Effects of Hawthorn on the Progression of Heart Failure in a Rat Model of Aortic Constriction

Hyun Seok Hwang, M.S., Marvin O. Boluyt, Ph.D., Kimber Converso, B.S., Mark W. Russell, M.D., and Barry E. Bleske, Pharm.D., FCCP

Study Objective. To determine the effects of hawthorn (Crataegus oxyacantha) on left ventricular remodeling and function in pressure overload–induced heart failure in an animal model.

Design. Randomized, parallel, dose-ranging animal study.

Setting. University research facility.

Animals. Seventy-four male Sprague-Dawley rats; 44 were included in the final analysis.

Intervention. Rats underwent a sham operation or aortic constriction. Rats subjected to the sham operation were treated with vehicle (10% agar-agar), and those subjected to aortic constriction were treated with vehicle or hawthorn (C. oxyacantha special extract WS 1442) 1.3, 13, or 130 mg/kg for 5 months.

Measurements and Main Results. Rats and their hearts were weighed, and echocardiographic measurements were performed at baseline and at 2, 3, 4, and 5 months after aortic constriction. Protein expression for markers of fibrosis and for atrial natriuretic factor was also measured. Aortic constriction increased the left ventricular:body weight ratio by 53% in vehicle-treated rats; Hawthorn treatment did not significantly affect the aortic constriction–induced increase in this ratio. Left ventricular volumes and dimensions at systole and diastole significantly increased 5 months after aortic constriction compared with baseline in rats given vehicle (> 20% increase, p<0.05) but not in those given hawthorn 130 mg/kg (< 10% increase). After aortic constriction, the velocity of circumferential shortening significantly decreased in the vehicle group but not in the medium- or high-dose groups. In the aortic constriction–vehicle group, the induced increases in messenger RNA expression for atrial natriuretic factor (~1000%) and fibronectin (~80%) were significantly attenuated by high-dose hawthorn treatment by approximately 80% and 50%, respectively.

Conclusion. Hawthorn treatment exhibited modest beneficial effects on cardiac remodeling and function during long-term, pressure overload–induced heart failure in rats.

Key Words: hawthorn, Crataegus oxyacantha, heart failure, atrial natriuretic factor, ANF; left ventricular volume, fibronectin, animal study.

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Heart failure is a clinical syndrome with diverse causes and has progressive cardiac dysfunction as its distinguishing characteristic.1 All patients with systolic heart failure have left ventricular dysfunction, but cardiac dysfunction itself does not always lead to symptoms of the disease. Left ventricular remodeling plays an important role in the progression of chronic heart failure. Early remodeling is an adaptive response to the increased load on the heart and/or the loss
of contractile components to maintain pumping capacity. However, prolonged overload taxes the limits of adaptive mechanisms, forcing compromises that result in progressive deterioration of pump function. Ultimately, the left ventricle enlarges, and its diastolic and/or systolic function is impaired.

Aortic constriction in rats causes chronic left ventricular pressure overload, progressive cardiac remodeling, and subsequent cardiac failure that mimics heart failure in humans.\(^2\)\(^{-}\)\(^4\) Pressure overload also induces an increase in cardiac fibrosis, resulting in diastolic dysfunction, again similar to findings in humans.\(^5\)\(^\text{,}^6\) In addition, this rat model leads to initial compensatory hypertrophy that progresses to heart failure in a predictable manner akin to the progression seen in people. Finally, effects of angiotensin-converting enzyme (ACE) inhibitors (and other drugs) in this model are similar to those observed in human heart failure.\(^4\) Overall, chronic aortic constriction–induced pressure overload is a good model to study the effects of drugs on the development and progression of heart failure.

At present, an estimated 38% of the adult population of the United States uses herbal therapy.\(^7\) Hawthorn (\textit{Crataegus oxycantha}) is a popular botanical treatment for heart disease, including heart failure, especially in Europe and China. Early trials demonstrated that hawthorn may offer modest beneficial effects in the treatment of heart failure.\(^8\)\(^,^9\) Recent trials, however, have not shown an especially striking benefit for heart failure that is well treated with standard drug therapy.\(^10\)\(^,^11\)

We recently demonstrated that short-term hawthorn treatment improved cardiac functions in pressure overload–induced cardiac remodeling.\(^12\) However, whether long-term treatment with hawthorn influences the progression of heart failure—specifically cardiac remodeling and fibrosis—is unknown. Therefore, the purpose of this study was to determine the effects of hawthorn on the progression of heart failure in a rat model of chronic pressure overload.

