

# Tai Chi Is a Promising Exercise Option for Patients With Coronary Heart Disease Declining Cardiac Rehabilitation

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**Background**—More than 60% of patients decline participation in cardiac rehabilitation after a myocardial infarction. Options to improve physical activity (PA) and other risk factors in these high-risk individuals are limited. We conducted a phase 2 randomized controlled trial to determine feasibility, safety, acceptability, and estimates of effect of tai chi on PA, fitness, weight, and quality of life.

**Methods and Results**—Patients with coronary heart disease declining cardiac rehabilitation enrollment were randomized to a “LITE” (2 sessions/week for 12 weeks) or to a “PLUS” (3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks) condition. PA (accelerometry), weight, and quality of life (Health Survey Short Form) were measured at baseline and 3, 6, and 9 months after baseline; aerobic fitness (stress test) was measured at 3 months. Twenty-nine participants (13 PLUS and 16 LITE) were enrolled. Retention at 9 months was 90% (LITE) and 88% (PLUS). No serious tai chi–related adverse events occurred. Significant mean between group differences in favor of the PLUS group were observed at 3 and 6 months for moderate-to-vigorous PA (100.33 min/week [95% confidence interval, 15.70–184.95 min/week] and 111.62 min/week; [95% confidence interval, 26.17–197.07 min/week], respectively, with a trend toward significance at 9 months), percentage change in weight, and quality of life. No changes in aerobic fitness were observed within and between groups.

**Conclusions**—In this community sample of patients with coronary heart disease declining enrollment in cardiac rehabilitation, a 6-month tai chi program was safe and improved PA, weight, and quality of life compared with a 3-month intervention. Tai chi could be an effective option to improve PA in this high-risk population.

**Clinical Trial Registration**—URL: <http://www.clinicaltrials.gov>. Unique identifier: NCT02165254. (*J Am Heart Assoc.* 2017;6:e006603. DOI: 10.1161/JAHA.117.006603.)

**Key Words:** cardiac rehabilitation • coronary heart disease • physical exercise • risk factor • secondary prevention

Approximately 1 of 4 survivors of an acute coronary event will experience another cardiovascular event or die within the following 5 years.<sup>1,2</sup> This unfavorable prognosis is, however, avoidable. The aftermath of an acute coronary event can represent, in fact, a critical window for behavioral change.<sup>3</sup> For example, smoking cessation rates of 60% are reported in long-term smokers in the period immediately after

a diagnosis of myocardial infarction.<sup>4</sup> Physical activity (PA) is another critical target for post-coronary event behavioral change. Substantial epidemiological evidence indicates that PA delays the development of atherosclerosis and is associated with reduced risk of future coronary events.<sup>5,6</sup> Each 1–metabolic equivalent increase in fitness is associated with a 19% decrease in the risk of cardiovascular mortality.<sup>7,8</sup> Cardiac rehabilitation (CR) programs leverage this critical window for behavioral change by offering exercise training and education on coronary risk factor modification and reduce both all-cause and cardiovascular mortality.<sup>9–12</sup> Given these substantial benefits, current guidelines recommend referral to CR after an acute coronary event, percutaneous coronary interventions, and coronary bypass surgery.<sup>13</sup>

The large majority of this high-risk population, however, misses the opportunity for healthy behavioral changes offered by the participation in CR, because <40% of eligible patients join these programs in the United States.<sup>14</sup> Older men, women of all ages, and ethnic minorities are particularly unlikely to enroll.<sup>15</sup> Reasons for nonenrollment include dislike

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## Clinical Perspective

### What Is New?

- Compared with a shorter intervention, a 6-month tai chi exercise intervention was safe, feasible, and enjoyable and increased moderate-to-vigorous physical activity among deconditioned patients with coronary heart disease who had declined enrollment in cardiac rehabilitation.
- Other benefits included weight loss and improvements in quality of life.
- This is the first study showing that tai chi may improve exercise behaviors in this high-risk population.

### What Are the Clinical Implications?

- Tai chi is a promising and safe exercise alternative for patients with coronary heart disease who are unable or unwilling to attend traditional cardiac rehabilitation.
- If proved effective in larger studies, tai chi could be offered as an alternative exercise option within existing cardiac rehabilitation programs or within the context of community-based rehabilitation programs.

of exercise, perception of exercise as tiring or painful, low exercise capacity, travel distance to the CR center, high copays, and psychosocial factors, such as depression and low social support.<sup>16–19</sup> In addition, CR programs have high attrition rates for reasons ranging from the limited number of available exercise options to the lack of social interactions during programs.<sup>18,20</sup> These discouraging data have prompted calls for the “design of evidence-based, alternative approaches to traditional CR that help provide affordable access to clinically effective secondary prevention interventions (p. 2954).”<sup>14</sup>

