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Cardiac Adaptations to High-Intensity Aerobic Training in Premenopausal and Recent Postmenopausal Women: The Copenhagen Women Study

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Background—We examined the role of menopause on cardiac dimensions and function and assessed the efficacy of exercise training before and after menopause.

Methods and Results—Two groups of healthy premenopausal (n=36, 49.4±0.3 years) and postmenopausal (n=37, 53.5±0.5 years) women with no history of cardiovascular disease and with a mean age difference between groups of only 4 years were studied. Cardiac dimensions and systolic and diastolic function were determined by transthoracic echocardiography with tissue Doppler imaging and 2-dimensional speckle tracking. Measurements were performed at baseline and after a 12-week period of high-intensity aerobic cycle training. LV internal diastolic diameter and LV mass were similar in the 2 groups at baseline and increased by ≈2% to 8% (P=0.04–0.0007) with training in both groups. Left atrial end-diastolic and end-systolic volumes were similar for both groups and increased by 23% to 36% (P=0.0006–0.0001) with training. Systolic function assessed by mean global strain was similar in both groups at baseline and increased by ≈8% (P=0.0004) with training in the postmenopausal group. LV displacement increased by ≈3% (P=0.04) in the premenopausal women only. Diastolic function assessed by E/A ratio was similar at baseline and increased by ≈7% (P=0.01) in the premenopausal group and 11% (P=0.0001) in the postmenopausal group with training.

Conclusions—These results suggest that training-induced cardiac adaptations are preserved in the early postmenopausal phase. Furthermore, the hormonal changes associated with the menopausal transition do not appear to affect cardiac dimensions and function.

Clinical Trial Registration—URL: http://www.clinicaltrials.gov. Unique identifier: NCT02135575. (J Am Heart Assoc. 2017;6: e005469. DOI: 10.1161/JAHA.117.005469.)

Key Words: E/A ratio • exercise training • menopause • transthoracic echocardiography • women

Cardiovascular disease is the leading cause of death in women globally and constitutes a major socioeconomic burden for society.1,2 Risk factors for cardiovascular disease include smoking, physical inactivity, age, metabolic syndrome, and loss of estrogen in women.2 Estrogen has been shown to have a protective effect on the cardiovascular system in women and the risk for cardiovascular events increases markedly after the menopausal transition when substantial hormonal changes occur, including the loss of estrogen production.3 However, the role of age versus hormonal changes for cardiovascular risk after menopause in women remains debated,4 in part because of the fact that comparisons of premenopausal and postmenopausal women are often conducted with significant age differences between the groups.5 We and others have recently provided support for the notion that the menopausal transition per se leads to rapid impairments in vascular function.6,7 In Nyberg et al,7 leg vascular function, assessed by infusion of vasoactive compounds, was markedly impaired in recent postmenopausal women compared with age-matched premenopausal women. Although evidence for altered cardiac structure and function exists in older postmenopausal women,8,9 it is unclear to what extent these alterations are detectable in the first few years after the menopausal transition.

Aerobic exercise training is well known to induce improvements in maximal oxygen uptake (VO2max) in sedentary...
Cardiac Adaptations, Menopause, and Exercise

Clinical Perspective

What Is New?
- This study shows that aerobic training improves cardiac structure and function in middle-aged women irrespective of menopausal status.

What Are the Clinical Implications?
- Implementation of aerobic training can be used as a safe prophylactic strategy to oppose deteriorations in cardiac function known to be prevalent in older postmenopausal women.

methods of early changes. Moreover, cardiac dysfunction determined by TDI has been shown to be an independent predictor of mortality.

The aim of the present study was 2-fold: (1) to compare cardiac dimensions and systolic and diastolic function in premenopausal and early postmenopausal women with a small age difference and (2) to assess the efficacy of 12 weeks of high-intensity aerobic training on cardiac adaptations in premenopausal and recent postmenopausal women. We hypothesized that recent postmenopausal women would have similar cardiac dimensions but impaired systolic and diastolic function compared with premenopausal women and that high-intensity aerobic exercise training would be more effective in inducing cardiac adaptations in premenopausal than postmenopausal women.

