Fibrosis in left atrial tissue of patients with atrial fibrillation with and without underlying mitral valve disease

A Boldt, U Wetzel, J Lauschke, J Weigl, J Gummert, G Hindricks, H Kottkamp, S Dhein

Objective: To examine the hypothesis that major extracellular matrix (ECM) proteins are expressed differently in the left atrial tissue of patients in sinus rhythm (SR), lone atrial fibrillation (AF), and AF with underlying mitral valve disease (MVD).

Design: Case-control study.

Patients: 118 patients with lone AF, MVD+AF, and SR.

Main outcome measures: Collagen I, collagen III, and fibronectin protein expression measured by quantitative western blotting techniques and immunohistochemical methods.

Results: Protein concentrations increased in patients with AF (all forms) compared with those in SR (all forms): collagen I (1.15 (0.11) vs 0.45 (0.28), respectively; p = 0.002), collagen III (0.74 (0.05) vs 0.46 (0.11); p = 0.002), and fibronectin (0.88 (0.06) vs 0.62 (0.13); p = 0.08). Especially, collagen I was similarly enhanced both in lone AF (1.49 (0.15) and MVD+AF (1.53 (0.16) compared with SR (0.56 (0.28); both p = 0.01). Collagen III was not significantly increased in lone AF but was significantly increased in AF combined with MVD (0.84 (0.07) both compared with SR (0.46 (0.11); p = 0.01). The concentration of fibronectin was not significantly increased in lone AF and MVD+AF (both compared with SR). Furthermore, there was a similar degree of enhanced collagen expression in paroxysmal AF and chronic AF.

Conclusions: AF is associated with fibrosis. Forms of AF differ from each other in collagen III expression. However, there was no systematic difference in ECM expression between paroxysmal AF and chronic AF. Enhanced concentrations of ECM proteins may have a role in structural remodelling and the pathogenesis of AF as a result of separation of the cells by fibrotic depositions.

METHODS

Patients

The study group consisted of patients with lone AF (n = 56: 30 PAF, 26 CAF) and patients with both AF and MVD (n = 46: 10 MVD-PAF, 36 MVD-CAF). Patients in SR (n = 8) and patients with SR combined with MVD (n = 8) formed the control group and were matched to the AF groups according to age, left atrial size, and left ventricular function. Patients were included in the study only if they had preserved left ventricular function. The lone AF group had a left atrial size of ≤45 mm as assessed by echocardiography. The surgical procedure and the concept of intraoperative ablation of AF have been described in detail.

All patients gave written informed consent. The institutional Ethical Committee approved the study. The investigation conforms with the principles outlined in the Declaration of Helsinki.

Atrial tissue from all patients was obtained from the left atrial free wall near the interatrial septum (about 5 × 5 mm from atriotomy) during cardiac surgery, quickly frozen in liquid nitrogen, and stored at −80°C until use.

Abbreviations: AF, atrial fibrillation; CAF, chronic atrial fibrillation; ECM, extracellular matrix; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; MVD, mitral valve disease; PAF, paroxysmal atrial fibrillation; SR, sinus rhythm
Table 1  Summary of patient characteristics

<table>
<thead>
<tr>
<th></th>
<th>Lone AF</th>
<th>MVD-AF</th>
<th>Control group</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>56</td>
<td>46</td>
<td>15</td>
<td></td>
</tr>
<tr>
<td>PAF</td>
<td>30</td>
<td>10</td>
<td>–1</td>
<td></td>
</tr>
<tr>
<td>CAF</td>
<td>26</td>
<td>36</td>
<td>–1</td>
<td></td>
</tr>
<tr>
<td>Cardiac surgery</td>
<td>IRAAF</td>
<td>MVR+IRA AF</td>
<td>MVR (7)</td>
<td>CABG (6)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>48 (13)</td>
<td>64 (10)</td>
<td>59 (7)</td>
<td>&lt;0.05 lone v MVD+AF</td>
</tr>
<tr>
<td>LVEF (%)</td>
<td>60 (6)</td>
<td>57 (16)</td>
<td>55 (13)</td>
<td>&lt;0.05 SR v lone AF</td>
</tr>
<tr>
<td>Left atrium (mm)</td>
<td>43 (6)</td>
<td>55 (11)</td>
<td>44 (7)</td>
<td>&lt;0.001 lone AF v MVD+AF</td>
</tr>
<tr>
<td>Calcium antagonists (all)</td>
<td>8 (14%)</td>
<td>6 (13%)</td>
<td>3 (19%)</td>
<td>NS SR v MVD+AF</td>
</tr>
<tr>
<td>Digitalis</td>
<td>16 (29%)</td>
<td>32 (69%)</td>
<td>2 (13%)</td>
<td></td>
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<tr>
<td>β Blocker</td>
<td>41 (93%)</td>
<td>45 (98%)</td>
<td>10 (63%)</td>
<td></td>
</tr>
<tr>
<td>Antiarrhythmic drugs</td>
<td>22 (39%)</td>
<td>6 (13%)</td>
<td>0</td>
<td></td>
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<tr>
<td>ACE inhibitors</td>
<td>23 (41%)</td>
<td>43 (93%)</td>
<td>9 (56%)</td>
<td></td>
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<tr>
<td>Spironolactone</td>
<td>1 (2%)</td>
<td>8 (17%)</td>
<td>0</td>
<td></td>
</tr>
</tbody>
</table>