**Methods**

**Animals**

Male Sprague-Dawley rats, each weighing 250–275 g, were obtained from Charles River Laboratories (Wilmington, MA). The rats were housed two or three to a cage, fed a standard laboratory diet (5001; PMI Nutrition International, Richmond, IN), given water ad libitum, and maintained on a cycle of 12 hours light and 12 hours dark. The University Committee on Use and Care of Animals at the University of Michigan approved all animal protocols, which conformed to the \textit{Guiding Principles in the Care and Use of Animals} of the American Physiological Society.

**Experimental Design**

The rats were subjected to a sham operation or aortic constriction. Rats subjected to the sham operation were treated with vehicle (10% agar-agar), and those subjected to aortic constriction were treated with vehicle or three doses of hawthorn (\textit{C. oxycantha} special extract WS1442; Dr. Willmar Schwabe Pharmaceuticals GmbH & Co. KG, Karlsruhe, Germany) for 5 months beginning 1 week after aortic constriction. \textit{Crataegus oxycantha} special extract WS 1442 is a dry extract derived from hawthorn leaves with flowers, with a raw material:extract ratio of 4–6.6:1 using 45% (weight/weight) ethanol as extraction solvent. The extract is adjusted to contain 17.3–20.1% oligomeric procyanidins. The batch we used (Ch. 289 N001) contained 18.8% oligomeric procyanidins, 2.9% flavonoids (calculated as hyperoside), 0.4% oleanolic acid, and 1% cyanidin (calculated as cyanidin chloride).

The doses and administration schedule for hawthorn were similar to those previously described.\(^12\) In brief, the recommended dosage of hawthorn in humans is 900 mg/day. This is approximately 13 mg/kg/day for a 70-kg person. As a result, we selected a human dose of 13 mg/kg as our middle dose and used doses 10-fold higher and 10-fold lower to achieve a reasonable dose-response range. Therefore, doses were 1.3
EFFECT OF HAWTHORN ON HEART FAILURE IN A RAT MODEL  Bleske et al

Hawthorn was administered by oral gavage at approximately 5:00 PM.

After 1 week of aortic constriction, the initial treatment groups consisted of the following: sham operation–vehicle group (12 rats), aortic constriction–vehicle group (15), aortic constriction–low-dose group (17), aortic constriction–medium-dose group (15), and aortic constriction–high-dose group (15). We terminated the study when the mortality rate reached approximately 50% in any of the treated aortic-constriction groups. Therefore, the study ended at 5 months. The decision to terminate the trial at a 50% mortality rate was made to ensure that we had adequate power at the end of the study and that a transition to decompensated heart failure could occur. Before the study ended, we conducted a final echocardiographic assessment. Afterward, we euthanized the rats and excised and dissected their hearts to collect the right ventricle, septum, left ventricular free wall, left atrium, and right atrium. Tissues were stored at -80°C for subsequent isolation of RNA and protein.

Aortic Constriction

Aortic constriction was conducted as described previously. In brief, we performed left thoracotomy with the animal under anesthesia induced with pentobarbital 50 mg/kg and exposed the ascending aorta. A Weck hemoclip was placed around the ascending aorta 4–6 mm superior to the aortic valve, with the gap of the Weck hemoclip applicator set at 1.02 mm.

Outcome Measures

We assessed body and ventricular weights, mortality rates, relative wall thickness, and cardiac function (left ventricular dimension in systole and diastole, stroke volume, velocity of circumferential shortening, and fractional shortening). In addition, we examined the expression of messenger RNA (mRNA) for atrial natriuretic factor (ANF), β-myosin heavy chain, and fibronectin, as well as the abundance of collagen type III.

