Kallikrein Gene Delivery Improves Cardiac Reserve and Attenuates Remodeling After Myocardial Infarction

Jun Agata, Lee Chao, Julie Chao

Abstract—In this study, we used the somatic gene delivery approach to explore the role of the kallikrein-kinin system (KKS) in cardiac remodeling and apoptosis after myocardial infarction (MI). Rats were subjected to coronary artery ligation to induce MI, and adenovirus carrying the human tissue kallikrein or luciferase gene was injected into the tail vein at 1 week after surgery. Cardiac output gradually decreased from 2 to 6 weeks after MI, whereas delivery of the kallikrein gene prevented this decrease. Cardiac responses to dobutamine-induced stress were improved in rats receiving kallikrein gene as compared with rats receiving control virus at 6 weeks after MI. Kallikrein significantly improved cardiac remodeling by decreasing collagen density, cardiomyocyte size, and left ventricular internal perimeter and increasing capillary density in the heart at 6 weeks after MI. Kallikrein gene transfer attenuated myocardial apoptosis, which was positively correlated with remodeling parameters in the heart at 2 weeks after MI. Endothelial dysfunction, characterized by increased vascular resistance, decreased left ventricular blood flow, and decreased cardiac nitric oxide levels, existed in remodeled hearts at 2 weeks after MI, whereas kallikrein gene transfer improved these parameters. Kallikrein gene delivery improved cell survival parameters as shown by increased phospho-Akt and reduced caspase-3 activation at 2 weeks after MI. This study indicates that the kallikrein-kinin system plays an important role in preventing the progression of heart failure by attenuating cardiac hypertrophy and fibrosis, improving endothelial function, and inhibiting myocardial apoptosis through the Akt-mediated signaling pathway. (Hypertension. 2002;40:653-659.)

Key Words: myocardial infarction ■ remodeling ■ kallikrein-kinin systems ■ genes ■ apoptosis

Angiotensin-converting enzyme inhibition was first shown to improve the survival of coronary ligation–induced myocardial infarction (MI) in a rat model.1 Treatment of the ACE inhibitor enalapril increased survival in rats with congestive heart failure.2 Subsequent studies have confirmed beneficial effects of ACE inhibitors in reduction of morbidity and mortality rates and improvement in the quality of life in patients.3,4 Therefore, treatment based on ACE inhibition has become an established therapy for patients with chronic heart failure (CHF), systolic left ventricular (LV) dysfunction, and MI.5–8 The effects of ACE inhibitors on CHF are mostly attributed to the blockade of angiotensin II (Ang II) production, which induces cardiomyocyte hypertrophy after MI or heart failure.9 It has been shown that Ang II stimulates collagen synthesis in cultured cardiac fibroblasts10 and induces apoptosis in cardiomyocytes and endothelial cells.11–13 Collectively, these observations support a role of Ang II in promoting the progression of cardiac remodeling and suppressing cardiac function. Since ACE is the same enzyme as kininase II, a kinin-degrading enzyme, inhibition of ACE not only results in reduced Ang II levels but also decreased kinin breakdown. This results in the accumulation of kinin in plasma and tissues. Previous reports have shown that a kinin B2 receptor antagonist, icatibant, partially abolished the protective effect of ACE inhibition in cardiac remodeling, implicating a role of kinin in cardioprotection.14–16 Binding of intact kinin to the kinin B2 receptor activates second messengers such as nitric oxide (NO)/cGMP and prostacyclin/cAMP and triggers many biological effects. Therefore, ACE inhibition could also be attributed to kinin-mediated protective effects in cardiac hemodynamics and remodeling. However, the potential mechanisms of the effects of kinin in cardiac remodeling have not been established.

Apoptosis in cardiomyocytes is one of the major factors that contributes to the progression of heart failure after MI.17,18 Recently, we reported that kallikrein gene delivery results in increased cardiac kinins and cGMP levels and reduced apoptosis after acute myocardial ischemia and reperfusion, and the effect is abolished by icatibant.19 However, the role of KKS on cardiac remodeling and apoptosis in progression of CHF is not well elucidated. In the present study, we investigated the potential role and mechanisms of the KKS in cardiac remodeling and apoptosis after MI by a somatic gene transfer approach.

Methods

Preparation of Adenovirus Carrying the Human Tissue Kallikrein Gene

Adenovirus containing the human tissue kallikrein gene under the control of cytomegalovirus promoter (Ad.CMV-cHK) was generated...
as previously described. Large quantities of high-titer Ad.CMV-cHK and control virus containing the luciferase gene (Ad.CMV-Luc) were prepared and purified for in vivo gene delivery.

