathy. Experiments were carried out as described in Figure 1. P, V, tension and rate of rise in tension at caffeine threshold concentration, respectively; P0, tension evoked by 20 mmol/l caffeine; V0, rate of rise in tension developed by incubating at pCa 5.0; X0, tension evoked by incubating at pCa 5.0. *p<0.005.

Figure 5 shows caffeine threshold distribution of normal human ventricular cardiac fibers and DCM: 100% of the fibers of normal myocardium had a caffeine threshold lower than 0.50 mmol/l, whereas 100% of the fibers of DCM had a caffeine threshold higher than 0.50 mmol/l.

A second parameter of comparison of caffeine-induced Ca²⁺ release from the sarcoplasmic reticulum was related to the isometric tension developed by skinned fibers at threshold concentration of caffeine (P) normalized to the maximum tension evoked by 20 mmol/l caffeine (P0), which releases almost all the Ca²⁺ accumulated in sarcoplasmic reticulum. P/P0 is, therefore, an indirect measure of the extent of caffeine-induced Ca²⁺ release at the threshold. Table 3 shows that P/P0 was 0.18±0.01 in DCM and 0.20±0.02 in normal myocardium, with no statistical difference.

A third parameter studied was the rate of rise in tension elicited by caffeine at threshold concentration and at 20 mmol/l, which is assumed to be related to the rate of Ca²⁺ release from the sarcoplasmic reticulum. The values were normalized to the rate of rise in tension measured at pCa 5.0, when the rate of rise in tension is a function only of the diffusion rate of the externally applied Ca²⁺ and when the contractile machinery is fully activated by Ca²⁺. Table 3 shows that the rate of rise in tension—that is, the rate of Ca²⁺ release from the sarcoplasmic reticulum—induced by threshold concentrations of caffeine was 0.08±0.03 and 0.06±0.01 in the normal group and DCM, respectively, with no statistical difference, whereas at 20 mmol/l caffeine, it was 0.36±0.07 and 0.32±0.04 in the normal group and DCM, respectively. In summary, these results indicate that the sarcoplasmic reticulum Ca²⁺ efflux channel is less sensitive to caffeine in DCM than in the normal control group. On the other hand, there were no differences in extent and rate of caffeine-induced Ca²⁺ release between DCM and the normal control group.

Because caffeine-induced calcium release seems to be dependent on the Ca²⁺ loading by the sarcoplasmic reticulum, it is possible that the increased caffeine threshold in DCM could be a result of a lower Ca²⁺ loading of sarcoplasmic reticulum. This possibility was tested by measuring tension devel-
oped by skinned fibers at 20 mmol/l caffeine (P0) normalized to that measured at pCa2+ 5.0 (X0) when the contractile machinery is fully activated by external Ca2+. Since 20 mmol/l caffeine releases all Ca2+ accumulated in the sarcoplasmic reticulum, P0/X0 is an index of Ca2+ accumulated in the sarcoplasmic reticulum.21 Table 3 indicates that P0/X0 was 0.71±0.06 in normal human ventricle and 0.65±0.07 in DCM, with no statistical difference. This conclusion is supported by a direct measure of maximum Ca2+ uptake by sarcoplasmic reticulum. This was done on two biopsies from normal human ventricle and two biopsies from DCM. The maximum Ca2+ uptake at pCa 7.0 was 1.81 and 2.68 nmol Ca2+/mg fiber protein in the normal group and DCM, respectively. In conclusion, the results suggest that differences in sarcoplasmic reticulum Ca2+ loading cannot account for the altered caffeine threshold of DCM.

Discussion

DCM is characterized by myocardial contractile failure.2 Many etiopathogenetic hypotheses have been proposed: the viral,2 immune,3-5 and alcoholic.6 This variety of etiopathogenetic hypotheses suggests that DCM may represent the final stage of various myocardial diseases. The contractile failure may be caused by changes in the contractile apparatus, changes in sarcoplasmic reticulum Ca2+ uptake and release, or both. Our results indicate that in DCM, there is an alteration of Ca2+ release from sarcoplasmic reticulum, whereas contractile activity seems to be unaffected.

Calcium Sensitivity of Contractile Apparatus

We were unable to detect any significant differences in either Ca2+ sensitivity, expressed as pCa50, n, in ventricular skinned fibers obtained from hearts with DCM compared with the normal group. In addition, the maximal tension and the rate of rise in tension, normalized to the cross-sectional area, were similar in normal and DCM fibers. This may indicate that the mechanisms of modulation of the contractile process in DCM are spared. These results are in good agreement with previous results on skinned fibers27 and on papillary muscles,28 showing that force development at different external Ca2+ concentrations was not significantly diminished in DCM. This suggests that, at least in vitro, myocardium from cardiomyopathic hearts can generate a normal contractile force.

Furthermore, we also examined the possibility that the myocardial contractile failure could result from an alteration of the relation between force of contraction and diastolic volume (Frank-Starling’s law). The cellular basis for Starling’s law is still not clear. It has been shown that stretching causes an increase in Ca2+ sensitivity of skinned fibers, as demonstrated by the shift to the left of the pCa/tension relation.20 Recent results show that troponin C is an important factor in producing stretch-related changes of Ca2+ sensitivity of cardiac muscle.29-31 Our results show that in DCM, the effect of stretching on the pCa/tension relation is comparable to the one seen in the normal myocardium, demonstrating that there are no major impairments of the mechanism of modulation of Starling’s law.