Methods

Participants

Participants were recruited from the Copenhagen area through advertisement in local newspapers. Study participants were premenopausal or postmenopausal women with an age range of 45 to 57 years (Table 1). The participants were all informed about potential risks and discomforts associated with the study and written consent was signed before the study experiments.

The study was approved by the ethics committee of Copenhagen and Frederiksberg municipalities Region H (H-1-2012-150) and conducted in accordance with the guidelines of the Declaration of Helsinki. All participants signed an informed consent form before participation in the study. The study was registered at ClinicalTrials.gov (NCT02135575).

The premenopausal women were all healthy with regular menstrual cycles. The postmenopausal women were all healthy and had not experienced a menstrual cycle during the previous 12 months but were less than 8 years past their final menstrual period. Menopausal status was verified by measurements of hypothalamic and reproductive hormones from a blood sample (estradiol <0.20 nmol/L and follicle-stimulating hormone: 22–138 IU/L, indicative of postmenopausal status). If the blood samples were indicative of perimenopause hormone levels, the women were excluded. Inclusion criteria were age 45 to 57 years, body mass index <30, and <2 hours of light to moderate physical activity per week of age. Exclusion criteria were smoking, use of hormonal contraceptives, hormone replacement treatment, prescription medicine, hypertension, cardiovascular disease, renal dysfunction, insulin resistance, diabetes mellitus, and hypercholesterolemia.

None of the participants had pathological ECG findings that were incompatible with the exercise intervention on examination. To ensure that all participants were normotensive,
blood pressure (BP) was measured 7 consecutive times by an automatic upper-arm BP monitor (M7, OMRON) after at least 15 minutes of rest in the supine position. Heart rate (HR) was measured during the BP monitoring.

The data presented are part of a larger study on the effects of exercise training on women in the menopausal transition funded by the University of Copenhagen Excellence Programme for Interdisciplinary Research and the data presented in Table 1 have previously been published.7,34

Study Design

The study was part of a larger study on the effect of menopause and physical activity on cardiovascular and metabolic health. The overall study design was a prospective intervention study with one group of premenopausal and one group of postmenopausal women. The women were selected to be close in age to minimize age as a confounder. All women underwent a 12-week period of high-intensity aerobic training with 3 sessions per week. For the present part of the study, the women underwent echocardiography measurements, a test for VO2max, and measurements of BP before and after the intervention period.

Exercise Training Intervention

All participants underwent the exercise training intervention. The training was performed on a cycle ergometer (Body Bike). Instructors from the research group supervised 2 training sessions per week and instructors from a local fitness center supervised 1 session. HR was monitored during all training sessions (TEAM2 Wearlink+, Polar). The training sessions were characterized by ≈50 minutes of intermittent high-intensity intervals where the participants reached HRs over 85% of maximum HR (Table 2). For more information and details on the participants’ variation in exercise intensities during the training see Nyberg et al7 and Mandrup et al.34

Echocardiography

Transthoracic echocardiography was performed using a GE Vivid 9 ultrasound machine with a 2.5-MHz transducer (GE
Table 2. Echocardiographic measures before and after 12 weeks of high aerobic intensity cycle training

