Aggravation of cardiac myofibroblast arrhythmogeneicity by mechanical stress

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Aims

Myofibroblasts (MFBs) as appearing in the myocardium during fibrotic remodelling induce slow conduction following heterocellular gap junctional coupling with cardiomyocytes (CMCs) in bioengineered tissue preparations kept under isometric conditions. In this study, we investigated the hypothesis that strain as developed during diastolic filling of the heart chambers may modulate MFB-dependent slow conduction.

Methods and results

Effects of defined levels of strain on single-cell electrophysiology (patch clamp) and impulse conduction in patterned growth cell strands (optical mapping) were investigated in neonatal rat ventricular cell cultures (Wistar) grown on flexible substrates. While 10.5% strain only minimally affected conduction times in control CMC strands (+3.2%, n.s.), it caused a significant slowing of conduction in the fibrosis model consisting of CMC strands coated with MFBs (conduction times +26.3%). Increased sensitivity to strain of the fibrosis model was due to activation of mechanosensitive channels (MSCs) in both CMCs and MFBs that aggravated the MFB-dependent baseline depolarization of CMCs. As found in non-strained preparations, baseline depolarization of CMCs was partly due to the presence of constitutively active MSCs in coupled MFBs. Constitutive activity of MSCs was not dependent on the contractile state of MFBs, because neither stimulation (thrombin) nor suppression (blebbistatin) thereof significantly affected conduction velocities in the non-strained fibrosis model.

Conclusions

The findings demonstrate that both constitutive and strain-induced activity of MSCs in MFBs significantly enhance their depolarizing effect on electrotonically coupled CMCs. Ensuing aggravation of slow conduction may contribute to the precipitation of strain-related arrhythmias in fibrotically remodelled hearts.

Keywords

Arrhythmia • Slow conduction • Fibrosis • Myofibroblast • Strain

1. Introduction

Old age, genetic predisposition, and insults to the heart like mechanical overload and infarction are well-established causes of fibrotic remodelling of the working myocardium.1 Remodelled tissue is characterized by the presence of excess amounts of collagen that compromises mechanical pump function and promotes arrhythmogenesis by disruption of the normally uniform electrical substrate for impulse propagation.2 Excess secretion of extracellular matrix proteins is primarily attributed to ‘activated’ fibroblasts (myofibroblasts, MFBs) that appear in the working myocardium of diseased hearts.3 Apart from contributing to structural tissue remodelling, MFBs have been shown to exert direct arrhythmogenic effects on cardiomyocytes (CMCs) following the establishment of heterocellular gap-junctional coupling based on connexin 43 (Cx43) and connexin 45 (Cx45) in cell culture systems.4 In the presence of heterocellular gap-junctional coupling, experiments showed that depolarizing current flow from moderately polarized MFBs to well-polarized CMCs causes the latter to undergo partial depolarization thereby inducing slow conduction, precipitation of ectopic activity, and initiation of re-entrant arrhythmias.5–7

In contrast to cell cultures grown on rigid substrates that undergo isometric contractions, intact cardiac tissue exhibits phasic length changes during the pump cycle with maximal strain present in end-diastole. With few exceptions, it has generally been found that stretching healthy cardiac tissue within physiological limits (from slack length to the length of maximal tension development) causes a proportional increase in conduction velocities ($\theta$) when measured in observer coordinates ($\theta_{\text{observ}}$) as obtained, e.g. from optical recordings, while conduction...
times between two specific reference points within the stretched tissue \((C_{\text{prep}})\) remain largely unchanged (for detailed review cf.\cite{8}). While these findings are normally discussed in the framework of CMC stretch sensitivity, work by Kamkin et al. suggested that cardiac fibroblasts may be involved in the response of the myocardium to stretch as well. They reported that mechanical stress causes changes in fibroblast polarization due to activation of mechanosensitive channels \((\text{MSCs})\) that may, if electrotonically coupled to CMCs, affect the electrophysiology of the latter.\cite{9} Similarly, Kohl et al. have shown in a model that strain-induced fibroblast depolarization may affect the discharge rate of electrotonically coupled sinoatrial pacemaker cells, thereby contributing to the adaptation of the heart rate to atrial filling pressure.\cite{10}