Animals and Treatments

Wistar rats (male, 225 to 250 g body weight, Sprague-Dawley Harlan, Indianapolis, Ind) were subjected to ligation of the left coronary artery as previously described. One week after coronary artery ligation, the rats were randomly divided into two groups and injected through the tail vein with 1.2×10^10 plaque-forming units of adenovirus harboring either human tissue kallikrein (MI-cHK group) or luciferase gene (MI-Luc group). The sham rats underwent the same surgical procedure without ligation of the left coronary artery. At 2 or 6 weeks after coronary artery ligation, the hemodynamic parameters were analyzed and the animals were then euthanized. Tissues were harvested for morphological and biochemical analyses.

Hemodynamic Parameters

At the end of the study (2 or 6 weeks after coronary artery ligation), rats were anesthetized with sodium pentobarbital (50 mg/kg IP) and cardiac function and regional blood flow were measured by a fluorescent microsphere assay. The right carotid artery (RCA) and the left femoral artery (LFA) were catheterized with PE-50 tubing. The distal end of the cannula of the RCA was connected to a physiological pressure transducer (Statham Laboratories) coupled with a model 7E polygraph (Grass Instrument Co). After measurements of heart rate (HR) and mean arterial pressure (MAP), the tip of the RCA catheter was gently advanced into the LV, and LV end-diastolic pressure (LVEDP) was recorded. Then, 0.2 mL of solution containing fluorescent microsphere beads (2.0×10^10 beads, Molecular Probes, Inc) was injected within 10 seconds into the LV, and regional blood flow (mL/min per g) was determined by the Polytom (Brinkmann Instruments) in 500 μL of buffer (pH 7.4) containing 1% Triton X-100, 0.1% SDS, 2 mmol/L EDTA, and 1% protease inhibitor cocktail (Sigma) and centrifuged at 14 000 rpm for 30 minutes at 4°C. The supernatants were used for Western blot analysis, with specific antibodies used for phospho-Akt, total Akt, and cleaved caspase-3 (Cell Signaling Technology). Nitrite/nitrate levels in the tissue extracts were measured by a fluorometric assay as previously described, and protein concentrations were determined by Lowry’s method.

Statistical Analysis

Values are expressed as mean±SEM. Statistical comparisons were performed with the use of 1-way ANOVA followed by the Fisher’s PLSD test for multiple comparisons. Regression analysis was used to compare the relation between TUNEL-positive myocytes and cardiomyocyte size or LV internal perimeter. A value of $P<0.05$ was considered statistically significant.

Results

Effects of Kallikrein Gene Delivery on Infarct Size and Physiological and Hemodynamic Parameters After MI

Our results show that mean infarct size did not differ significantly between the groups injected with the kallikrein or luciferase gene (37.9±2.3% in MI-Luc versus 38.5±3.2% in MI-cHK at 2 weeks after MI, 42.3±2.5% in MI-Luc versus 43.1±2.9% in MI-cHK at 6 weeks after MI). The Table shows physiological and hemodynamic parameters in hearts at 2 and 6 weeks after MI. There were no differences in BW among the groups at 6 weeks after MI. However, BW in the MI-Luc and MI-cHK groups was reduced as compared with the sham group at 2 weeks after MI. MI resulted in increases in BW, HW/BW, LVV/Wt, and LVV/BW, especially at 6 weeks.
Physiological and Hemodynamic Parameters at 2 and 6 Weeks After Myocardial Infarction