<table>
<thead>
<tr>
<th>Echocardiographic Variables</th>
<th>Premenopausal n=36</th>
<th>Training Effect</th>
<th>Postmenopausal n=37</th>
<th>Training Effect</th>
<th>Group Differences</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Unit</td>
<td>Baseline</td>
<td>12 Weeks</td>
<td>P Value</td>
<td>Unit</td>
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<tr>
<td>LV morphology</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LVIDD cm</td>
<td>4.5±0.1</td>
<td>4.6±0.1</td>
<td>0.0007</td>
<td>4.5±0.1</td>
<td>4.6±0.1</td>
</tr>
<tr>
<td>LVIDS cm</td>
<td>2.8±0.8</td>
<td>2.7±0.8</td>
<td>0.064</td>
<td>2.9±0.8</td>
<td>2.8±0.8</td>
</tr>
<tr>
<td>LVEDV mL</td>
<td>93±2.6</td>
<td>101.7±2.6</td>
<td>0.0006</td>
<td>95.4±2.5</td>
<td>97.6±2.5</td>
</tr>
<tr>
<td>LVESV mL</td>
<td>37.5±1.7</td>
<td>42.0±1.7</td>
<td>0.054</td>
<td>40.3±1.7</td>
<td>41.8±1.7</td>
</tr>
<tr>
<td>LV stroke volume mL</td>
<td>55.6±1.9</td>
<td>59.7±1.9</td>
<td>0.064</td>
<td>55.1±1.9</td>
<td>55.8±1.9</td>
</tr>
<tr>
<td>LV mass index g/m²</td>
<td>76.9±2.5</td>
<td>82.1±2.5</td>
<td>0.001</td>
<td>81.4±2.4</td>
<td>87.6±2.4</td>
</tr>
<tr>
<td>LV mass g</td>
<td>136±5</td>
<td>145±5</td>
<td>0.001</td>
<td>142±5</td>
<td>152±5</td>
</tr>
<tr>
<td>IVSD cm</td>
<td>0.92±0.2</td>
<td>0.92±0.2</td>
<td>0.91</td>
<td>0.94±0.02</td>
<td>0.97±0.02</td>
</tr>
<tr>
<td>IVSS cm</td>
<td>1.32±0.04</td>
<td>1.37±0.04</td>
<td>0.23</td>
<td>1.35±0.04</td>
<td>1.37±0.04</td>
</tr>
<tr>
<td>LV mass W / g</td>
<td>1.64±0.05</td>
<td>1.80±0.05</td>
<td>0.002</td>
<td>1.74±0.05</td>
<td>1.78±0.05</td>
</tr>
<tr>
<td>LV morphology</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>EF, biplane</td>
<td>%</td>
<td>59.3±1.3</td>
<td>58.8±1.3</td>
<td>0.79</td>
<td>57.9±1.3</td>
</tr>
<tr>
<td>GLS %</td>
<td>−19.8±0.4</td>
<td>−20.2±0.4</td>
<td>0.27</td>
<td>−19.3±0.4</td>
<td>−20.8±0.4</td>
</tr>
<tr>
<td>s' TDI, mean cm/s</td>
<td>6.1±0.2</td>
<td>6.3±0.3</td>
<td>0.41</td>
<td>5.7±0.2</td>
<td>5.8±0.2</td>
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<tr>
<td>Pulsed TDI s', mean cm/s</td>
<td>8.3±0.2</td>
<td>8.0±0.2</td>
<td>0.12</td>
<td>7.6±0.2</td>
<td>7.4±0.2</td>
</tr>
<tr>
<td>LV displacement mm</td>
<td>12.4±0.2</td>
<td>12.8±0.2</td>
<td>0.04</td>
<td>11.8±0.2</td>
<td>12.1±0.2</td>
</tr>
<tr>
<td>Systolic function</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mitral inflow velocity E/A ratio</td>
<td>1.29±0.05</td>
<td>1.38±0.05</td>
<td>0.01</td>
<td>1.20±0.04</td>
<td>1.32±0.04</td>
</tr>
<tr>
<td>Mitral deceleration time ms</td>
<td>207±8</td>
<td>185±8</td>
<td>0.02</td>
<td>190±8</td>
<td>196±8*</td>
</tr>
<tr>
<td>E' / e'</td>
<td>...</td>
<td>6.5±0.3</td>
<td>7.1±0.3</td>
<td>0.001</td>
<td>7.0±0.3</td>
</tr>
<tr>
<td>Pulsed TDI e', mean cm/s</td>
<td>9.1±0.2</td>
<td>8.9±0.2</td>
<td>0.33</td>
<td>9.2±0.2</td>
<td>9.0±0.2</td>
</tr>
<tr>
<td>Pulsed TDI e', mean cm/s</td>
<td>11.6±0.3</td>
<td>11.3±0.3</td>
<td>0.18</td>
<td>10.1±0.3</td>
<td>10.1±0.3</td>
</tr>
<tr>
<td>IVRT, mean ms</td>
<td>86±2</td>
<td>87±2</td>
<td>0.54</td>
<td>96±2</td>
<td>94±2</td>
</tr>
<tr>
<td>Mitral valve E velocity m/s</td>
<td>0.8±0.02</td>
<td>0.87±0.02</td>
<td>0.0002</td>
<td>0.77±0.02</td>
<td>0.82±0.02</td>
</tr>
<tr>
<td>Mitral valve A velocity m/s</td>
<td>0.64±0.02</td>
<td>0.65±0.02</td>
<td>0.54</td>
<td>0.66±0.02</td>
<td>0.64±0.02</td>
</tr>
<tr>
<td>Left atrium</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LAEDV mL</td>
<td>22.9±1.5</td>
<td>28.4±1.5</td>
<td>0.0006</td>
<td>23.5±1.5</td>
<td>31.6±1.5</td>
</tr>
<tr>
<td>LAEDV index mL/m²</td>
<td>12.9±0.8</td>
<td>16.3±0.8</td>
<td>0.0004</td>
<td>13.3±0.8</td>
<td>18.2±0.8</td>
</tr>
<tr>
<td>LAESV index mL/m²</td>
<td>29.4±1.2</td>
<td>36.1±1.2</td>
<td>&lt;0.0001</td>
<td>30.3±1.2</td>
<td>37.8±1.2</td>
</tr>
<tr>
<td>LAESV mL</td>
<td>51.9±2.2</td>
<td>63.3±2.2</td>
<td>&lt;0.0001</td>
<td>51.2±2.2</td>
<td>65.6±2.2</td>
</tr>
<tr>
<td>Right ventricle</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>TAPSE cm</td>
<td>2.59±0.1</td>
<td>2.62±0.1</td>
<td>0.63</td>
<td>2.55±0.1</td>
<td>2.55±0.1</td>
</tr>
<tr>
<td>Pulsed TDI s' cm/s</td>
<td>13.6±0.3</td>
<td>13.9±0.4</td>
<td>0.34</td>
<td>13.4±0.3</td>
<td>13.3±0.3</td>
</tr>
</tbody>
</table>