Echocardiography

We echocardiographically measured cardiac function at baseline (1 wk before aortic constriction) and at 2, 3, 4, and 5 months after aortic constriction. Two-dimensional, guided, M-mode recordings were obtained from the short-axis view at the level of the papillary muscles by using a system (Acuson Sequoia; Siemens Medical Solutions, Malvern, PA) equipped with a 15-MHz linear-array transducer or a system (Vivid 7; GE Medical Systems, Milwaukee, WI) equipped with a 10-MHz phased-array transducer (S10-MHz; GE Medical Systems).

For each M-mode measurement, at least three consecutive cardiac cycles were sampled, as previously described. A single echocardiographer performed all measurements in accordance with the conventions of the American Society of Echocardiography.

Relative wall thickness was defined as 2 times the thickness of the posterior wall in diastole divided by the dimension of the left ventricle during diastole. Relative wall thickness was used instead of absolute wall thickness to facilitate comparisons among rats because of inherent variances from rat to rat.

RNA Blotting

Blotting for RNA was performed as described previously, with modifications. We isolated RNA from the left ventricle and size-fractionated 10 µg of total RNA by electrophoresis through 1% agarose gels. Samples were transferred electrophoretically at 5 V/cm to a nylon membrane (Nytran SPC; Whatman Inc., Piscataway, NJ) and hybridized with 32P-radiolabeled probes overnight at 68°C for complementary DNA probes and at 42–45°C for oligonucleotide probes by using a hybridization buffer (PerfectHyb Plus; Sigma-Aldrich Corp., St. Louis, MO).

Hybridization intensity was quantified with a Personal Phosphorimager FX (Bio-Rad Laboratories, Hercules, CA). Signals visualized on computer screen were identified by their position relative to migration of 18S and 28S ribosomal RNA. They were delineated by rectangles and quantified after subtracting the background. Each blot was subsequently stripped and reprobed. The signal from each sample was normalized to the signal obtained with an oligonucleotide specific for the 3′ untranslated region of glyceraldehyde-3′-phosphate dehydrogenase. The probe for β-myosin heavy chain was an oligonucleotide described previously. Complementary DNA probes for ANF and fibronectin were synthesized from a template by the random prime method (Promega Corp., Madison, WI).
Immunoblotting for Collagen Type III

Portions of the frozen left ventricle were processed for immunoblotting, as described elsewhere. Relative levels of collagen type III were measured protein using a monoclonal antibody directed at collagen type III (Santa Cruz Biotechnology, Santa Cruz, CA). Signals were quantified by using enhanced chemiluminescence.

Statistical Analysis

Data are expressed as percentages or mean ± standard error of the mean. Mortality rates were determined by Kaplan-Meier analysis. End-point variables were analyzed by using a 1-way analysis of variance (ANOVA), and group differences were determined with the least significant difference post hoc procedure. Contrasts were performed to compare the aortic constriction–vehicle group with the three hawthorn dose groups combined to determine the overall effect of hawthorn.

Echocardiographic data were analyzed by using 2-way ANOVA (group x time) with repeated measures, and group differences were determined with the least significant difference post hoc procedure. When the interaction for group x time was significant, a 1-way ANOVA procedure with repeated measures was conducted for each group separately, followed by the least significant difference post hoc procedure to determine differences between time points within a group. Differences were considered statistically significant when α was 0.05 or less.

Results


Table 1. Postmortem Analysis During 5 Months of Pressure Overload by Treatment Group

<table>
<thead>
<tr>
<th>Variable</th>
<th>Sham Operation, Vehicle Group (n=12)</th>
<th>Aortic Constriction</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Vehicle Group (n=9)</td>
<td>Low-Dose Group (n=10)</td>
</tr>
<tr>
<td>Mortality rate</td>
<td>0%</td>
<td>40% (6/15)</td>
</tr>
<tr>
<td>Mean ± SEM Initial body weight (g)</td>
<td>262 ± 6.6</td>
<td>261 ± 8.5</td>
</tr>
<tr>
<td>Final body weight (g)</td>
<td>561 ± 22</td>
<td>555 ± 22</td>
</tr>
<tr>
<td>LV dry:LV wet ratio (mg:mg)</td>
<td>0.23 ± 0.003</td>
<td>0.23 ± 0.002</td>
</tr>
<tr>
<td>RV:body weight ratio (mg:g)</td>
<td>0.51 ± 0.02</td>
<td>0.53 ± 0.02</td>
</tr>
</tbody>
</table>

<sup>a</sup>SEM = standard error of the mean; LV = left ventricular; RV = right ventricular.
<sup>b</sup>End-point variables were evaluated by using 1-way analysis of variance followed by the least significant difference post hoc procedure. Mortality was determined by Kaplan-Meier analysis.
<sup>c</sup>For all aortic-constriction groups vs sham operation–vehicle group.
<sup>d</sup>For all comparisons over the course of the experiment.