Given these findings, the question arises to which extent MFBs may modulate conduction velocities in diseased fibrotic myocardia subjected to stretch and relaxation. Because a direct investigation of this question in intact tissue exhibiting a complex cellular composition and architecture is not feasible with presently available methodologies, we developed an in vitro cell culture system with controlled geometry and defined cellular composition that permits the direct assessment of the differential contribution of CMCs and MFBs to changes of \(\theta\) during application of defined levels of stretch and relaxation. Using this model, we investigated the hypotheses that (i) immunocytochemically defined cardiac MFBs respond to stretch with a reduction in membrane polarization, that (ii) stretch-induced depolarization of MFBs aggravates slow conduction in strands of electrotonically coupled CMCs, and that (iii) mechanical strain exerted by contractile MFBs on adjacent CMCs may directly affect propagation. Our findings show that physiological levels of longitudinal fibre strain as observed in intact healthy tissue during diastolic filling \((< 10\%)\) have no significant effects on \(C_{\text{prep}}\) in preparations consisting predominantly of CMCs. In the presence of electrotonically coupled MFBs, however, \(C_{\text{prep}}\) increases proportionally to applied stretch, suggesting that MFBs act as potent sensors of mechanical stress that cause substantial slowing of conduction in electrotonically coupled CMCs. Such MFB-mediated sensitization of fibrotic cardiac tissue to strain may contribute to the mechanistic understanding of arrhythmogenesis in fibrotically remodelled hearts subjected to mechanical stress.

2. Methods

2.1 Cell culture model

Experiments were conducted in agreement with the relevant institutional and Swiss Federal guidelines for animal experimentation. Primary cultures of 1-day-old Wistar neonatal rat ventricular CMCs and MFBs were established using previously published methods.\cite{12} Animals were decapitated, hearts removed, and the ventricular tissue was dissociated with trypsin. The resulting cell suspension containing CMCs and non-CMCs was sub-hearts removed, and the ventricular tissue was dissociated with trypsin. This preparation was used to quantify the time course of the change in the wrinkle area during experiments. For stretch/relaxation experiments, recordings were obtained within 25 min after application of defined levels of strain.

2.2 Optical measurement of impulse conduction

Impulse propagation along strand preparations was assessed optically using the voltage sensitive dye di-8-ANEPPS \((\text{Biotium})\). Experiments were conducted at 36 °C and signals were recorded after pre-stimulation of the preparations for 10 s at 2 Hz. Recordings were made with a \( \times \) 20 objective \((\text{Spatial resolution: } 50 \mu \text{m})\). Signals acquired at 20 kHz were digitally filtered prior to analysis \((f_{\text{c}}: 0.5 \text{ kHz})\). Optically recorded action potential amplitudes were normalized \%APA\) and maximal upstroke velocities \((dv/dt_{\text{max}})\) were calculated in units of \%/APA/ms. For the case of an APA of 100 mV, values given in units of \%/APA/ms are identical to those in units of V/s.

2.3 Patch-clamp recording

Cell electrophysiology of CMCs and MFBs was assessed using standard whole-cell patch-clamp techniques \((\text{HEKA EPC-10})\). Signals were filtered \((1 \text{ kHz})\), digitized \((2.9 \text{ kHz})\), and stored for off-line analysis. The pipette filling solution contained \((\text{in mmol/L})\): K-aspartate \(120\), NaCl \(10\), MgATP \(3\), CaCl\(_2\) \(1\), EGTA \(10\), and HEPES \(5\) \((\text{pH 7.2})\). Pipette resistances ranged from 4 to 6 M\(\Omega\). Series resistance and, after rupturing of the patch, cell capacitance were compensated and voltage values were corrected for liquid junction potentials \((\text{for } 12.4 \text{ mV})\). \(I-V\) relationships of CMCs and MFBs were obtained with voltage ramp protocols. Whole-cell currents were normalized to cell capacitance and are reported as pA/pF. For stretch/relaxation experiments, recordings were obtained within 25 min after application of defined levels of strain.

2.4 Wrinkle experiments

Substrates for the wrinkle assay \((\text{Excellness, Switzerland})\) were coated with collagen type IV, sterilized with UV, and coated with MFBs at low density \((100 \text{ cells/mm}^2)\). During experiments, preparations were continuously superfused at 36 °C with Hank’s balanced salt solution \((\text{HBSS})\) containing 1% neonatal calf serum \((\text{NCS})\). Time lapse video recordings served to assess the changes of wrinkle patterns following drug addition. A custom-made software \((\text{MatLab; The MathWorks, Natick, MA, USA})\) was used to quantify the time course of the change in the wrinkle area during experiments.