Data are mean (SD) or number (%). ACE, angiotensin converting enzyme; AVR, aortic valve replacement; CABG, coronary artery bypass grafting; CAF, chronic atrial fibrillation; IRAAF, intraoperative radiofrequency ablation of atrial fibrillation; LVEF, left ventricular ejection fraction; MVD, mitral valve disease; MVR, mitral valve repair or replacement; NS, not significant; PAF, paroxysmal atrial fibrillation; SR, sinus rhythm.

Histology
Sirius red staining
Formalin fixed, paraffin embedded atrial tissue sections of 5 μm thickness were deparaffinised in xylol and a descending alcohol sequence (100%, 96%, 75%) and brought into distilled water. Subsequently, the slices were exposed to a picro sirius red solution (Hollborn & Söhne, Leipzig, Germany) for one hour. Tissue sections were then washed in diluted acetic acid for 10 minutes. Slices were dehydrogenated in an ascending alcohol sequence (75%, 96%, 100%) and xylol. Lastly, slices were embedded in a mounting medium (Pertex; Histolab, Vastra Frölunda, Sweden).

Immunohistochemistry
Formalin fixed, paraffin embedded atrial tissue sections of 5 μm thickness were deparaffinised. Subsequently, slices were blocked with 5% milk powder tris buffered saline solution and incubated with primary antibodies: mouse anti-human collagen type I, mouse anti-human collagen type III (both from Medicorp, Montreal, Quebec, Canada), and mouse anti-human fibronectin (Santa Cruz Biotechnology, Santa Cruz, California, USA). The staining steps for collagen type I and fibronectin (APAAP staining method) and for collagen type III (AEC staining method) were performed as described by Dako Corp (Carpinteria, California, USA). The specificity was controlled by omitting the primary antibodies.

Western blot analysis
For electrophoresis, 20 μg of total protein was separated in a sodium dodecyl sulphate polyacrylamide gel and blotted on to a cellulose membrane (Roth, Karlsruhe, Germany). Membranes were blocked in 5% milk powder (Roth) in tris buffered saline with 0.3% tween 20 for one hour. After washing, membranes were incubated with primary antibodies for two hours. Goat anti-human collagen I, goat anti-human collagen III (Santa Cruz Biotechnology) and rabbit anti-human fibronectin (DPC Biermann, Bad Nauheim, Germany) were used as primary antibodies. As a reference we used mouse anti-human glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (Hytest, Turku, Finland). After washing, membranes were incubated with the secondary antibodies rabbit anti-goat IgG (for collagen I and III), goat anti-rabbit IgG (for fibronectin) or rabbit anti-mouse IgG (for GAPDH), all conjugated with horseradish peroxidase, for one hour (all secondary antibodies from Sigma, Deisenhofen, Germany). Subsequently, membranes were developed with Super Signal Reagent (Pierce, Rockford, Illinois, USA).

Densitometric analysis
Immunoblots were exposed to x ray film (Eastman Kodak Co, developed, and analysed by ONE-Dscan 1.0 Software (Scanalytics, Los Angeles, California, USA). The relative amount of collagen or fibronectin (target proteins) in each sample was investigated by comparison of the grey scale value of target proteins with the grey scale signal of GAPDH. The GAPDH value was used as an adjusting factor to assure that the same amount of cellular proteins was analysed in each sample. The ratio of target protein to GAPDH from each patient was used to calculate possible differences in target protein synthesis between the patients.

Statistical analysis
All data are mean (SEM). Statistical evaluation was by two way multivariate analysis of variance with a subsequent post hoc Tukey HSD test. Values of p < 0.05 were considered significant.

RESULTS
Patients
Table 1 summarises the clinical characteristics of the patient population. Patients in the lone AF group were younger than those with MVD+AF and all patients in SR. Left atrial sizes in the lone AF and SR group were smaller than in MVD+AF group.

