Circulating concentrations of cardiac proteins indicate the severity of congestive heart failure

T Goto, H Takase, T Toriyama, T Sugiura, K Sato, R Ueda, Y Dohi

Objective: To test the hypothesis that myocardium specific proteins may be useful markers for evaluating the severity of congestive heart failure.

Methods: Serum concentrations of myosin light chain I (MLC-I), heart fatty acid binding protein (H-FABP), creatine kinase isoenzyme MB (CK-MB), and troponin T (TnT) and plasma concentrations of brain natriuretic peptide (BNP) were determined in 48 patients with acute deterioration of congestive heart failure, both before and after effective treatment.

Results: Before treatment, MLC-I (mean (SEM) 3.2 (2.2) µg/l), H-FABP (9.0 (3.5) µg/l), TnT (30 (21) ng/l), and BNP (761 (303) ng/l) were higher than the normal reference range, and concentrations of CK-MB (5.4 (2.9) µg/l) were near normal. Treatment of congestive heart failure with conventional medication significantly decreased the concentrations of MLC-I (1.2 (0.3) µg/l, p < 0.0001), H-FABP (6.0 (2.0) µg/l, p < 0.0001), CK-MB (2.9 (1.5) µg/l, p < 0.0001), TnT (9 (1) ng/l, p < 0.001), and BNP (156 (118) ng/l, p < 0.0001). The decreases in H-FABP and CK-MB concentrations after treatment correlated with the decrease in BNP concentrations (p < 0.05). The absolute concentrations of MLC-I, H-FABP, CK-MB, and TnT correlated positively with those of BNP (p < 0.01).

Conclusions: These findings suggest that MLC-I, H-FABP, CK-MB, and TnT may be used as reliable markers for the evaluation of the severity of congestive heart failure.
were: 4.0
range (99th centile of the control population) of the assay
and interassay coefficients of variation, and normal reference
Data are expressed as mean (SEM). A paired Student’s
Statistical analysis
2
ately frozen and stored at
blood sampling in a precooled centrifuge and were immedi-
and plasma samples were prepared within 30 minutes of
TnT, 5 ml of blood was transferred to plastic tubes. Serum
For measurement of MLC-I, H-FABP, CK-MB, and
2
Biochemical measurements
For BNP measurement, 3 ml of blood was transferred to a
lists medications during the stable phase.
the second biochemical evaluation was 52 (5) days. Table 2
evaluation was repeated after stabilisation of clinical findings
and symptoms (stable phase). The time between the first and
Medications
a
Angiotensin converting enzyme inhibitor 19
Angiotensin II type 1 receptor blocker 15
B Blocker 5
β Blocker 8
Calcium channel blocker 9
Diuretics 48
Digitalis 29
Pimobendan 3

<table>
<thead>
<tr>
<th>Table 1 Characteristics of patients</th>
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<tbody>
<tr>
<td>Acute phase</td>
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<tr>
<td>-------------</td>
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<tr>
<td>NYHA class (I/II/III/IV)</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
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<tr>
<td>Systolic blood pressure (mm Hg)</td>
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<td>Diastolic blood pressure (mm Hg)</td>
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<tr>
<td>Heart rate (beats/min)</td>
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<tr>
<td>Cardiacoratic ratio (%)</td>
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<tr>
<td>Left atrial diameter (mm)</td>
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<tr>
<td>Diastolic left ventricular diameter (mm)</td>
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<tr>
<td>Systolic left ventricular diameter (mm)</td>
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<tr>
<td>Ejection fraction (%)</td>
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<td>Fractional shortening (%)</td>
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</tbody>
</table>

*p<0.05, **p<0.001, ***p<0.0001 versus acute phase (paired Student’s t test).

Statistical analysis

Data are expressed as mean (SEM). A paired Student’s t test was used to compare and analyse differences in continuous variables that had a normal distribution between the acute and stable phases. Because the distributions of MLC-I, H-FABP, CK-MB, TnT, and BNP concentrations were skewed rightwards, median concentrations were computed for these parameters and expressed as median value (median absolute deviation). The significance of any difference in medians was assessed by the Wilcoxon signed rank test and correlations between these variables were assessed by the Spearman rank correlation test. A probability value of p < 0.05 was considered significant.

RESULTS

On admission, all patients had worsening CHF with increased cardiothoracic ratios on chest radiographs and decreased left ventricular systolic function on echocardiograms (table 1). The circulating concentrations of BNP were higher than the normal reference range in all patients (fig 1). Biochemical analysis showed that serum concentrations of proteins derived from myocardial cells (MLC-I, H-FABP, and TnT) were higher than the normal reference range, and the concentrations of CK-MB were near normal (fig 1).

All patients were successfully treated, resulting in improvement of their symptoms and physiological findings; the NYHA functional class, cardiothoracic ratio, and findings obtained from echocardiograms all improved (table 1). Effective medical treatment significantly decreased not only BNP but also MLC-I, H-FABP, CK-MB, and TnT (fig 1). The decrease in H-FABP and CK-MB, but not in MLC-I and TnT, concentrations after treatment significantly correlated with the decrease in BNP concentrations (fig 2). In addition, the absolute concentrations of MLC-I, H-FABP, CK-MB, and TnT correlated positively with those of BNP in patients with CHF (fig 3).

DISCUSSION

Structural changes associated with severe CHF are characterised by degeneration of subcellular organelles in viable myocytes and multiple foci of myocardial cell death. Thus, we hypothesised that circulating concentrations of specific cardiac proteins may be useful as markers for the evaluation of the severity of CHF. The present study has shown that circulating concentrations of the cardiac proteins MLC-I, H-FABP, CK-MB, and TnT correlated with the concentrations of BNP in patients with CHF. Since circulating BNP concentrations are increased in patients with CHF in proportion to the severity of the disease, the results of this study suggest that these specific cardiac proteins may be useful as powerful markers for determining the severity of CHF.