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Sham (n=13)</th>
<th>MI-Luc (n=16)</th>
<th>MI-cHK (n=16)</th>
<th>Sham (n=17)</th>
<th>MI-Luc (n=11)</th>
<th>MI-cHK (n=15)</th>
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</thead>
<tbody>
<tr>
<td>BW, g</td>
<td>319±7</td>
<td>292±10*</td>
<td>276±8*</td>
<td>428±7</td>
<td>414±12</td>
<td>409±7</td>
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<tr>
<td>HW, g</td>
<td>0.87±0.02</td>
<td>0.94±0.03</td>
<td>0.87±0.03</td>
<td>1.01±0.02</td>
<td>1.16±0.04*</td>
<td>1.13±0.02*</td>
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<tr>
<td>HW/BW, g/kg</td>
<td>2.73±0.04</td>
<td>3.26±0.12*</td>
<td>3.17±0.07*</td>
<td>2.35±0.04</td>
<td>2.81±0.09*</td>
<td>2.76±0.04*</td>
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<tr>
<td>LW, g</td>
<td>0.70±0.02</td>
<td>0.72±0.02</td>
<td>0.70±0.02</td>
<td>0.81±0.02</td>
<td>0.90±0.03*</td>
<td>0.86±0.01</td>
</tr>
<tr>
<td>LW/BW, g/kg</td>
<td>2.20±0.03</td>
<td>2.48±0.05*</td>
<td>2.52±0.04*</td>
<td>1.90±0.03</td>
<td>2.19±0.06*</td>
<td>2.10±0.03*</td>
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<tr>
<td>LVIP, mm</td>
<td>11.2±1.0</td>
<td>16.6±0.5*</td>
<td>13.3±0.8*</td>
<td>12.4±0.4</td>
<td>18.5±1.1*</td>
<td>15.6±0.5†</td>
</tr>
<tr>
<td>LVLA, mm</td>
<td>10.1±0.2</td>
<td>11.7±0.2*</td>
<td>11.1±0.2*</td>
<td>10.5±0.2</td>
<td>13.3±0.5*</td>
<td>11.8±0.2†</td>
</tr>
<tr>
<td>MAP, mm Hg</td>
<td>124±4</td>
<td>100±4*</td>
<td>98±4*</td>
<td>124±3</td>
<td>109±4*</td>
<td>111±3*</td>
</tr>
<tr>
<td>HR, bpm</td>
<td>418±9</td>
<td>399±9</td>
<td>373±12*</td>
<td>409±7</td>
<td>398±8</td>
<td>376±8*</td>
</tr>
<tr>
<td>LVEDP, mm Hg</td>
<td>2.1±0.5</td>
<td>11.9±1.3*</td>
<td>8.3±1.0†*</td>
<td>2.1±0.4</td>
<td>12.5±1.5*</td>
<td>7.5±1.51*</td>
</tr>
<tr>
<td>CO, mL/min</td>
<td>102±4</td>
<td>83±5*</td>
<td>81±4*</td>
<td>98±4</td>
<td>70±3*</td>
<td>80±2†*</td>
</tr>
<tr>
<td>CI, mL/min/kg</td>
<td>324±15</td>
<td>286±15</td>
<td>303±17</td>
<td>230±9</td>
<td>171±10*</td>
<td>197±6†*</td>
</tr>
</tbody>
</table>

TPRI, mm Hg · min · kg/mL

Data represent mean±SEM. BW indicates body weight; HW, heart weight; LWV, left ventricular weight; LVIP, left ventricular internal perimeter; LVLA, left ventricular long axis diameter; MAP, mean arterial pressure; HR, heart rate; LVEDP, left ventricular end diastolic pressure; CO, cardiac output; CI, cardiac index; TPRI, total peripheral resistance index.

*P<0.05 vs Sham; †P<0.05 vs MI-Luc.

Kallikrein Gene Transfer Improves Cardiac Responses to Dobutamine-Induced Stress

Although there was no significant difference in dP/dt before dobutamine infusion among sham, MI-Luc, and MI-cHK groups, positive dP/dt in MI-Luc after dobutamine infusion was significantly lower than that of the sham group (8141±1144 versus 12 270±1306 mm Hg/s, n=5, P<0.05), whereas no significant difference was observed between MI-cHK (10 045±858 mm Hg/s, n=6) and sham groups (Figure 1A). The cardiac response to dobutamine in MI-Luc on positive dP/dt was reduced as compared with that of the sham group (+8.0±5.4 versus +36.6±6.5%, n=5, P<0.05), whereas kallikrein gene delivery improved the response to dobutamine-induced stress (+30.7±11.2%, n=6) (Figure 1B). Similarly, although there was no significant difference in CI before dobutamine infusion between sham, MI-Luc, and MI-cHK, CI in MI-Luc after dobutamine infusion was significantly lower than that of sham (249.6±21.6 versus 357.9±21.6 mL/min per kilogram, n=5, P<0.05), whereas no significant difference was observed between MI-cHK (328.5±31.0 mL/min per kilogram, n=6) and sham (Figure 1C). The cardiac response to dobutamine on CI in MI-Luc rats was reduced as compared with that of sham rats.
Kallikrein Gene Delivery Attenuates Cardiac Remodeling After MI