Figure 1. Cardiac hypertrophy after 5 months of pressure overload assessed as left ventricular (LV) mass (panel A) and the LV:body weight ratio (panel B) at the end point of the experiment. Data are mean ± standard error of the mean. AC = aortic constriction. *p<0.05 vs sham operation–vehicle group (1-way analysis of variance followed by the least significant difference post hoc procedure).
constriction–medium-dose group (6), and aortic constriction–high-dose group (7). Increases in body weight from before the operation to 5 months after the operation were similar in the sham operation–vehicle group and aortic constriction group–vehicle group. This result indicated that neither hawthorn treatment nor aortic constriction influenced the increase in body weight that occurred over the course of the experiment (Table 1).

After 5 months, left ventricular mass was 47% greater in the aortic-constriction group than in the sham operation–vehicle group (p<0.05; Figure 1A). Hawthorn treatment significantly attenuated the increase in left ventricular mass caused by aortic constriction. Aortic constriction increased the left ventricular:body weight ratio by 53% in vehicle-treated rats (p<0.001). The influence of hawthorn on the aortic constriction–induced increase in this ratio was not statistically significant, although the trend was similar to changes observed for left ventricular mass alone (Figure 1B).

During 5 months of aortic constriction–induced pressure overload, the mortality rate was 40% for vehicle-treated rats. No significant differences