2.5 Immunocytochemistry

The presence of MFBs was confirmed by staining the preparations for \(\alpha\)-smooth muscle actin and counterstaining the nuclei with DAPI using standard protocols.

2.6 Solutions

In all experiments, preparations were superfused at 2–3 mL/min with HBSS containing 1% NCS and 10 mmol/L of HEPES \((\text{pH 7.40})\). For gadolinium \((\text{Gd}^{3+})\) experiments, a solution devoid of phosphate and bicarbonate was used that contained \((\text{in mmol/L})\): NaCl \(140\), KCl \(5.4\), CaCl\(_2\) \(1.8\), MgCl\(_2\) \(1.2\),
glucose 20, HEPES 5 (pH 7.40). Drugs used were obtained from Sigma, except thrombin (Biopur AG, Switzerland).

2.7 Statistical analysis
Values are given as mean ± SD in the text and in the bar graphs. The number of samples refer to independent experiments. Data were compared using the two-tailed Student’s t-test (homoscedastic or heteroscedastic where appropriate), and differences between data sets were considered significant at P < 0.05.

A detailed description of the methods used can be found in Supplementary material online.

3. Results

3.1 Validation of the experimental model
The effects of stretch and relaxation on action potential propagation were investigated in tissue-engineered cell strands grown on silicone membranes that were fixed to the arms of a sliding digital caliper (Figure 1A). During experiments, the entire assembly was placed on the stage of an inverted microscope, the preparation was superfused at 36°C and cell strands under investigation were stimulated with an extracellular electrode (Figure 1B). A typical example of optically recorded action potential upstrokes during propagated activity along a non-strained CMC cell strand is shown in Figure 1C. Experimental preparations consisted either of CMCs (CMC cell strands exhibiting a low degree of ‘contamination’ with MFBs; Figure 2Aa) or of CMC cell strands uniformly coated with MFBs (CMC–MFB cell strands; Figure 2Ab). Important in the context of patch-clamp experiments, fibroblasts seeded on silicone membranes displayed a rapid phenotype switch to MFBs as shown by abundant expression of α-smooth muscle actin containing stress fibres after 2 days in culture (Figure 2Ac). As shown in Figure 2B in a time series of images recorded with a high resolution camera (2048 × 2048 pixels, Ximea), strand preparations retained their stretched morphology beyond the maximal duration of the experiments. Being initially stretched by 5.5%, the example shown relaxes only slightly by 0.3% during 31 min of maintained stretch. As summarized in Figure 2C, this behaviour was typical for all four types of preparations used in this study, i.e. preparations retained their stretched geometry during prolonged application of static stretch (CMC strands: 0.01 ± 0.31%, n = 16; CMC–MFB strands: −0.07 ± 0.29%, n = 22; single-cell CMC preparations: −0.21 ± 0.70%, n = 21; single-cell MFB preparations: 0.26 ± 0.80%, n = 10; no significant differences among different types of preparations). Optical determinations of the dependence of conduction velocity on strain in CMC and CMC–MFB strands consisted of initial control measurements followed by recordings during constant application of 5% relaxation or 5% stretch, respectively. This was followed by control recordings after returning to initial conditions. Preparations were included in the analysis only if measurements obtained after returning to initial lengths were not statistically different from initial control recordings.

3.2 Effects of acute length changes on impulse conduction velocities and conduction times in cell strands
Conduction velocities in observer coordinates ($\theta_{\text{observ}}$) were assessed at control lengths and immediately after stretching and relaxing the silicone membranes by 5% each (overall length change of 10.5%). As shown in Figure 3A, changing the overall length of CMC cell strands by 10.5% caused $\theta_{\text{observ}}$ to increase significantly by 7.1% from 325.7 ± 26.4 to 348.8 ± 39.9 mm/s ($n = 25$, $P < 0.05$). With 335.8 ± 27.8 mm/s, $\theta_{\text{observ}}$ recorded under non-strained control conditions fell exactly between $\theta_{\text{observ}}$ measured during relaxation and stretch, respectively. Maximal upstroke velocities (dV/dt$_{\text{max}}$) of propagating action potentials in CMC cell strands were not significantly affected by either manoeuvre (control: 71.4 ± 3.0%APA/ms; 5% stretch: 71.6 ± 5.0%APA/ms; 5% relaxation: 71.5 ± 4.0%APA/ms; $n = 25$, n.s.). A completely opposite behaviour was observed in CMC–MFB cell strands where changing the overall length by 10.5% caused $\theta_{\text{observ}}$ to decrease significantly by -12.5% from 291.4 ± 48.7 mm/s to 255.0 ± 46.6 mm/s ($n = 20$, $P < 0.005$). With 276.7 ± 44.2 mm/s, $\theta_{\text{observ}}$ recorded under non-strained control conditions fell between $\theta_{\text{observ}}$ measured during relaxation and stretch, respectively. Maximal upstroke velocities were not significantly affected by the different strain conditions (control: 45.9 ± 13.4%APA/ms; 5% stretch: 45.6 ± 13.4%APA/ms; 5% relaxation: 47.6 ± 11.3%APA/ms; $n = 20$, n.s.).