Figures 2 and 3 show the effect of kallikrein gene delivery on cardiac remodeling parameters in LV including collagen density, cardiomyocyte size, and capillary density. Histological and morphometric analyses showed that collagen density of rats receiving the luciferase gene (MI-Luc) markedly increased compared with the sham group (3.7±0.4 vs 1.9±0.1%, n=5 and 9, P<0.05), whereas kallikrein gene delivery (MI-cHK) significantly reduced collagen density (2.9±0.1%, n=6) (Figures 2A and 3A). Similarly, cardiomyocyte size in MI-Luc was remarkably increased at 6 weeks after MI compared with sham rats (2.9±0.1% vs 1.9±0.1%, n=5 and 9, P<0.05), and kallikrein gene delivery significantly reduced cardiomyocyte size (528±18.0 μm², n=6) (Figures 2B and 3B). Capillary density in MI-Luc was markedly decreased at 6 weeks compared with sham rats (1188±63 vs 1949±55/mm², n=5 and 9, P<0.05), and kallikrein gene delivery significantly increased capillary density (1589±77/mm², n=6) (Figures 2C and 3C).

Effect of Kallikrein Gene Delivery on Blood Flow, Vascular Resistance, and Nitrite/Nitrate Levels in LV

Left ventricular blood flow in MI-Luc was significantly decreased compared with sham rats (3.2±0.5 vs 5.9±1.0 mL/min/g, n=6 and 5, P<0.05), whereas kallikrein gene delivery significantly increased blood flow (4.8±0.4 mL/min/g, n=6) (Figure 4A). LV resistance increased in MI-Luc compared with the sham control (39.1±8.9 vs 5.9±21.9±3.7 mm Hg/min/g per milliliter, n=6 and 5, P<0.05), and kallikrein gene delivery significantly decreased this parameter (21.3±2.7 mm Hg/min/g per milliliter, n=6, P<0.05) (Figure 4B). Nitrite/nitrate levels in LV in MI-Luc were significantly lower than sham (157±10 vs 201±10 pmol/mg of protein, n=6 and 5, P<0.05), and kallikrein gene delivery increased nitrite/nitrate production in LV (198±75 pmol/mg of protein, n=6) (Figure 4C).

Apoptosis and Apoptosis-Related Protein Expression in LV

TUNEL-positive myocytes in the MI-Luc group at 2 weeks after MI were significantly increased compared with the sham group, whereas kallikrein gene delivery significantly reduced TUNEL-positive myocytes (n=6, P<0.05) (Figure 5A). There were positive correlations between TUNEL-positive myocytes and cardiomyocyte size (n=16, R=0.85, P<0.01) or LV internal perimeter (n=16, R=0.65, P<0.01) (Figures 5B and 5C). MI significantly reduced the phosphorylation of the survival factor Akt, whereas kallikrein gene delivery increased phospho-Akt but had no effect on total Akt (Figure 6A). MI resulted in increased cleaved caspase-3, whereas kallikrein gene delivery reduced caspase-3 activation (Figure 6B).

Discussion

In the present study, we showed that kallikrein gene delivery improves cardiac function and protects against cardiac remodeling in CHF. The beneficial effects of kallikrein gene transfer in the heart after MI include (1) attenuation of cardiac hypertrophy, fibrosis, and LV enlargement; (2) increase in capillary density; (3) improvement of cardiac responses to dobutamine infusion; (4) inhibition of myocardial apoptosis; and (5) enhancement of endothelial function including increased blood flow, decreased vascular resistance, and increased NO production. These combined results indicate that
the KKS prevents progression of heart failure and improves endothelial function through reduction of cardiac hypertrophy, fibrosis, and apoptosis.

In this study, we used dobutamine, a \(\beta_1\)-adrenergic agonist, to mimic exercise-induced stress and showed that hearts remodeled after MI exhibit a low response to dobutamine. Interestingly, kallikrein gene delivery improved cardiac responses against dobutamine infusion (Figure 1). These results indicate that hearts remodeled after MI have low cardiac reserve, and kallikrein gene delivery can improve cardiac reserve. Patients with heart failure must limit their exercises or activities because heart failure symptoms are induced by such stress conditions caused by low cardiac reserve. Therefore, cardiac responses against stress are more important than the basal level of cardiac function. Taken together, these findings indicate that KKS may play an important role in improving exercise capacity for patients with heart failure after MI.