<table>
<thead>
<tr>
<th>Measure</th>
<th>Sham Operation, Vehicle Group (n=12)</th>
<th>Vehicle Group (n=9)</th>
<th>Hawthorn Low-Dose Group (n=10)</th>
<th>Hawthorn Medium-Dose Group (n=6)</th>
<th>Hawthorn High-Dose Group (n=7)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stroke volume (ml/beat)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Baseline</td>
<td>205 ± 19.4</td>
<td>197 ± 19.9</td>
<td>190 ± 17.7</td>
<td>192 ± 16.7</td>
<td>228 ± 26.0</td>
</tr>
<tr>
<td>Month 2</td>
<td>399 ± 25.4b</td>
<td>297 ± 40.2b</td>
<td>303 ± 34.2b</td>
<td>245 ± 33.5</td>
<td>282 ± 40.8</td>
</tr>
<tr>
<td>Month 3</td>
<td>371 ± 19.7b</td>
<td>388 ± 49.7b</td>
<td>382 ± 20.6b</td>
<td>303 ± 25.3b</td>
<td>306 ± 40.0</td>
</tr>
<tr>
<td>Month 4</td>
<td>429 ± 27.7b</td>
<td>296 ± 35.4bc</td>
<td>361 ± 34.5b</td>
<td>289 ± 34.7bc</td>
<td>335 ± 31.7b</td>
</tr>
<tr>
<td>Month 5</td>
<td>418 ± 24.0b</td>
<td>319 ± 17.4bc</td>
<td>413 ± 36.1b</td>
<td>344 ± 32.2b</td>
<td>319 ± 32.5b</td>
</tr>
<tr>
<td>Percent change</td>
<td>+103</td>
<td>+62</td>
<td>+117</td>
<td>+79</td>
<td>+40</td>
</tr>
<tr>
<td>LV dimension, diastole (mm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Baseline</td>
<td>6.29 ± 0.23</td>
<td>6.20 ± 0.20</td>
<td>6.05 ± 0.21</td>
<td>6.25 ± 0.18</td>
<td>6.59 ± 0.21</td>
</tr>
<tr>
<td>Month 2</td>
<td>7.78 ± 0.21b</td>
<td>7.30 ± 0.34b</td>
<td>7.25 ± 0.38b</td>
<td>6.70 ± 0.37</td>
<td>6.99 ± 0.31</td>
</tr>
<tr>
<td>Month 3</td>
<td>7.81 ± 0.16b</td>
<td>7.84 ± 0.34b</td>
<td>7.87 ± 0.21b</td>
<td>7.07 ± 0.20b</td>
<td>7.20 ± 0.45</td>
</tr>
<tr>
<td>Month 4</td>
<td>8.15 ± 0.20b</td>
<td>7.30 ± 0.33b</td>
<td>7.81 ± 0.32b</td>
<td>7.05 ± 0.28b</td>
<td>7.44 ± 0.28</td>
</tr>
<tr>
<td>Month 5</td>
<td>8.26 ± 0.19b</td>
<td>7.58 ± 0.20b</td>
<td>8.20 ± 0.28b</td>
<td>7.72 ± 0.30b</td>
<td>7.20 ± 0.38</td>
</tr>
<tr>
<td>Percent change</td>
<td>+31</td>
<td>+22</td>
<td>+37</td>
<td>+24</td>
<td>+9</td>
</tr>
<tr>
<td>LV dimension, systole (mm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>3.71 ± 0.17</td>
<td>3.60 ± 0.13</td>
<td>3.25 ± 0.24</td>
<td>3.77 ± 0.17</td>
<td>3.94 ± 0.17</td>
</tr>
<tr>
<td>Month 2</td>
<td>4.88 ± 0.21b</td>
<td>4.39 ± 0.37b</td>
<td>4.33 ± 0.44b</td>
<td>3.83 ± 0.45</td>
<td>4.06 ± 0.26</td>
</tr>
<tr>
<td>Month 3</td>
<td>4.73 ± 0.19b</td>
<td>4.73 ± 0.34b</td>
<td>4.59 ± 0.36b</td>
<td>3.72 ± 0.23</td>
<td>4.07 ± 0.57</td>
</tr>
<tr>
<td>Month 4</td>
<td>4.85 ± 0.24b</td>
<td>4.54 ± 0.41b</td>
<td>4.78 ± 0.45b</td>
<td>3.95 ± 0.36</td>
<td>4.31 ± 0.30</td>
</tr>
<tr>
<td>Month 5</td>
<td>5.28 ± 0.21b</td>
<td>4.84 ± 0.29b</td>
<td>5.13 ± 0.37b</td>
<td>4.65 ± 0.58</td>
<td>4.07 ± 0.28</td>
</tr>
<tr>
<td>Percent change</td>
<td>+42</td>
<td>+34</td>
<td>+58</td>
<td>+23</td>
<td>+3</td>
</tr>
<tr>
<td>Fractional shortening (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>40.8 ± 1.7</td>
<td>41.8 ± 0.3</td>
<td>47.1 ± 2.5</td>
<td>39.7 ± 2.3</td>
<td>40.0 ± 2.3</td>
</tr>
<tr>
<td>Month 2</td>
<td>37.9 ± 1.5</td>
<td>37.5 ± 3.3</td>
<td>41.2 ± 3.1</td>
<td>43.3 ± 4.1</td>
<td>41.9 ± 3.3</td>
</tr>
<tr>
<td>Month 3</td>
<td>39.6 ± 1.6</td>
<td>40.1 ± 2.9</td>
<td>42.1 ± 3.3</td>
<td>47.5 ± 2.6</td>
<td>44.6 ± 4.8</td>
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<tr>
<td>Month 4</td>
<td>40.7 ± 2.0</td>
<td>38.5 ± 3.7</td>
<td>39.6 ± 3.9</td>
<td>44.0 ± 4.3</td>
<td>42.3 ± 2.1</td>
</tr>
<tr>
<td>Month 5</td>
<td>36.3 ± 1.4</td>
<td>36.7 ± 2.4</td>
<td>37.7 ± 2.8</td>
<td>40.3 ± 5.4</td>
<td>43.3 ± 2.6</td>
</tr>
<tr>
<td>Percent change</td>
<td>-9</td>
<td>-12</td>
<td>-20</td>
<td>+2</td>
<td>+8</td>
</tr>
</tbody>
</table>

Data are mean ± standard error of the mean. Baseline = 1 wk before aortic constriction.