Histological results
Sirius red and immunohistochemical staining for collagen I, collagen III, and fibronectin from left atrial tissue of patients in SR and patients in AF showed clear differences in the occurrence and deposition of connective tissue. In patients with AF, the muscle bundles were surrounded by thick connective tissue fibres. These thick fibres were also present between the single muscle cells or cardiomyocytes separating cells from each other (fig 1A,C,E,G). In the tissue of the
control group there was only a fine network of collagen between the separate muscle bundles and no connective tissue was deposited between the single cells (fig 1B, D, F, H).

After detecting clear histological differences between patients in SR and patients with AF, we quantified the differences by Western blot techniques. Morphometric analysis of the cells showed that the mean size of the cells did not differ between SR and AF (18.5 (0.5) vs 18.9 (0.4) μm, respectively).

**Influence of AF on connective tissue synthesis**

Western blot analysis of collagen I, collagen III, and fibronectin showed a clear increase of the collagens and to a lower extent of fibronectin in patients with AF (all AF) compared with control group patients in SR (all SR). The protein concentration of collagen type I (1.15 (0.11); n = 101) was significantly (about twofold) increased compared with the SR control group (0.45 (0.28), p = 0.002; n = 16). Likewise, type III collagen was significantly enhanced by about 60% in patients with AF (all forms) (0.74 (0.05); n = 89) in comparison with patients in SR (0.46 (0.11), p = 0.022; n = 16). Fibronectin was also, but to a lesser extent, increased in patients with AF (0.88 (0.06); n = 86) compared with the control group (0.62 (0.13); n = 14), without reaching the level of significance (p = 0.08) (fig 2A). The factors coronary heart disease and age of the patient had no significant influence on the development of fibrosis (coronary heart disease: p = 0.8; age: p = 0.62).
Influence of MVD on ECM synthesis

We also analysed the influence of MVD on collagen and fibronectin synthesis. Comparing patients without MVD (SR and all lone AF) with patients with MVD (MVD+AF and all MVD+AF) we did not detect any significant differences either in collagen type I (1.38 (0.15); n = 64 v 1.59 (0.16); n = 54) or in fibronectin protein concentration (0.87 (0.07); n = 57 v 0.79 (0.08); n = 45). Collagen type III was only slightly increased in patients with MVD (0.79 (0.06); n = 48) compared with patients without MVD (0.62 (0.06); n = 55) with borderline significance (p = 0.045) (fig 2B).

Differences between lone AF and MVD+AF

Subsequently, we determined the influence of lone AF (all forms) and MVD+AF (all forms) on collagen and fibronectin production in comparison with that in patients in SR (with and without MVD). There were significant differences in collagen type I (analysis of variance, p = 0.009). The protein concentration of collagen I was increased in patients with lone AF (1.49 (0.15); n = 56) compared with the SR group (0.56 (0.28), p = 0.01; n = 16) and was also increased in the MVD+AF group (1.53 (0.16), p = 0.01; n = 46). We did not find differences between lone AF and MVD+AF (fig 3A).

Collagen type III concentrations changed significantly in comparing SR versus lone AF versus MVD+AF by analysis of variance (p = 0.01). Especially, the increase was significant in patients with MVD+AF (0.84 (0.07); n = 41), about 80%, compared with SR (0.46 (0.11), p = 0.01; n = 14). No significant changes were detected between SR and lone AF (0.66 (0.06); n = 48) or between lone AF and MVD+AF (fig 3A).

The protein concentration of fibronectin tended to be enhanced in the lone AF group (0.93 (0.08); n = 48) and in the MVD+AF group (0.81 (0.09); n = 38) compared with SR group (0.62 (0.13); n = 16), without reaching the level of significance (analysis of variance, p = 0.13; fig 3C).

Influence of paroxysmal and chronic forms of AF

To investigate the influence of the PAF and CAF forms of AF, we compared patients with lone PAF and lone CAF with the SR group and patients with MVD+PAF and MVD+CAF with patients with MVD+SR.

The protein concentration of collagen type I increased in the left atrium of patients with lone PAF (1.31 (0.21); n = 29) and in patients with lone CAF (1.71 (0.22); n = 26) compared with the SR control group (0.57 (0.4); n = 8). Similarly, in patients with MVD collagen type I concentration increased both in the MVD+PAF group (1.7 (0.36); n = 10) and in the MVD+CAF group (1.48 (0.18); n = 36) compared with the MVD+SR group (0.54 (0.4); n = 8) (fig 4A). However, the differences between the PAF and CAF subgroups did not reach significance.