Because preparations underwent defined levels of stretch and relaxation, changes of conduction times in preparation coordinates (CT$_{\text{prep}}$)
during stretch and relaxation relative to control can be inferred by normalizing \( \theta_{\text{observed, intervention}} \) to \( \theta_{\text{observed, control}} \), followed by multiplication of the results with the length changes applied (0.95 for relaxation and 1.05 for stretch). The respective data shown in Figure 3B demonstrate that conduction times in CMC cell strands were not significantly affected by an overall lengthening of 10.5%. In contrast, the same amount of strain caused a highly significant increase of CT\(_{\text{prep}}\) in CMC–MFB cell strands (\( +26.3\% \); \( P < 0.0001 \)).

### 3.3 Strain-induced mechanosensitive currents in MFBs and CMCs

To investigate whether modulation of conduction by strain is based on changes in resting polarization of CMCs and/or MFBs secondary to activation of MSCs, we assessed to which extent basic electrophysiological characteristics of the two cell types are affected by stretch. For this purpose, cells were grown at low densities on silicon membranes and were subjected to whole-cell patch-clamp recordings either under non-strained conditions or immediately after stretching the substrate by 5% (for detailed clamp protocols cf. Supplementary material online). In accordance with the protocols used for optical experiments, patch-clamp recordings were limited to 25 min following onset of static stretch. Mean I–V relationships obtained from MFBs under control conditions (\( n = 13 \)) and during application of 5% stretch (\( n = 8 \)) are shown in Figure 4Aa and b with Figure 4Ac, showing the superposition of the two I–V relationships. As can be gathered from the difference between the two relationships (Figure 4Ad), 5% stretch induced an outward rectifying current that reversed polarity at \(-12.8\) mV. At \(-65\) mV, a potential typically observed in CMC coupled to MFB,\(^{13}\) stretch nearly doubled the inward current present under control conditions from \(-0.20 \pm 0.11\) to \(-0.36 \pm 0.3\) pA/pF. Neither cell capacitance (125.7 \pm 67.9 vs. 125.3 \pm 74.5 pF) nor input resistance (1.5 \pm 0.6 vs. 1.1 \pm 0.6 GΩ) of MFBs were significantly affected by stretch. The same type of experiments conducted with CMCs yielded, as illustrated by the stretch-induced difference current in Figure 4Bd (difference between the mean I–V relationships of 12 control and 13 stretched cells), a more complex response with no consistent stretch-induced changes at
potentials positive to −50 mV. Below this value, 5% stretch induced an inward current that reached a maximum at −76 mV (−0.2 pA/pF) before declining again and turning into an outward current at potentials negative to −88 mV. Cell capacitance of CMCs was not significantly affected by stretch (18.4 ± 6.6 vs. 17.6 ± 9.6 pF), whereas their input resistance increased from 1.0 ± 0.5 to 2.4 ± 0.7 GΩ (P < 0.005). As...
shown in Figure 4C, strain-induced inward currents caused a significant reduction in the membrane potential of MFBs from $-35.6 \pm 5.9 \text{ mV} \ (n = 13)$ to $-26.1 \pm 3.1 \text{ mV} \ (n = 8; P < 0.001)$ and CMCs from $-78.0 \pm 3.2 \text{ mV} \ (n = 12)$ to $-70.8 \pm 8.3 \text{ mV} \ (n = 13; P < 0.05)$.