aFor all measures, 2-way analysis of variance with a repeated-measures, least significant difference post hoc procedure yielded the following p values: group p≥0.05, time p<0.01, and group x time p≥0.05.
bp<0.05 compared with baseline value within same treatment group.
cp<0.05 compared with sham operation–vehicle value at same time point.
dCalculated as (value at 5 months – baseline value)/baseline value.
were seen when these rats were compared with hawthorn-treated rats (Table 1). Aortic constriction significantly increased relative wall thickness by 40% in vehicle-treated rats. Over time, low- and medium-dose hawthorn treatment significantly diminished the increase in constriction-induced relative wall thickness (Figure 2).

Left ventricular dimension in systole and diastole significantly increased by greater than 20% over baseline in the aortic constriction–vehicle group; however, no significant increase (< 10%) was observed in the aortic constriction–high-dose group (Table 2). Likewise, left ventricular volumes at end of systole and end of diastole did not significantly increase at 5 months after aortic constriction in the aortic constriction–high-dose group. Significant increases were observed in the aortic constriction–vehicle group (Figure 3A and 3B). In aggregate, the data indicated that high-dose hawthorn modified aortic constriction–induced cardiac remodeling most significantly.

Aortic constriction attenuated the increase in stroke volume that occurred over time in vehicle-treated rats (Table 2). Constriction-induced attenuation of the increase in stroke volume over time was counteracted, especially by low and medium doses of hawthorn. Velocity of circumferential shortening significantly decreased in the aortic constriction groupstreated with vehicle or low-dose hawthorn. However, medium and high doses did not significantly lower velocity of circumferential shortening after 5 months of aortic constriction (Figure 3C).

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**Figure 2.** Cardiac remodeling, assessed as relative wall thickness, during 5 months of pressure overload. Data are mean ± standard error of the mean. Data were analyzed by using a 2-way analysis of variance with repeated measures followed by the least significant difference post hoc procedure. AC = aortic constriction; baseline = 1 week before AC. ap<0.05 vs sham operation–vehicle group at 5 months; bp<0.05 vs aortic constriction–vehicle group at 5 months.

**Figure 3.** Cardiac function during 5 months of pressure overload evaluated as left ventricular volume (LV) at end of systole (panel A), left ventricular volume at end of diastole (panel B), and velocity of circumferential shortening in the left ventricle (panel C). Data are mean ± standard error of the mean. Data were analyzed by using a 2-way analysis of variance with repeated measures followed by the least significant difference post hoc procedure. AC = aortic constriction; baseline = 1 week before AC. ap<0.05 vs baseline within the same treatment group.
For the sham operation–vehicle, aortic constriction–vehicle, and aortic constriction–low-dose groups, fractional shortening was lower at 5 months compared with baseline (Table 2). In contrast, fractional shortening seemed to improve in the medium- and high-dose groups. Overall, the data suggested that hawthorn given at a high dose maintained cardiac function better than vehicle after 5 months of aortic constriction.

At 5 months, levels of ANF mRNA significantly rose by 1000% in the hearts of rats who received aortic constriction and vehicle (Figure 4A). Each dose of hawthorn almost completely counteracted the increase in ANF mRNA caused by long-term aortic constriction. At 5 months after sham operation or aortic constriction, levels of mRNA for β-myosin heavy chain did not significantly differ among any of the five groups (Figure 4B). After pressure overload, expression of fibronectin mRNA in the left ventricle significantly increased by 78% (Figure 4C). Each of the three doses of hawthorn markedly attenuated the aortic constriction–induced increase in mRNA expression for fibronectin.

The abundance of collagen type III protein was significantly and similarly elevated in the three aortic constriction groups treated with vehicle or medium or high doses of hawthorn compared with the sham operation–vehicle group (Figure 5). Collagen type III protein levels in the aortic constriction–low-dose group were not significantly greater than those of the sham-vehicle group, but they were also not significantly lower than the aortic constriction–vehicle levels. Taken together, these data showed that hawthorn treatment had little, if any, effect on the aortic constriction–induced increase in the abundance of collagen III protein.