EF indicates ejection fraction; GLS, global longitudinal strain; IVRT, isovolumic relaxation time; IVSD, left ventricular interventricular septum diastole; IVSS, left ventricular interventricular septum systole; LAEDV, left atrial diastolic volume; LAESV, left atrial systolic volume; LV, left ventricular; LVIDD, left ventricular end-diastolic diameter; LVIDS, left ventricular end-systolic volume; LVIDD, left ventricular end-diastolic diameter; LVSD, left ventricular ejection fraction; LVSS, left ventricular end-systolic volume; LVESV, left ventricular posterior wall thickness systole; TAPSE, tricuspid annular plane systolic excursion; TDI, tissue Doppler imaging.

Values are expressed as mean±SEM. *P<0.05: significant delta change with training between group.
Healthcare). The echocardiographic examinations were performed by 2 investigators but the examination before and after the training intervention were always performed by the same investigator for each participant. During the examination, participants rested in a supine position on their left side in a darkened room. All examinations were analyzed offline in unidentifiable random order, using the Echo Pac software version BT 13.0 (GE Healthcare). To reduce analytical variability, all echocardiographic examinations were analyzed by one of the investigators.

All participants were examined with the same protocol as previously described. In short, cardiac structure was evaluated from parasternal long axis 2-dimensional recordings according to current guidelines. The calculated left ventricular (LV) mass was indexed according to body surface area ($m^2 = 0.20247 \times \text{height} [m]^{0.725} \times \text{weight} [kg]^{0.425}$, DuBois formula). LV volume and LV ejection fraction (%) were evaluated using Simpson’s biplane method. Peak transmirtal blood inflow velocity in early (E) and late (A) diastole and the corresponding E/A ratio was measured using pulsed-wave Doppler in the apical 4-chamber view. Peak systolic velocity ($s'$; cm/s), peak early diastolic velocity ($e'$; cm/s), and peak late diastolic velocity ($a'$; cm/s) were measured with pulsed-wave TDI in the 6 LV segments at the level of the mitral annulus. The value of $e'$ represented the average of the segments of early peak diastolic velocities. Color TDI (TDI color) at a frame rate $>120/s$ was evaluated at the 6 mitral annular sites and values were averaged. Measurements included peak systolic velocity ($s'_\text{TDI color}$), peak early diastolic velocity ($e'_\text{TDI color}$), and peak late diastolic velocity ($a'_\text{TDI color}$). LV displacement was evaluated using tissue tracking. LV systolic function was evaluated by speckle tracking analysis where a semiautomated function tracks speckles from frame to frame identified within a specified region of interest and reported as the absolute value of global longitudinal strain.