### 3.4 Contribution of MSCs to conduction under non-stretched conditions

The finding that relaxation of CMC–MFB cell strands caused a significant decrease of conduction times suggested that MSCs of MFBs were contributing to conduction also under non-stretched control conditions. Accordingly, we investigated the effects of the MSC blockers streptomycin (SM, 50 μmol/L) and Gd3+ (50 μmol/L; dissolved in appropriate low-phosphate buffers) on the membrane voltage of CMCs and MFBs cultured under non-stretched conditions. As shown in Figure 5A, the drugs had no effect on the resting polarization of CMCs (SM: $-75.4 \pm 1.3 \text{ mV; } n = 6$, n.s.; Gd3+: $-76.1 \pm 2.9 \text{ mV; } n = 11$, n.s.), indicating that MSCs of CMCs were not active under non-stretched conditions. In contrast, both MSC blockers caused MFBs to undergo a significant hyperpolarization (SM: $-29.4 \pm 7.8 \text{ mV; } n = 9$, $P < 0.005$; Gd3+: $-32.2 \pm 11.6 \text{ mV; } n = 9$, $P < 0.005$), suggesting that MSCs contribute importantly to the membrane polarization of MFBs under non-stretched conditions. In accordance with these single-cell data and as shown in Figure 5B, SM had no effect on $u$ (control: $336.3 \pm 24.4 \text{ mm/s; } n = 58$, n.s.) and $dV/dt_{\text{max}}$ (control: $75.2 \pm 6.1\%\text{APA/ms; } n = 58$, n.s.) in non-stretched CMC cell strands. In contrast, SM caused a significant increase of both $u$ (173.8 $\pm$ 68.2 to $224.9 \pm 63.3 \text{ mm/s; } n = 54; P < 0.005$) and $dV/dt_{\text{max}}$ (29.8 $\pm$ 13.3 to $38.5 \pm 14.3\%\text{APA/ms; } n = 54; P < 0.005$) in CMC–MFB cell strands. Because Gd3+ was substantially more effective in hyperpolarizing MFBs than SM, this compound was expected to surpass the effect of SM in increasing $u$ in CMC–MFB strands. However, both CMC and CMC–MFB strands consistently showed conduction blocks in the presence of Gd3+, which is likely explained by its capacity to cause significant depression of sodium and calcium inward.
currents in CMCs. Overall, the results demonstrate that MFBs exhibit basal MSC activity under non-strained conditions that adds to their depolarized phenotype and, accordingly, contributes to slow conduction in CMC–MFB cell strands also under non-strained control conditions.

3.5 Effects of acute changes in the contractile state of MFBs on their membrane potential

Tension development is a characteristic feature of MFBs and, in the heart, is well established to be responsible for infarct scar consolidation. The finding of basal MSC activity in non-strained MFBs raises the question of whether tension exerted by contractile MFBs on their substrate may feed back onto their own cell membrane in a manner that causes auto-activation of MSCs. This hypothesis was investigated by exposing MFBs to a blocker (blebbistatin) and an activator (thrombin) of MFB contractility. The extent and time course of change of tension development by MFBs following addition of these drugs was assessed using a wrinkle assay, where the extent of deformation of the extracellular substrate (wrinkle formation) is a qualitative measure of tension exerted by adherent cells on the substrate. In the example shown in Figure 6A, a single MFB caused extensive wrinkles in the substrate under control conditions. Following addition of 10 μmol/L blebbistatin to the superfusate, the area occupied by wrinkles declined by ~50% within 10 min which reflects the drug-induced decrease of tension exerted by MFBs on their substrate (cf. also corresponding time lapse movie in Supplementary material online). Overall and as shown in Figure 6B, exposure of MFBs to 10 μmol/L of blebbistatin for 20 min caused a significant reduction in the wrinkle area by 30.3 ± 23.7% (n = 9, P < 0.005). Vice versa, exposure to 1 U/mL of thrombin caused the wrinkle area to increase by 75.2 ± 43.1% (n = 10; P < 0.005), which reflects a substantial increase of tension developed by MFBs. Time courses of the decrease (blebbistatin) and increase (thrombin) of the wrinkle area are shown in Supplementary material online, Figure S1. Determination of the resting polarization of MFBs exposed for a similar time to identical concentrations of the drugs demonstrated that blebbistatin had no significant effect on the membrane potential (control: −31.6 ± 11.2 mV;
blebbistatin: $-34.2 \pm 11.8$ mV; $n = 9$, n.s.), whereas thrombin, contrary to the hypothesis, caused a hyperpolarization despite increasing the tension developed by MFBs (control: $31.0 \pm 6.4$ mV; thrombin: $-34.2 \pm 9.2$ mV; $n = 9, P < 0.05$). These findings suggest that MFB contractility does not directly contribute to basal activity of MSCs in non-constrained preparations by auto-activation of these channels.