Discussion

Our survival data indicate that our model of pressure overload likely caused a progression of heart failure, which increased mortality rates in rats undergoing aortic constriction compared with those receiving sham operation plus vehicle. The most potent effects of hawthorn were observed for variables that reflected the nature of cardiac remodeling. These included marked suppression of the aortic constriction–induced increase in the abundance of collagen III protein.
increase in mRNA expression for ANF and fibronectin, as well as alterations in structural remodeling. All doses of hawthorn positively influenced ANF and fibronectin mRNA expression. At the highest dose, hawthorn favorably affected ventricular size and systolic function. The results suggest that hawthorn exerted important effects on remodeling and on the progression of heart failure in a pressure overload model of heart failure.

To our knowledge, this was the first study to show that hawthorn treatment influences cardiac remodeling in the context of long-term pressure overload. Hawthorn attenuated the increase in left ventricular mass after aortic constriction. Although the left ventricular:body weight ratio was not significantly attenuated, the trend was similar to that observed with left ventricular mass. Given other findings for remodeling parameters, although the trend is not statistically significant, it is likely to be clinically significant. The mean percentage increase in left ventricular systolic and diastolic dimensions was less than 10% (p>0.05 vs baseline) in the high-dose hawthorn group after 5 months of aortic constriction compared with an increase of greater than 20% (p<0.05 vs baseline) in the aortic constriction–vehicle group. We noted similar findings regarding left ventricular systolic and diastolic volumes. In support of these results, ANF expression, which increases in response to pressure and stretch, was significantly lower in the hawthorn groups than in the aortic constriction–vehicle group.

Possible mechanisms for the benefits of hawthorn on cardiac remodeling are likely multifactorial given its wide range of reported effects on the myocardium and on signaling pathways. These include inotropic actions on cardiac muscle that may indirectly benefit remodeling of the heart, as well as a vasodilatory property of hawthorn that may be attributed to nitric oxide. Another factor to consider is that hawthorn exhibited protective effects against ischemia-reperfusion injury by conferring mitochondrial protection. If true, prevention of mitochondrial damage and apoptosis would be consistent with improved remodeling because it would prevent loss of myocytes that can lead to weakening and fibrosis.

Additional mechanisms—or mechanisms that may explain the observed effects of hawthorn—may also be proposed on the basis of observed effects of the agent or its individual constituents (flavonoids and polyphenolic compounds). These additional mechanisms may include effects on reactive oxygen species, nitric oxide (endothelial and inducible nitric oxide synthases), and specific signaling pathways involving p38 mitogen-activated protein kinase, nuclear factor-κ B, and extracellular signal-regulated kinases 1 and 2. The intrinsic ability of the ventricles to contract was maintained most with the medium and high doses of hawthorn after aortic constriction, as evidenced by the lack of change in the velocity of circumferential shortening and fractional shortening. By comparison, these parameters decreased in the vehicle and low-dose groups after aortic constriction. These findings were consistent with our observations regarding ventricular remodeling in this study. Together, the data suggest that hawthorn may attenuate the transition from compensated to decompensated heart failure after long-term pressure overload. The apparent effects of hawthorn to lessen a maladaptive form of hypertrophy that we found were consistent with our previous results from a short-term model of pressure overload. In that model, hawthorn maintained a more adaptive form of hypertrophy after aortic constriction than vehicle, on the basis of the velocity of circumferential shortening and relative wall thickness.

Potential mechanisms to account for the effects on systolic function include beneficial effects on apoptosis, reactive oxygen species, endothelial nitric oxide synthase, and inducible nitric oxide synthase, along with inotropic actions of hawthorn. Previous studies have shown that hawthorn increases the inotropic activity of cardiac muscle. Moreover, treatment of failing human myocardium with hawthorn increases both the calcium ion transient and the force production of papillary muscles from these hearts. The mechanism of these inotropic effects is partly attributed to the ability of hawthorn to inhibit Na⁺-K⁺-adenosine 5′-triphosphatase.

