Therapy of Ischemic Cardiomyopathy With the Immunomodulating Agent Pentoxifylline

Results of a Randomized Study

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Background—Inflammatory immune activation commonly occurs in heart failure and may perpetuate this syndrome. We sought to determine whether the immunomodulating agent pentoxifylline enhances left ventricular function in patients with ischemic cardiomyopathy. We also investigated the effect of therapy on levels of brain natriuretic peptide, C-reactive protein, tumor necrosis factor-alpha, and the marker of apoptosis, Fas/Apo-1.

Methods and Results—In a single-center, prospective, randomized, double-blind, placebo-controlled study, 38 patients with ischemic cardiomyopathy received pentoxifylline 400 mg TID or placebo in addition to standard therapy. Clinical assessment, radionuclide ventriculography, echocardiography, and blood analyses were performed at baseline and after 6 months. There were no differences in baseline characteristics between the groups. Five patients died (4 in the placebo group). Pentoxifylline treatment resulted in an improvement in functional class (P<0.005) and an increase in systolic blood pressure (P<0.05) and left ventricular radionuclide ejection fraction (P<0.05) compared with the placebo-treated group. There were reductions in plasma concentrations of CRP, NT-pro BNP, TNF-alpha, and Fas/Apo-1 in the pentoxifylline compared with the placebo-treated group.

Conclusions—In patients with heart failure due to ischemic left ventricular dysfunction, the addition of pentoxifylline to standard therapy results in improvements in clinical status and radionuclide ejection fraction, which are accompanied by reductions in plasma markers of inflammation, prognosis, and apoptosis. (Circulation. 2004;109:750-755.)

Key Words: heart failure ■ cardiomyopathy ■ pentoxifylline ■ natriuretic peptides ■ C-reactive protein
Methods

Study Design and Patient Enrollment

The Ethics Committee of the University of the Witwatersrand approved the protocol (approval number M960718). All patients gave informed consent before study entry. This was a single-center, prospective, double-blind, randomized, placebo-controlled trial with 2 parallel arms. Inclusion criteria were as follows: (1) age ≥18 and ≤70 years; (2) stable New York Heart Association functional class II or III congestive heart failure secondary to coronary artery disease, as defined by the presence of ≥2-vessel disease on angiography; (3) left ventricular ejection fraction <40% by radionuclide scintigraphy; (4) sinus rhythm; and (5) eligible patients in whom high-quality echocardiographic images could be obtained. Exclusion criteria were (1) clinical conditions other than cardiomyopathy that could influence cytokine levels (i.e., rheumatoid arthritis, AIDS); (2) pregnancy; (3) severe exercise-induced malignant ventricular arrhythmia; (4) myocardial infarction within the last 12 months; (5) recent myocardial revascularization (<6 months); and (7) any clinical condition that according to the investigators precluded inclusion into the study.

All patients were undergoing optimal medical therapy for 3 months before randomization. Patients received diuretics, spironolactone, the ACE inhibitor perindopril, and the β-adrenoceptor blocker carvedilol. Target doses were 8 mg of perindopril once daily and 25 mg of carvedilol twice daily. After 3 months of stable medical therapy, patients whose left ventricular ejection fraction was <40% as assessed by radionuclide ventriculography were randomized to receive either pentoxifylline 400 mg TID (n = 20) or a matching placebo (n = 18) for 6 months. Pentoxifylline and an identical-looking placebo were purchased from Aventis Pharma. Patients were randomly assigned according to a computer-generated randomization list. At randomization, patients’ medical therapy consisted of ACE inhibitors (100%), β-blockers (100%), diuretics (93%), and spironolactone (50%). Medication remained unchanged throughout the study period. Monthly visits were scheduled for clinical assessment and evaluation of compliance. Clinical examination, exercise tests, echocardiographic and radionuclide studies, and determinations of plasma concentrations of TNF-α, Fas/Apo-1, high-sensitivity CRP, and NT-pro BNP were performed at baseline and then repeated 6 months after randomization. The primary end point was left ventricular ejection fraction assessed by the multiple gated equilibrium cardiac blood scintigraphic technique. To show a significant difference in the change in left ventricular ejection fraction at 6 months between pentoxifylline and placebo groups with 80% power, assuming this difference to be 15±12%, a sample number of 11 patients was required in each group.