3.6 Effects of modulation of MFB contractility on impulse propagation in hybrid cell strands

For hybrid cell strands consisting of MFBs cultured on top of CMCs, mechanical tension developed by MFBs is likely to be transmitted by

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**Figure 6** Modulation of MFB tension development by blebbistatin and thrombin. (A) Phase contrast images of wrinkle patterns before and after exposure of the preparation to blebbistatin (10 μmol/L). The panel below illustrates the time course of the reduction of the area occupied by wrinkles. (B) Effects of blebbistatin (left panels) and thrombin (right panels) on wrinkle formation and membrane potential of cardiac MFBs.
adhesion junctions to underlying CMCs. Ensuing forces acting on the cell membrane of CMCs may modulate conduction by activation of MSCs in CMCs. Accordingly, increasing (thrombin) or decreasing (blebbistatin) tension developed by MFBs attached to CMCs is expected to slow and accelerate conduction, respectively. As shown in Figure 7A, exposure of CMC cell strands to 10 μmol/L of blebbistatin for ≥20 min caused a slight reduction of $u$ from 375.7 ± 35.6 to 365.3 ± 42.9 mm/s ($n = 47; P < 0.05$) that was accompanied by an equally slight decrease of $dV/dt_{\text{max}}$ from 78.8 ± 5.6 to 76.7 ± 10.1%APA/ms ($n = 47; P < 0.05$). In CMC–MFB cell strands, blebbistatin had no effects on $u$ (control: 230.6 ± 34.8 mm/s; blebbistatin: 225.2 ± 35.1 mm/s; $n = 55$; n.s.) and $dV/dt_{\text{max}}$ (control: 52.2 ± 10.4%APA/ms; blebbistatin: 51.8 ± 7.8%APA/ms; $n = 55$; n.s.). As shown in Figure 7B, exposure of CMC cell strands to thrombin at 1 U/mL for ≥20 min had no effects on $u$ (control: 352.7 ± 34.2 mm/s; thrombin: 348.2 ± 44.1 mm/s; $n = 45$; n.s.) and $dV/dt_{\text{max}}$ (control: 74.8 ± 8.5%APA/ms; thrombin: 73.4 ± 10.9%APA/ms; $n = 45$; n.s.). Similarly, in CMC–MFB strands, thrombin did not affect $u$ (control: 246.6 ± 53.8 mm/s; thrombin: 258.6 ± 45.9 mm/s; $n = 47$; n.s.) nor $dV/dt_{\text{max}}$ (control: 45.8 ± 14.4%APA/ms; thrombin: 49.8 ± 11.9%APA/ms; $n = 47$; n.s.). The finding that neither blebbistatin nor thrombin affected conduction in hybrid cell strands suggests that tension exerted by MFBs on adjacent CMCs is not sufficient to modulate MSC activity of CMCs to an extent large enough to affect impulse conduction in hybrid CMC–MFB preparations.