Right ventricular function was evaluated as tricuspid annular plane systolic excursion and pulsed-wave TDI derived peak systolic velocity ($s'$).

**Determination of VO$_2$max**

VO$_2$max was measured with an Oxycon Pro (Intramedic, Denmark). The protocol was an incremental exercise test on a cycling ergometer (Monark, E9). The participants started with a 10-minute warm-up and thereafter the test was initiated with a start load of 50 W and increased by 25 W/min until volitional fatigue. Criteria for determination of VO$_2$max were: a plateau in VO$_2$, even with increased workload and/or respiratory exchange ratio $>1.1$ and/or an HR $>90\%$ of expected value. Two of 3 criteria had to be attained before the test was approved. The VO$_2$max tests as well as the echocardiography measurements were conducted in the weeks before the initiation of training and between 2 and 5 days after the training.

**HR Monitoring and Compliance of Training**

The participants had an individual HR monitor (TEAM2 Wearlink+, Polar) to record HR during the training sessions and ensure the training quality and that intervals reached a target HR $>85\%$ of maximum HR. All training sessions were registered throughout the training period.

**Statistical Analysis**

Our hypotheses could be directly evaluated as differences between groups and within groups in a linear mixed model framework. Fixed-effects factors were “group” (premenopausal, postmenopausal) and “time” (before training, after training). Between-subject variation was modeled using random effects. Model assumptions on homogeneity of variance and normal distribution were confirmed through residual and Q–Q plots. Pairwise differences were performed using post hoc t tests (based on the mixed models); no adjustment for multiplicity was applied. Data are reported as mean±SEM. The statistical analysis was executed with R version 3.2.2 (R Core Team, 2015) through the interface RStudio (RStudio Team [2015]. RStudio: Integrated Development for R. RStudio, Inc.) and the extension packages mice4 and multcomp. A significance level of 0.05 was used. Sample size was calculated on the basis of detecting a 10% change in E/A ratio with training, with a power of 0.8 and a significance level of 0.05.

**Results**

**Participant Characteristics**

A total of 83 participants (43 premenopausal and 40 postmenopausal) were initially recruited. A CONSORT flow diagram for the participants has been included in Mandrup et al. Ten participants were either excluded or dropped out because of illness (1), during run-in (4), because of insufficient training adherence (2), and because of pregnancy (1). Two premenopausal women were not assessed by echocardiography for logistic reasons. Therefore, a total of 36 premenopausal and 37 postmenopausal women were included in the study. All of the included participants fulfilled the training intervention and participated in the echocardiography examination both before and after the intervention period. The premenopausal women participated on average in 37±6 training sessions (≈93%) and the postmenopausal women in
38±5 training sessions (≈95%). Systolic arterial BP was similar in the 2 groups at baseline and after the training period. In the postmenopausal group, diastolic BP decreased (P=0.03) during the training period (by –4.3%) with no change in systolic BP. BP did not change significantly with training in the premenopausal group. It is important to point out that BP data were previously reported on these groups of women and were found to be unaltered with training. The reason for this discrepancy is likely that the number of included participants was lower in the present group and there was a slight difference in statistical approach.

HR at rest was similar in the 2 groups both at baseline and after the training period. There was a decrease (P=0.0001) in resting HR after the training period in the postmenopausal group (5.2%), but the change in HR at rest in the premenopausal group did not reach statistical significance. Data on low-density lipoprotein cholesterol, high-density lipoprotein cholesterol, and triglycerides are presented elsewhere.