Many studies have shown that ANF mRNA levels increase in response to a wide range of hypertrophic stimuli, including aging, long-term β-adrenergic stimulation, aortic constriction, thyroid hormone, hypertension, exercise training, and coronary artery occlusion. Therefore, repeat induction of gene expression for ANF in the left ventricle is one of the most reliable markers of cardiac hypertrophy. Levels of ANF mRNA were significantly elevated in spontaneously hypertensive rats compared with normotensive Wistar-Kyoto rats and increased even further when hypertension-induced hypertrophy progressed to heart failure. In the
present study, hawthorn almost completely abolished the increase in ANF mRNA levels caused by 5 months of pressure overload. These data helped confirm our findings that hawthorn significantly affects ventricular remodeling.

Excessive accumulation of proteins of the extracellular matrix, fibronectin, and collagen causes myocardial fibrosis that results in stiffening of the heart and diastolic dysfunction. Type III collagen is one of the two major types of collagen expressed in the heart. Type III collagen accounts for approximately 20% of the total collagen in rat heart tissue. Levels of mRNA for collagen type III are transiently elevated in the heart after aortic constriction in chronic hypertension and during the transition from compensated hypertrophy to heart failure. Fibronectin is a large protein of the extracellular matrix that binds together various components and cells of the matrix, contributing to its overall integrity. Fibronectin gene expression increases in response to advancing age, aortic constriction, chronic hypertension, cardiomyopathy, and heart failure.

In our study, 5 months of aortic constriction increased levels of type III collagen by 35% and increased fibronectin mRNA levels by 78%, consistent with previous findings. Hawthorn treatment did not significantly attenuate the aortic constriction–induced increase in type III collagen protein. Hawthorn did, however, attenuate the constriction-induced increase in fibronectin mRNA expression, suggesting that it may have modified the nature of extracellular matrix to some extent. The lack of effect on the abundance of collagen type III protein suggests that hawthorn did not mitigate the aortic constriction–induced increase in the total amount of extracellular matrix. Effects of hawthorn on fibronectin expression may contribute, at least in part, to modest changes in end-diastolic dimension and volume, as we observed in some of the hawthorn-treated groups.

One obvious limitation was that this investigation was an animal study and that mechanisms for the results found were not well defined. However, ANF was suppressed in the hawthorn-treated groups, and this finding appears to confirm our results regarding structural remodeling. Another limitation was that this was not a mortality trial; therefore, inferences from our mortality results are limited. Although mortality rates seemed to be higher in the hawthorn groups than in the others, we found no clear dose response, and these findings were likely the result of chance. Given the few animals studied, one death can dramatically alter interpretation of the data.

Also of note, echocardiography was performed at five time points, and data for rats that died before the 5-month end point were excluded from the final analysis. When we examined the last echocardiographic measurements obtained from rats that died before this point, we noted no obvious indications of heart failure (e.g., marked reduction in fractional shortening or left ventricular ejection fraction). Autopsies were not conducted for rats that died before the end-point analysis. Therefore, although heart failure due to pressure overload was the assumed cause of death, no autopsy or echocardiographic data were available to confirm this assumption.

From a clinical perspective, a potential limitation of this study was the fact that hawthorn was not administered with standard-of-care therapies such as ACE inhibitors and β-blockers. On the other hand, this design might actually have been the strength of the study in that hawthorn could demonstrate its effects on the progression of heart failure. One can easily reason that powerful remodeling drugs such as ACE inhibitors and β-blockers may mask or attenuate the effects of hawthorn on remodeling. In clinical terms, this may have important therapeutic implications, particularly in patients who cannot tolerate or who are not candidates for standard-of-care ACE inhibition or β-blockade. Because hawthorn has shown remodeling effects similar to those of both these treatments, high-risk patients who cannot receive this standard of care are most likely to benefit from hawthorn therapy, especially with regard to ventricular remodeling. Perhaps an important niche for complementary and alternative medicine may be the treatment of patients who cannot tolerate or who are not candidates for standard approaches.

Conclusion

Our data suggest that hawthorn attenuates maladaptive ventricular remodeling and decreases in systolic function associated with longstanding pressure overload. Of note, these findings are consistent with the observation that hawthorn attenuates increases in ANF, a marker related to cardiac hypertrophy, and in fibronectin, a marker associated with fibrosis. Our study demonstrated the importance of herbal therapy in a rat model of heart failure. Further studies in humans are now warranted to determine the effect of hawthorn on ventricular remodeling.
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