Biochemistry

Plasma TNF-α and Fas/Apo-1 concentrations were determined as described previously.5,6 Fifteen milliliters of blood was withdrawn from an antecubital vein and collected into prechilled evacuated tubes that contained EDTA. Plasma was separated by centrifugation at 2500 rpm for 12 minutes within 15 minutes of collection. Aliquots were stored at −70°C. TNF-α measurements were performed with a commercially available high-sensitivity enzyme-linked immunosay (Amersham). The average of undiluted determinations performed in triplicate was calculated. Fas/Apo-1 was measured with a nonisotopic quantitative immunosay (Calbiochem) appropriately diluted. High-sensitivity CRP and NT-pro BNP plasma concentrations were determined by a commercially available ELISA (Roche Diagnostics).

Functional Class and Exercise Test

A physician who was blinded to the treatment assigned to each patient assessed the functional class of the patients during the baseline and follow-up visits. The same physician evaluated all patients. Exercise tests were performed with a modified Naughton protocol.12 The same cardiac technologist performed all tests.

Statistical Analysis

ANCOV techniques adjusting for baseline values were used to compare data at 6 months and changes from baseline to 6 months between the groups. Fisher’s exact test was used where appropriate. Data are presented as unadjusted values and expressed as mean±SD. Significance was assumed at a 2-tailed value P<0.05.

Results

Baseline characteristics of the 18 patients randomized to receive placebo were not different from the 20 patients randomized to receive pentoxifylline (Table 1). Twelve patients in the placebo group and 14 in the pentoxifylline group had CABG surgery (P=NS). None of the patients had the revascularization procedure in the previous 18 months. There were no differences in the dose of carvedilol either at baseline (20±14 mg daily in the placebo-treated group versus 21±18 mg daily in the pentoxifylline-treated group) or at 6 months between the 2 groups. For the duration of the study, all patients received perindopril 8 mg/d. During the 6-month study period, 5 patients died (4 in the placebo group; P=NS). These 5 patients were therefore not included in the follow-up analysis at 6 months. Except for ejection fraction (P<0.05) and peak ejection rate (P<0.05), there were no significant differences at baseline between the groups after exclusion of these 5 patients (Table 2).

| TABLE 1. Baseline Characteristics of Total Study Population |
|-----------------|-----------------|-----------------|
|                  | Placebo (n=18)  | Pentoxifylline (n=20) |
| Age, y           | 53±11           | 57±10            |
| Males, %         | 67              | 75               |
| Functional class I/II/III, n | 1/6/9/2          | 1/7/12/0         |
| Exercise time, min | 9.7±5.4         | 9.2±3.5          |
| Systolic BP, mm Hg | 117±18          | 119±19           |
| Diastolic BP, mm Hg | 74±13           | 77±14            |
| Heart rate, bpm  | 78±14           | 82±15            |
| Left ventricular EDD, mm | 61.5±8.5       | 60.7±6.8         |
| Ejection fraction (MUGA), % | 23.0±10.2       | 27.4±7.2         |
| E/A ratio        | 1.7±1.0         | 1.3±0.9          |
| Deceleration time, ms | 138±40          | 150±46           |
| TNF-α, pg/mL     | 7.7±3.5         | 7.0±3.3          |
| Fas/Apo-1, U/mL  | 31.6±17.0       | 26.4±19.8        |

BP indicates blood pressure; EDD, end-diastolic diameter; and MUGA, multiple gated equilibrium cardiac blood scintigraphic technique.
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Pentoxifylline treatment produced a decrease in both plasma pentoxifylline- and placebo-treated groups (Table 3). Plasma TNF-α and Fas/Apo-1 concentrations were higher in the pentoxifylline- and placebo-treated groups (Table 3). Pentoxifylline treatment produced a decrease in both plasma TNF-α and Fas/Apo-1 concentrations over the study period compared with the placebo-treated group (Table 3; Figure 2). Hence, plasma TNF-α and Fas/Apo-1 concentrations were lower in the pentoxifylline-treated group than in the placebo-treated group at 6 months (Table 3; Figure 2).