Cardiac Structure and Function

Intervariability and intravariability of measurements

A sample of 10 participants was reanalyzed for intraobserver and interobserver variability by 2 investigators. The results showed a variation between 0% and 15.2% for the different common variables for interobserver and between 0% and 6.7% for the intraobserver variation.

Cardiac dimensions

Left atrial end-diastolic volume index and left atrial end-systolic volume index were similar in both groups at baseline (Table 2 and Figure 1A). Training increased left atrial end-diastolic volume index by ≈26% (P<0.0004) and 36% (P<0.0001) and left atrial end-systolic volume index by ≈23% (P<0.0001) and 25% (P<0.0001) in the premenopausal and postmenopausal groups, respectively. LV anatomy (determined as LV end-diastolic volume), LV internal diastolic diameter, and LV mass index were similar in the 2 groups at
Baseline (Table 2 and Figure 1). Training increased LV end-diastolic volume by \( \approx 9\% (P=0.0006) \) in the premenopausal group (Figure 2B); LV internal diastolic diameter by \( \approx 3\% (P=0.0007) \) and 2\% (0.04) in the premenopausal and postmenopausal groups, respectively (Figure 2C); and LV mass index by \( \approx 7\% (P=0.001) \) and 8\% (0.001) in the premenopausal and postmenopausal groups, respectively (Figure 2D).

Interventricular septum width at diastole and LV posterior wall thickness diastole were similar in both groups at baseline and these variables remained unaltered after the intervention (Table 2).

**Systolic function**

LV ejection fraction was similar in the 2 groups at baseline and was not altered by the training period in either group (Figure 2A). Global longitudinal strain was similar in both groups at baseline and increased \( (P=0.0004) \) in the postmenopausal group only (by \( \approx 8\% \), Figure 2B). LV longitudinal displacement was similar in both groups at baseline (Figure 2D). In the premenopausal group, LV displacement was significantly higher \( (P=0.04) \) after the training period (by \( \approx 3\% \)), whereas no change was detected in the postmenopausal group. LV displacement was \( \approx 6\% \) lower \( (P=0.04) \) in the postmenopausal compared with the premenopausal group after the training period. Color TDI (s’ TDI color) was similar in both groups at baseline and after the training intervention (Table 2).

**Diastolic function**

The E/A ratio was similar in both groups at baseline and was higher in \( \approx 7\% (P=0.01) \) and 11\% \( (P=0.0001) \) in the groups after the training period (Figure 3A). Mitral valve deceleration time was similar in both groups at baseline (Table 2). None of the participants had detectable diastolic dysfunction.
Mitral valve deceleration time was significantly lower ($P=0.02$) in the premenopausal ($\approx 11\%$) but not in the postmenopausal group after the training period. The training-induced change in mitral valve deceleration time was different ($P=0.03$) between the 2 groups. The $e'$ was similar in the 2 groups before training and remained unaltered with training (Figure 3B and 3C). At baseline, isovolumetric relaxation time (IVRT) was $\approx 12\%$ lower ($P=0.002$) in the premenopausal compared with the postmenopausal group (Table 2). The training intervention did not alter IVRT in either group.

**Right ventricular function**

Tricuspid annular plane systolic excursion (Figure 2C) and right ventricular pulsed-wave TDI' was similar in both groups at baseline and remained unaltered with the training intervention.

**Discussion**

The present study compared cardiac structure and functions in premenopausal and postmenopausal women and assessed the effect of a high-intensity aerobic training period in the 2 groups. Overall, the main findings were that cardiac dimensions and cardiac systolic and diastolic functions were similar in premenopausal and postmenopausal women separated by only 4 years of age. Moreover, a period of high-intensity aerobic training induced adaptations in cardiac dimensions as well as systolic and diastolic function in the 2 groups. These findings indicate that cardiac structure and function are not altered in the early postmenopausal phase and that cardiac adaptations to high-intensity aerobic training are preserved in recent postmenopausal women.