4. Discussion

The results of this study demonstrate that impulse conduction in bioengineered strands of CMCs responds to moderate levels of relaxation and stretch highly similar to intact tissue, i.e. conduction times remain largely unchanged for physiological length changes as those encountered in end-diastole where maximal strain along the direction of the fibres amounts to ~10%. In contrast, in the presence of MFBs simulating a fibrotically remodelled myocardium, impulse propagation is rendered highly sensitive to strain as reflected by a 3% increase of conduction time per each percent of lengthening. Given that the myocardium is maximally strained in end-diastole, these findings suggest that adverse electrotonic interactions between MFBs and CMCs are equally maximal at the time of electrical activation of the myocardium, thereby causing aggravation of slow conduction beyond values reported before for models of the fibrotically remodelled myocardium kept under isometric conditions.
undertaken in the present study, a similar U-shaped mechanosensitive current observed in murine ventricular CMC was shown to be due to stretch-dependent inactivation of $I_{K1}$, which would explain the observation that strain significantly increased the input resistance of CMCs.\textsuperscript{23,24} Assuming that CMCs forming the cell strands underwent the same degree of strain-induced depolarization, the finding that it had no major effect on $\theta$ is likely explained by the circumstance that conduction in cardiac tissue is little affected by changes in resting polarization in the range of $-80$ to $-70$ mV because of the presence of supernormal conduction.\textsuperscript{25-27} This is illustrated schematically in Figure 8, where the grey curve depicts the dependence of conduction velocity on the resting membrane potential of CMCs as derived from a previous study with cultured strands of CMCs.\textsuperscript{27} Typical for supernormal conduction, $\theta$ increases with increasing CMC depolarization from approximately $-85$ to $-75$ mV before declining rapidly with further depolarization. When inserting the resting potentials measured in this study for non-stained CMCs (approximately $-78$ mV) and CMCs subjected to 5% strain (approximately $-71$ mV) into this graph, it becomes evident that the substantial and significant depolarization of CMCs induced by 5% strain has a very limited effect on $\theta$ which is reflected by the experimental finding that conduction times remained unchanged during this intervention. Additional factors contributing to the modest dependence of $\theta$ on strain may include stretch-dependent modulation of sodium currents, membrane capacity, and intercellular resistance.\textsuperscript{8,28-30} Important for this study, the results show that conduction in bioengineered strands of neonatal CMCs behaves identical to intact cardiac tissue from adult animals, i.e. conduction remains largely unchanged during application of physiological levels of strain which suggests that this preparation is a suitable model for investigating the effects of strain on conduction at the cellular level.\textsuperscript{8,31,32}

4.2 Effect of strain on conduction in the fibrosis model

In contrast to the lack of significant effects of strain on conduction in CMC cell strands, applying identical strains to hybrid CMC–MFB cell strands caused a significant increase of conduction times by 26.3%. This contrasting result suggests that MFBs ‘sensitize’ CMCs to strain. The combination of three basic mechanisms as summarized schematically in Figure 8 is likely to explain this observation: (i) as shown before, gap junctional coupling of moderately polarized MFBs to well-polarized CMCs causes convergence of the membrane potentials of the two cell types based on electronic current flow from MFBs to CMCs.\textsuperscript{1} The resulting depolarization of CMCs to values less negative than $-70$ mV pushes non-stained networks of CMCs beyond membrane potentials supporting the peak of supernormal conduction, i.e. into a range where every additional depolarization further slows conduction due to increasing levels of sodium channel inactivation (Figure 8—green dashed lines).\textsuperscript{25} (ii) As shown in this study, such additional depolarization is produced in CMCs by activation of MSC-dependent inward currents at potentials below $-50$ mV that aggravate the pre-existing, MFB induced depolarized state of CMCs. (iii) Additionally, strain applied to MFBs evokes mechanosensitive inward currents at potentials less negative than $-12.8$ mV. The resulting depolarization of MFBs will accentuate depolarizing current flow to electrotonically coupled CMCs, thereby further reducing the membrane voltage of CMCs (Figure 8—green solid lines). Overall, strain-induced activation of MSCs in both CMCs and MFBs that occur on top of the ‘background’ depolarization of CMCs by electrotonically coupled MFBs is likely to form the basis for the increased sensitivity of the fibrotic tissue model to strain. In
analogy, strain sensitivity of conduction may also be increased in other pathological conditions like hyperkalaemia or ischaemia where cardiac tissue is depolarized beyond membrane potentials supporting the peak of supernormal conduction.