Blood Pressure, Functional Class, and Exercise Tolerance

Patients treated with pentoxifylline had a higher systolic blood pressure at 6 months than those treated with placebo (Table 2). However, diastolic blood pressure was unchanged. There was an improvement in functional class of heart failure in the pentoxifylline-treated group compared with the placebo-treated group (Table 2). Pentoxifylline did not influence exercise time (Table 2).

Left Ventricular Dimensions and Function

After adjustment for differences at baseline, pentoxifylline treatment produced an increase in radionuclide ejection fraction over the study period compared with the placebo-treated group (Table 2; Figure 1). The mean change in ejection fraction from baseline to 6 months was 9.2 ± 11.4 in the pentoxifylline group versus 1.1 ± 4.2 in the placebo group. Furthermore, at 6 months, radionuclide ejection fraction was greater in the pentoxifylline-treated than the placebo-treated group (Table 2; Figure 1). Ten patients (53%) in the pentoxifylline group improved their ejection fraction by >10 absolute units compared with no patients in the placebo group (Figure 1; P<0.005). Pentoxifylline did not produce a significantly greater increase in either peak ejection or peak filling rate compared with the placebo-treated group (Table 2). No other differences were noted after 6 months of therapy between the pentoxifylline and placebo-treated groups (Table 2).

TNF-α and Fas/Apo-1 Concentrations

Plasma TNF-α and Fas/Apo-1 concentrations were higher in the study group than in a group of 20 healthy volunteers (TNF-α, 7.33 ± 3.33 versus 1.44 ± 1.30 pg/mL, respectively, P<0.0001; Fas/Apo-1, 28.89 ± 18.43 versus 0.84 ± 0.20 U/mL, respectively, P<0.0001). At baseline, plasma concentrations of TNF-α and Fas/Apo-1 were similar in the pentoxifylline- and placebo-treated groups (Table 3). Pentoxifylline treatment produced a decrease in both plasma TNF-α and Fas/Apo-1 concentrations over the study period compared with the placebo-treated group (Table 3; Figure 2). Hence, plasma TNF-α and Fas/Apo-1 concentrations were lower in the pentoxifylline-treated group than in the placebo-treated group at 6 months (Table 3; Figure 2).

Plasma CRP and NT-pro BNP

High-sensitivity CRP concentrations for the total study group (8.75 ± 5.65 mg/L) were elevated compared with the upper range of normal values for healthy age-matched control

Figure 1. Effects of 6 months of pentoxifylline or placebo on ejection fraction. *P<0.05, placebo vs pentoxifylline, change from baseline to 6 months; ANCOVA to correct for baseline differences.
The rationale for using immunomodulating agents to treat patients with heart failure is based on the fact that excessive enhancement of proinflammatory cytokines appears to mimic many aspects of the heart failure phenotype. In addition, inflammatory cytokines have been shown to play a key role in the pathogenesis of atherosclerosis and coronary artery disease. Furthermore, increased levels of inflammatory cytokines have been observed in patients with silent myocardial ischemia and in patients with ischemic cardiomyopathy. Hence, it has been suggested that sustained TNF-α expression after myocardial infarction and in persistent ischemia may have detrimental effects on the remodeling process. However, recent trials with anticytokine agents such as etanercept and infliximab showed time- and dose-dependent worsening of heart failure. These rather discouraging results may be explained by the mechanisms of action of these agents. Infliximab exerts its effects by fixing complement in cells, which in the heart is reported to lead to cardiac myocyte lysis. Etanercept stabilizes TNF-α and hence leads to an accumulation of TNF-α in the peripheral circulation. In comparison, the effects of pentoxifylline are to reduce the synthesis of TNF-α by blocking transcriptional activation. Furthermore, pentoxifylline has been shown to inhibit apoptosis in different human cell types in vitro and in vivo. Hence, pentoxifylline is likely to be a more promising anticytokine agent. Indeed, we were able to show improvements in cardiac function with pentoxifylline, which were associated with reductions in inflammatory markers. The observed improvement in systolic function in the present study was comparable to our previous work performed in patients with idiopathic dilated cardiomyopathy.