It is well established that estrogen has favorable effects on endothelial and cardiac function and loss of estrogen at menopause is associated with an increased risk of cardiovascular disease. In the present study, only minor differences in cardiac dimensions and systolic and diastolic function were detected between the premenopausal and recent postmenopausal women at baseline. This finding was therefore surprising and contrasts our own findings of reduced peripheral vascular function in postmenopausal women, as determined in a subgroup of the same premenopausal and postmenopausal women. The reason for this discrepancy is unclear; however, it could suggest that impairments in vascular function are manifested before changes in cardiac function are detectable. The current study was specifically designed to achieve a small age difference (4 years) between the groups, and therefore cardiac impairments caused by hormonal changes at menopause may not yet have occurred. This possibility is supported by a study showing that approximately half of postmenopausal women at a mean age of 62 years had mild diastolic dysfunction as indicated by E/A ratios $\leq 1.0$, whereas only very few women had E/A ratios $<1.0$ in the current study. Another possibility is that estrogen loss is more important for vascular function than for cardiac impairments, for which age may be a more influential factor. However, previous studies have shown that estrogen treatment improves diastolic function in postmenopausal women. Thus, at least with time, estrogen is likely to also impact cardiac function.

Figure 3. Left ventricular diastolic function evaluated as the ratio between early mitral inflow ($E$) and late ($A$) mitral inflow ($E/A$ ratio, $A$), peak early diastolic velocity ($e'$, $B$), and $E/e'$ ($C$) in premenopausal and postmenopausal women at baseline and after training. $*P<0.05$ significantly different from before training.
TDI and 2-dimensional speckle tracking used in the examinations of the participants allows for a more sensitive measure of early subclinical changes in systolic and diastolic function. Cardiac dysfunction determined by TDI has been shown to independently predict mortality. Previous studies have reported training-induced adaptations in systolic and diastolic function as assessed by TDI in men and premenopausal women, but no studies have examined training-induced cardiac adaptations in postmenopausal women. We compared a large number of parameters related to cardiac functionality in the contraction phase (systole) and the filling phase of the ventricles (diastole) before and after 12 weeks of high-intensity aerobic training. Within this relatively short time of training we were able to detect changes in a majority of the parameters determined, confirming the sensitivity of TDI for determination of cardiac functionality.

The finding of similar training-induced adaptations in the premenopausal and postmenopausal women rejects the study hypothesis of lesser adaptation in postmenopausal compared to premenopausal women. Our hypothesis was based on previous studies showing that postmenopausal women present lesser adaptations to training with regard to vascular function compared with age-matched men, and that improvements in vascular function only occur when combined with estrogen supplementation. However, one explanation for the discrepancy in findings could be differences in age, because the postmenopausal women in the study by Moreau et al and Pierce et al were older than the women in the present study. Numerous studies have shown that inactive aging has a marked influence on cardiac structure and function and it is plausible that the ability to respond to training accordingly is lost with age after menopause.

The magnitude of adaptations to training in the postmenopausal group, both in terms of dimensions and function, is similar to that observed after a period of football training in middle-aged and older men and premenopausal women, suggesting that high-intensity aerobic training provides an effective means of improving cardiac function in general, including in postmenopausal women. There may, however, be some difference in cardiac adaptation to training between men and women. This was evidenced in a previous study comparing the effect of 1 year of intensive endurance training on cardiac function and dimensions in young men and women. The study showed that the women had similar adaptations in cardiac function to men, whereas adaptations in cardiac dimensions were smaller and plateaued earlier in the women.

Conclusions

The present study demonstrates that cardiac dimensions and subclinical measures of cardiac function are similar in premenopausal and postmenopausal women with a small age difference. The finding indicates that, despite the known beneficial effects of estrogen on cardiac tissue, loss of estrogen and other hormonal changes at menopause, do not seem to alter cardiac dimensions or functionality in the near term. Moreover, a period of high-intensity aerobic exercise training led to significant and similar adaptations in cardiac dimensions and systolic and diastolic function in premenopausal and postmenopausal women. These findings contrast with the notion that postmenopausal women adapt less well to exercise training and clearly suggest that high-intensity aerobic training is an effective means of attaining beneficial cardiac adaptations in both premenopausal and recent postmenopausal women.

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Disclosures
None.

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