4.3 Constitutive activity of MSCs in MFBs

The finding that relaxation of cell strands by 5% caused a significant decrease of conduction times in hybrid CMC–MFB cell strands, but not in CMC strands, suggests that the membrane polarization of MFBs but not CMCs is affected by basal activity of MSCs present under non-strained conditions. In accordance with previous findings in acutely isolated atrial fibroblasts, the presence of constitutively active MSCs in MFBs but not CMCs was supported by the finding that two blockers of MSCs, SM and Gd³⁺, caused a significant hyperpolarization of MFBs, but failed to affect the membrane voltage of CMCs under non-strained conditions. Consistent with the presence of basal activity of MSCs in MFBs that accentuates membrane depolarization of electrotonically coupled CMCs under non-strained conditions, superfusion of hybrid CMC–MFB cell strands with SM caused a significant increase of θ and dV/dtₘₚₓ, but had no effect on CMC cell strands. While an even larger effect was expected to occur in the presence of Gd³⁺ that caused substantial hyperpolarization of MFBs, the finding that bioengineered preparations consisting predominantly of CMCs rather accurately reproduce the strain sensitivity of intact cardiac tissue consisting of electrotonically interacting CMCs and MFBs, the extent to which these mechanisms are operational in vivo has to await the development of experimental methods suitable to investigate this question directly in intact tissue. This concerns especially the lack of unequivocal proof for the presence of heterocellular electrotonic coupling between MFBs and CMCs in the working myocardium of fibrotically remodelled hearts. Also, it cannot be ruled out that fibroblasts present in intact hearts may add an additional layer of complexity to the response of cardiac tissue to stretch. Finally, the model used is based on neonatal rat ventricular cells. Whereas this limits extrapolations of results to intact human cardiac tissue, the finding that bioengineered preparations consisting predominantly of CMCs rather accurately reproduce the strain sensitivity of intact healthy tissue from adult mammals suggests that the biophysical principles governing the dependence of θ on strain are likely similar.

5. Study limitations

In contrast to intact cardiac tissue undergoing phasic variations of strain, methodological restrictions imposed by the technique used in this study required measurements to be performed under static strain conditions. However, given that mechanosensitive currents of CMCs from rats and humans were reported before to activate promptly with applied strain and to display virtually no time dependence, effects of phasic variations of strain on impulse conduction may be indirectly deduced from the observation that strain—θ relationship presented in this study. Further limitations of the study are related to the question as to which extent the in vitro model reflects the situation in vivo whereas the model used in this study is well characterized in terms of cellular morphology, heterocellular gap junctional coupling, ion current densities, and ratios of cell-to-cell contact of neighbouring MFBs (cf. Supplementary material online).

4.4 Effect of MFB contractility on MSC activity and conduction in the fibrosis model

Given the presence of basal activity of MSCs in non-strained MFBs, the question arises as to whether this activity is due to tonic tensile forces exerted by MFBs that feed back onto their own cell membrane thereby causing ‘auto-activation’ of MSCs. This hypothesis was tested by assessing changes in membrane voltage of MFBs superfused with established activators (thrombin) and blockers (blebbistatin) of MFB contractility. While, as shown by the wrinkle assay, both substances affected tension exerted by cardiac fibroblasts on their substrate according to expectations (increased tension with thrombin and reduced tension with blebbistatin), they did either not affect the membrane potential of MFBs (blebbistatin) or even caused a hyperpolarization (thrombin), which makes it unlikely that tension developed by MFBs causes auto-activation of MSCs.

Consistent with the small effects of blebbistatin and thrombin on the membrane potential of single MFBs, exposure of hybrid CMC–MFB cell strands to these drugs failed to affect θ and dV/dtₘₚₓ. These findings demonstrate that modulation of MFB contractility alone is unable to affect impulse conduction in hybrid CMC–MFB preparations. This conclusion is supported by previous findings showing that communication}

5. Study limitations

In contrast to intact cardiac tissue undergoing phasic variations of strain, methodological restrictions imposed by the technique used in this study required measurements to be performed under static strain conditions. However, given that mechanosensitive currents of CMCs from rats and humans were reported before to activate promptly with applied strain and to display virtually no time dependence, effects of phasic variations of strain on impulse conduction may be indirectly deduced from the strain—θ relationship presented in this study. Further limitations of the study are related to the question as to which extent the in vitro model reflects the situation in vivo: whereas the model used in this study is well characterized in terms of cellular morphology, heterocellular gap junctional coupling, ion current densities, and ratios of cell-to-cell contact of neighbouring MFBs (cf. Supplementary material online).

6. Conclusions

Whereas impulse conduction in healthy cardiac tissue shows little dependence on physiological levels of strain, the results of this study suggest that MFBs present in cardiac tissue undergoing fibrotic remodelling may convey increased strain sensitivity to the tissue with conduction velocities being inversely related to applied strain. Extrapolated to intact fibrotic tissue, this mechanism would imply that impulse conduction is slowest at the moment of electrical activation because activation coincides with the moment of maximal distension of the myocardium in end-diastole. Moreover, in diseased hearts displaying non-uniform mechanics, the mechanism may similarly increase non-uniformities of conduction. Pending verification in intact tissue, both effects would contribute to arrhythmogenesis in fibrotically remodelled myocardia.

Supplementary material

Supplementary material is available at Cardiovascular Research online.
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