Caffeine Sensitivity of Sarcoplasmic Reticulum

Our results show that in DCM, the sensitivity to caffeine of sarcoplasmic reticulum is lower than that of normal cardiac fibers, because higher concentrations of caffeine are required to release Ca2+. Caffeine causes Ca2+ release from the sarcoplasmic reticulum by direct interaction with the sarcoplasmic reticulum Ca2+ release channel.16,17,21-25 There are two possible explanations for the lower caffeine sensitivity: 1) a decreased Ca2+ loading of sarcoplasmic reticulum and 2) a decreased caffeine affinity of the Ca2+ release channel.

It has been reported that Ca2+ release from sarcoplasmic reticulum is dependent on sarcoplasmic reticulum Ca2+ loading.26 Thus, it is possible that the lower caffeine affinity of sarcoplasmic reticulum in DCM is caused by a reduced Ca2+ loading of sarcoplasmic reticulum, which, in turn, is caused by a decreased Ca2+ uptake activity of sarcoplasmic reticulum. This conclusion is supported by physiological and biochemical results. It has been suggested that elevated diastolic pressures in DCM might reflect an abnormality of left ventricular relaxation, as indicated by the decrease of the maximum rate of pressure decline (peak negative dP/dt) and of the mean velocity of circumferential fiber lengthening.32 A decreased rate of decline of myoplasmic Ca2+ in muscle from failing hearts has also been reported.33 These results are consistent with other studies that report a decrease in sarcoplasmic reticulum Ca2+ uptake activity on heart homogenates from patients with idiopathic DCM.34 Other studies, however, were unable to detect a decrease in sarcoplasmic reticulum Ca2+ uptake activity in isolated vesicle preparations.35 Our results suggest that in DCM, sarcoplasmic reticulum Ca2+ uptake activity is not decreased. In fact, the amount of Ca2+ released by 20 mmol/l caffeine, a concentration that releases almost all the accumulated Ca2+,12 is not significantly different from control myocardium. Moreover, the measurement of the amount of Ca2+ loaded into the sarcoplasmic reticulum of two biopsies from DCM was similar to that of control myocardium. These results are in agreement with the results of calcium uptake activity of sarcoplasmic reticulum in vesicle preparations,35 suggesting that abnormal Ca2+ handling in DCM is not the result of alteration of sarcoplasmic reticulum Ca2+ uptake. In conclusion, the most likely explanation for the higher caffeine threshold found in DCM seems to be a decreased caffeine affinity of sarcoplasmic reticulum Ca2+ release channel. The possibility cannot be ruled out, however, that the increased caffeine threshold is a result of enhanced resistance to the skimming procedure secondary to the modifica-
tion of lipid species and/or content in sarcoplasmic reticulum membrane.

On the other hand, we did not find differences between DCM and the normal group with regard to the rate and extent of Ca\(^{2+}\) release induced by caffeine either at threshold concentrations or at 20 mmol/l caffeine. These findings suggest that in DCM, there are no changes in the number and conductance of caffeine-sensitive channels of sarcoplasmic reticulum.

These results may have physiological significance, considering that caffeine activates the Ca\(^{2+}\)-induced Ca\(^{2+}\) release mechanism,\(^{18,21,22,26}\) Ca\(^{2+}\)-induced Ca\(^{2+}\) release is the mechanism by which Ca\(^{2+}\) release from the sarcoplasmic reticulum is triggered by a rapid increase of free Ca\(^{2+}\) at the outer surface of the sarcoplasmic reticulum.\(^{36-38}\) A recent study on isolated cardiomyocytes indicates that Ca\(^{2+}\) entry through the L-type of sarcolemmal Ca\(^{2+}\) channel is required to trigger the release of Ca\(^{2+}\) from the sarcoplasmic reticulum.\(^{39}\) Caffeine activates the Ca\(^{2+}\)-induced Ca\(^{2+}\) release by interacting directly with the Ca\(^{2+}\) release channel regulatory sites of the sarcoplasmic reticulum Ca\(^{2+}\)-release channel, increasing their affinity for Ca\(^{2+}\).\(^{16,17,21,22,27}\) Since Ca\(^{2+}\) and caffeine are thought to activate the same mechanism—the Ca\(^{2+}\)-induced Ca\(^{2+}\) release,\(^{18,21,22,26}\) our results suggest that in DCM, higher free Ca\(^{2+}\) concentrations are required for maximum activation of the sarcoplasmic reticulum Ca\(^{2+}\) release channel.

In conclusion, our results demonstrate that in DCM, the Ca\(^{2+}\) sensitivity of the contractile apparatus is unaffected, whereas the gating mechanism of the sarcoplasmic reticulum Ca\(^{2+}\) release channel is abnormal, suggesting a possible involvement of the excitation–contraction coupling in the pathogenesis of this disease.

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KEY WORDS: dilated cardiomyopathy • sarcoplasmic reticulum • contractility • calcium
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_Circulation_. 1992;85:518-525
doi: 10.1161/01.CIR.85.2.518

_Circulation_ is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
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Print ISSN: 0009-7322. Online ISSN: 1524-4539

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