Table 3. Plasma Inflammatory Markers, High-Sensitivity CRP, and pro-BNP at Baseline and at 6 Months for Patients Who Completed the Study Period

<table>
<thead>
<tr>
<th></th>
<th>Placebo (n=14)</th>
<th>Pentoxifylline (n=19)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Baseline</td>
<td>6 Months</td>
</tr>
<tr>
<td>TNF-α, pg/mL</td>
<td>7.9±3.9</td>
<td>9.3±6.2</td>
</tr>
<tr>
<td>Fas/Apo-1, U/mL</td>
<td>28.3±17.4</td>
<td>24.3±15.5</td>
</tr>
<tr>
<td>CRP, mg/L</td>
<td>6.9±5.7</td>
<td>9.9±7.0</td>
</tr>
<tr>
<td>NT-pro BNP, pmol/L</td>
<td>168±58</td>
<td>254±65</td>
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</tbody>
</table>

Abbreviations as in Table 1.

*Placebo vs pentoxifylline at 6 months, ANCOVA to correct for baseline differences.

†Placebo vs pentoxifylline change from baseline to 6 months, ANCOVA to correct for baseline differences.

Discussion

We have shown that pentoxifylline therapy as an adjunct to standard therapy results in improvements in systolic function in patients with ischemic cardiomyopathy. The increases in ejection fraction that were attributed to pentoxifylline therapy were associated with reductions in plasma concentrations of the inflammatory markers TNF-α and CRP, a marker of apoptosis (Fas/Apo-1), and a marker of ventricular dysfunction (NT-pro BNP).

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The beneficial effects of pentoxifylline observed in the present study are likely to have been mediated by several mechanisms. Patients treated with pentoxifylline showed a marginal decrease in plasma TNF-α concentrations but significant reductions in plasma Fas/Apo-1 concentrations. Because programmed cell death has been recognized as a contributing cause of myocyte loss in myocardial infarction, and TNF-α augments this process through the stimulation of apoptosis, the combination of reductions in TNF-α and Fas/Apo-1 concentrations in the present study may explain the clinical benefits observed with pentoxifylline therapy.

In addition to reductions in the inflammatory cytokine TNF-α, the acute-phase protein CRP was reduced in those patients who were treated with pentoxifylline. CRP has direct proinflammatory effects on endothelial cells, including the expression of adhesion molecules and monocyte chemotactic protein-1. Furthermore, CRP is implicated in the synthesis of TNF-α. Hence, a reduction in serum CRP concentrations could have beneficial effects on the progression of cardiac dysfunction.

We also observed reductions in NT-pro BNP in patients treated with pentoxifylline in the present study. Plasma levels of NT-pro BNP have been used in several clinical trials to assess the efficacy of medical therapy. The reduction in NT-pro BNP with pentoxifylline in the present study confirms the efficacy of this form of therapy in ischemic heart failure.

Although pentoxifylline did not abolish increments in circulating TNF-α concentrations in patients in the present study, experimental studies have suggested that physiological levels of TNF-α have cytoprotective effects on the heart during ischemic events. We therefore suggest that the use of pentoxifylline led to a moderate reduction of excessively elevated levels of TNF-α.

Although we are not aware of any large-scale study that has evaluated the safety of pentoxifylline in patients with heart failure, this pharmacological agent has been in clinical use for more than 25 years for conditions such as peripheral and cerebrovascular disease. Furthermore, because patients with peripheral vascular disease frequently also have coronary artery disease and heart failure, it is significant that large trials with more than 10,000 such patients have not reported increases in mortality in patients treated with pentoxifylline.

**Conclusions**

In patients with heart failure due to ischemic left ventricular dysfunction, the addition of pentoxifylline to standard therapy results in improvements in left ventricular ejection fraction, which are associated with reductions in markers of inflammation and markers of prognosis.

**Acknowledgments**

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**References**


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