Regarding collagen type III, we found a (non-significant) slight rise in both lone PAF (0.57 (0.08); n = 25) and in lone CAF (0.77 (0.08); n = 23) compared with SR (0.37 (0.15); n = 7). Furthermore, collagen type III was about twofold increased in the MVD+PAF group (1.14 (0.13), p = 0.04; n = 10) compared with MVD+SR (0.55 (0.15); n = 7), whereas the increase in the MVD+CAF group (0.74 (0.07); n = 36)
The present study shows for the first time a clear relation between human AF and the expression of major ECM components such as collagen I, collagen III, and fibronectin. We found an increase of about 100% in collagen I, an increase of about 50% in collagen III (which was confined to MVD+AF), and a smaller non-significant increase in fibronectin in left atrial tissue samples of patients with AF. Thus, our data seem to confirm a relation between AF and fibrosis in both lone AF and MVD+AF. Furthermore, interstitial fibrosis may be the cause of the electrophysiological and structural changes seen in patients with AF in several investigations. AF is associated with an increase of connective tissue between individual cells and with the deposition of large amounts of collagen and fibronectin. In patients with AF, the degree of fibrosis is augmented and tends to separate myocytes from each other. Advanced interstitial fibrosis in human AF would predict an impairment of atrial conduction at the microscopic level and may render the atrial myocardium discontinuous resulting in a branching structure. Such types of fibrosis may alter the biophysical properties of the tissue allowing the initiation and perpetuation of AF.

Collagen type I expression was significantly enhanced in both patients with lone AF and patients with MVD+AF compared with the control group. There were no significant differences between lone AF and MVD+AF. However, in both groups we observed a threefold increase (compared with SR) in collagen type II. Interestingly, we found an increase in collagen type III expression in patients with MVD+AF but not in patients with lone AF. The increase in patients with lone AF was only slight (about 40%) and non-significant compared with that in SR, whereas collagen III was significantly increased (about 80%) in the MVD+AF group compared with the SR group. Table 2 presents a simplified summary. Furthermore, left atrial diameter was larger in patients with MVD than in either the control patients or patients with lone AF.

These data support the hypothesis that remodelling with increased concentrations of collagen I occurs even in lone AF and that the mechanical stress of MVD causes further changes especially in collagen III. The hypothesis that fibrosis progresses systematically from PAF to CAF was not confirmed in either the lone AF group or the MVD+AF group.
Considering the data of these subgroups there was a slight increase in collagen from PAF to CAF in lone AF; however, the data varied too much to allow for a clear conclusion. It is known that systematic differences between PAF and CAF exist. Thus, patients with CAF exhibit a shorter cycle length and a higher degree of disorganised activity than do patients with PAF.\textsuperscript{24–27} Nevertheless, we could not detect significant differences between PAF and CAF regarding changes in ECM components. This is supported by other studies that also found no differences between PAF and CAF in human AF of different causes, such as AF with MVD, coronary artery disease, and lone AF, regarding other pathophysiological factors such as angiotensin and endothelin.\textsuperscript{26–27}

To the best of our knowledge this is the first study that determined the relation between the degree of fibrosis and ECM components and lone AF in human left atrium in patients with PAF and CAF, with or without MVD. Previous studies based on determinations in animals or in the human right atrial appendage did not consider subgroups such as PAF and CAF.\textsuperscript{28–30} Additionally, the human studies included very mixed patient populations in most cases. Since initiation and perpetuation of AF pathophysiologically depends on the left atrium, it seems important to investigate the left atrium.\textsuperscript{31} In fact, AF itself can lead to atrial dilatation and from our data we believe that AF is associated with the fibrotic process.\textsuperscript{32}

Conclusion

Forms of AF differ from each other in collagen III expression. However, there was no systematic difference in collagen expression between PAF and CAF. Furthermore, AF is associated with fibrosis. On the basis of cell-cell decoupling and isolation of the myocytes caused by fibrosis, we assume that fibrosis may be an important factor in the maintenance and progression of AF. Furthermore, the irreversible deposition of increased amounts of connective tissue in the atrial tissue, resulting in a modified electrophysiology, may support the theory of Wijffels and colleagues\textsuperscript{1} that “atrial fibrillation begets atrial fibrillation”: We conclude that the role of enhanced concentrations of ECM proteins may be important in the pathogenesis of AF due to separation of cells by fibrotic depositions.

Authors’ affiliations

A Boldt, J Gummert, S Dhein, Heart Centre, Cardiovascular Surgery, University of Leipzig, Leipzig, Germany
U Wietzel, J Lauschke, J Weigl, G Hindricks, H Kottkamp, Department of Electrophysiology, Cardiology, Leipzig, Germany

References