In the present study in patients with CHF, the concentrations of MLC-I, H-FABP, CK-MB, and TnT were higher on admission (acute phase) than under the stable conditions achieved after appropriate treatment. The results are compatible with previous reports that circulating concentrations of
Figure 1  Serum concentrations of myosin light chain I (MLC-I), heart fatty acid binding protein (H-FABP), creatine kinase isoenzyme MB (CK-MB), troponin T (TnT), and brain natriuretic peptide (BNP) before (acute) and after effective treatment (stable) in patients with congestive heart failure. Open circles: scattergram showing data in each patient; closed diamonds and vertical bars: median value (median absolute deviation). *p<0.001, **p<0.0001 versus acute phase.

Figure 2  Correlations between the decrease in the concentrations of specific cardiac proteins and BNP after effective treatment of patients with congestive heart failure. The decrease in the concentrations of H-FABP and CK-MB, but not of MLC-I and TnT, correlate with that in BNP.
Troponin I, TnT, and MLC-I are increased in advanced CHF. These proteins are specific to myocardial cells, suggesting that the increase in protein concentrations reflects cellular injury of the myocardium in advanced CHF. The presence of myocardial cytosolic proteins such as H-FABP and CK-MB in serum from patients with severe CHF implies that permeability of the membrane of injured myocytes has been compromised and that the release of the cytosolic proteins into the circulation may be caused by leakage. However, this concept does not explain the appearance of MLC-I or TnT in the circulation, as the majority of these proteins are present in the contractile apparatus and not in cytosol. In severe CHF, myofibrils may degenerate and, subsequently, the myofibrillar components may be released into the circulation. Indeed, degeneration of hypertrophied myocardial cells has been reported in human dilated cardiomyopathy. The mechanisms of ongoing myocardial damage in patients with CHF have not been established, but experimental models provide insight into the cellular and molecular mechanisms that contribute to the progression of CHF. Microcirculatory abnormalities of coronary arteries have been implicated in the aetiologies of human cardiomyopathy. These may be contributing factors to focal myocardial necrosis and replacement fibrosis. Recently, other mechanisms have been identified that also mediate the progressive impairment of cardiac structure and function including neurohormonal factors, oxidative stress, and some cytokines. Each of these factors can promote cardiac cell death by producing either myocyte necrosis or myocyte apoptosis through activation of specific genetic pathways. These structural changes may accelerate left ventricular dysfunction, which increases left ventricular filling pressure resulting in secretion of BNP. Thus, it is hardly surprising that circulating concentrations of these specific cardiac proteins correlate with those of BNP in CHF.

Appropriate treatment greatly decreased the concentrations of MLC-I, H-FABP, CK-MB, and TnT, suggesting that the cardiac markers are quite sensitive to changes in left ventricular haemodynamic dysfunction. Strikingly, the decrease in H-FABP and CK-MB concentrations correlated with the decrease in BNP concentrations. This indicates that concentrations of the marker proteins appear to be related to left ventricular filling pressures and can give clinically important information for the treatment of CHF. Since BNP concentrations may serve as a useful prognostic indicator in patients with CHF, the marker proteins may also provide useful information on the prognosis of CHF. BNP and the marker proteins may have a complementary role in the prognostic assessment because BNP and the marker proteins provide intrinsically different information. However, further studies are necessary to confirm this hypothesis.

The interpretation of the present results is limited by the small number of patients studied. However, the clear correlation between the absolute concentrations of the specific cardiac protein markers and of BNP and between the decrease in the concentrations of the cardiac markers and of BNP shows that cardiac proteins may serve as reliable markers for the evaluation of the severity of CHF or the effect of treatment of CHF. Since BNP concentrations do not appear to indicate the severity of CHF in some cases, measurement of the circulating concentrations of these cardiac proteins in combination with BNP may provide more useful information than measurements of BNP concentrations alone in the management of patients with CHF.

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A 67 year old man with chronic renal failure, hypertension and ischaemic heart disease was admitted for polymorphic ventricular tachycardia with syncope. Echocardiography revealed left ventricular hypertrophy with moderately impaired function. Amiodarone was administered for these arrhythmias, but recurrent episodes were experienced during haemodialysis. Therefore, an implantable cardioverter-defibrillator (ICD) was implanted at the left anterior chest on the other side of the shunt vein. Although the left cephalic vein was too small for insertion of a ventricular lead, an atrial lead was inserted from there, and the ventricular lead was inserted from the left subclavian vein. During the procedure, a persistent left superior vena cava (PLSVC) was detected incidentally, as both leads ran through an unusual left sided approach. The Medtronic model 6945 (65 cm) screw-in ventricular lead was successfully affixed to the right ventricular apex by forming the stylet into a U shape in the right atrium. The Intermedics model 418-35S (52 cm) atrial lead was fixed in the right atrium (see panels). Atrial and ventricular sensing and stimulation thresholds were acceptable. Ventricular fibrillation induced by a T wave shock was successfully defibrillated twice with 20 joules of energy.

PLSVC is present in approximately 0.5% of the population, and a transvenous pacemaker or ICD implantation is sometimes difficult or even impossible in those cases. If transvenous implantation via PLSVC is intended, it is recommended to use a screw-in type electrode with sufficient length so that a loop in the right atrium can be formed for appropriate fixation to the right ventricle.

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