Kallikrein gene transfer not only reduced cardiac hypertrophy and fibrosis but also increased capillary density after MI. Capillary density in the heart appeared to be especially important under stress conditions because the demand of oxygen and nutrition is increased in the heart during stress. A previous study by Liu et al.\(^{16}\) showed that MI induced the reduction of capillary density in non-MI area, and an ACE inhibitor prevented this reduction; however, kinin \(B_2\) receptor antagonist abolished the protective effect of ACE inhibition. These combined results indicate that kinin may promote an increase in capillary density. Increased Ang II levels in heart failure after MI may induce endothelial cell apoptosis and thus reduce capillary density in the heart.\(^{13}\) Our previous results together with the present study show that kallikrein gene delivery significantly increases cardiac kinin and NO levels.\(^{19}\) NO has been shown to inhibit human endothelial cell apoptosis.\(^{26}\) Therefore, kallikrein gene delivery may increase cardiac capillary density through suppression of endothelial cell apoptosis mediated by the kinin-NO pathway. Moreover, we have also reported that kallikrein gene delivery promotes spontaneous angiogenesis in hindlimb ischemia of rats.\(^{27}\) Thus, increased capillary density in hearts remodeled hearts after kallikrein gene transfer may contribute to enhanced endothelial function and reduced cardiac remodeling and apoptosis.

Myocardial apoptosis has been shown to be one of many important factors leading to cardiac remodeling and heart failure in several experimental and clinical heart failure stages.\(^{17,18}\) In this study, we observed positive correlations between TUNEL-positive myocytes and LV internal perim-
increase myocardial apoptosis in heart remodeled heart after phosphorylation of Akt and NO production also decreased in absence of repeated injections.

Therefore, the second set of data (week 6) emphasizes the long-term accumulated effects of kallikrein gene delivery. The results observed at week 2 after surgery represent the early effect of kallikrein gene delivery. The results observed at week 6 after surgery represent the acute and chronic effects of Ad-kallikrein injection as compared with control animals injected with the vehicle.

We evaluated the expression level of recombinant human tissue kallikrein in rats at only one point (3 days after gene delivery) to check whether the virus injection was successful. The time course of recombinant human tissue kallikrein levels in serum was not monitored continuously. However, we have shown that recombinant human tissue kallikrein in rat serum remains detectable on day 36 after tail vein injection of Ad-kallikrein. The current study was designed to analyze the acute and chronic effects of Ad-kallikrein gene delivery. Thus the time course selected was 2 and 6 weeks after coronary artery ligation or 1 and 5 weeks after gene delivery. Thus the time course selected was 2 and 6 weeks after coronary artery ligation or 1 and 5 weeks after gene delivery. The results observed at week 2 after surgery represent the early effect of kallikrein gene delivery. The results observed on week 6 after surgery represent the long-term accumulated effects of Ad-kallikrein gene delivery. Therefore, the second set of data (week 6) emphasizes the sustained effects of a single kallikrein gene delivery in the absence of repeated injections.

We showed that endothelial dysfunction exists and that phosphorylation of Akt and NO production also decreased in remodeled hearts. This may be one of the mechanisms to increase myocardial apoptosis in heart remodeled heart after MI because Akt and NO are well known cell survival factors, and endothelial dysfunction may enhance the decrease of Akt phosphorylation and NO production. Cardiac expression of the human tissue kallikrein mRNA in rats after kallikrein gene delivery by tail vein injection was previously identified by RT-PCR followed by Southern blot analysis. A continuous supply of exogenous kallikrein-kinin resulted in increased phosphorylation of Akt leading to increased Ca2+-independent activation of eNOS and thus NO formation. Phospho-Akt can inhibit apoptosis by phosphorylating the Bad component of the Bad/Bcl-X or Bad/Bcl-2 complex. Bad has been shown to be a proapoptotic member of the Bcl-2 family that binds to Bcl-2 and Bcl-XL, resulting in cell death through the release of cytochrome c from mitochondria and the activation of caspases. Increased kinin-NO production may also prevent the degradation of the pro-survival members of the bcl-2 family, the release of cytochrome c from mitochondria, and the inhibition of caspase activities. Taken together, these results indicate that kallikrein-kinin may inhibit myocardial apoptosis through the Akt-Bad or Akt-NO-cGMP pathways.

In conclusion, this study shows that KKS plays an important role in attenuating heart failure and improving cardiac reserves by the reduction of cardiac remodeling after MI through the enhancement of endothelial function and the inhibition of myocardial apoptosis through the Akt signaling pathway.

Acknowledgments

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