Tai chi, a traditional Chinese martial art based on gentle body movements accompanied by relaxation and breathing exercises,<sup>21</sup> could be a promising exercise option for patients not attending CR. First, the exercise intensity during tai chi can be adjusted by varying session duration, body position during practice, and training style to meet the needs of elderly and deconditioned individuals.<sup>22</sup> Second, because tai chi is safe even in high-risk patients,<sup>23</sup> it can be offered in community centers,<sup>24,25</sup> thus overcoming transportation barriers associated with CR. Third, tai chi gentle training may be particularly attractive for women, who are less likely than men to attend CR and typically dislike the type of exercise offered at CRs.<sup>19</sup> Fourth, tai chi energy expenditure can reach 4.6 metabolic equivalents (moderate-intensity aerobic activity)<sup>26,27</sup> and has been shown to improve self-reported PA<sup>28–30</sup> and other coronary risk factors.<sup>31</sup> Studies have also suggested that tai chi may improve cardiac fitness.<sup>28</sup> Last, the emphasis placed on breathing and relaxation exercises during tai chi may improve psychological distress.<sup>31</sup> Recent research

has shown that the integration of stress reduction within a comprehensive CR program results in additional clinical benefits compared with CR alone.<sup>32</sup>

To our knowledge, no study has to date evaluated whether tai chi could be offered as an alternative exercise option for patients who declined enrollment in CR. In preparation for a future large efficacy trial, we conducted a phase 2 study to determine which dose of tai chi is most feasible, acceptable, and safe (primary outcome) and to obtain preliminary estimates of effect of each dose on PA and aerobic fitness (secondary outcomes). Other secondary outcomes included quality of life (QOL) and body weight.

## Methods

### Population

Methods for this study have been described in detail elsewhere.<sup>33</sup> Briefly, participants were recruited via flyers placed in cardiology practices, community centers, and other public venues; on-line resources; and advertisements on local radio programs and community newspapers. Because tai chi was offered in a class format, we recruited 3 consecutive cohorts of participants. Inclusion criteria were as follows: (1) history of coronary heart disease; (2) ability to understand and speak English; (3) aged  $\geq 21$  years; and (4) being physically inactive (7 days PA Recall Questionnaire)<sup>34</sup> and defined as not meeting current recommendations for PA<sup>35</sup> (ie, 30 minutes of moderate-intensity aerobic PA 5 d/week or 20 minutes of vigorous-intensity activity 3 d/week). Exclusion criteria included the following: (1) inability or unwillingness to give informed consent; (2) unstable angina; (3) blood pressure  $>200/110$  mm Hg or orthostatic systolic blood pressure decrease  $>20$  mm Hg; (4) uncontrolled atrial or ventricular arrhythmias; (5) third-degree AV block; (6) pericarditis or myocarditis; (7) recent thrombophlebitis; (8) abnormal stress test result without study cardiologist's clearance; (9) medical conditions likely to limit life span to  $<1$  year; (10) New York Heart Association functional class IV; (11) severe cognitive impairment (Blessed Orientation-Memory-Concentration test scores  $\geq 10$ )<sup>36</sup>; (12) orthopedic problems prohibiting tai chi practice; (13) ongoing tai chi or other mind-body training; (14) current enrollment in CR; (15) severe depression (Hospital Anxiety and Depression Scale depression subscale scores,  $>14$ ); and (16) conditions likely to affect study compliance (ie, moving out of study area and substance use).

Once interest and preliminary eligibility were confirmed, participants were invited for a screening visit. After providing in-person informed consent, participants completed baseline assessments and entered a run-in period of 1 week during which they wore accelerometers for the assessment of

baseline level of PA and underwent a symptom-limited exercise stress test for evaluation of cardiac risk<sup>37</sup> and baseline aerobic fitness. A cardiologist (W-C.W.) gave the final clearance for the participant's enrollment in the study. Participants received a \$40 stipend at each study visit. The Rhode Island Hospital (Providence) Institutional Review Board approved the study protocol (docket 599729).

The randomization schedule (1:1 ratio) was generated in "R" (<http://cran.us.r-project.org>)<sup>38</sup> on the basis of a permuted block randomization procedure using small randomized blocks. The allocation table was uploaded to an Access database; once all assessments were completed, the research assistant (J.K.) assigned each participant to either condition by clicking the "randomize" button.

## Tai Chi Interventions

Classes were held in a room located in the CR building. For each cohort, we offered 3 1-hour tai chi classes/week during weeks 1 to 12. Patients in the LITE (2 sessions/week for 12 weeks) group were invited to come twice a week, and patients in the PLUS (3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks) group were invited to attend 3 times a week. Participants in the PLUS group continued with 2 classes weekly during week 13 to 16, followed by 1 maintenance class every other week (weeks 17–24). Both groups were instructed to practice tai chi at home at least 3 times per week using a DVD (30-minute duration) containing the same sequence of tai chi exercises taught in class. Regardless of dose assignment, the content of the intervention was identical in the 2 groups.

A panel of experts in CR, cardiovascular prevention, tai chi, and exercise physiology reviewed the protocol used in previous studies conducted by our group in patients with heart failure and chronic obstructive lung disease.<sup>29,39</sup> On the basis of the panel's recommendations, a tai chi research expert (P.M.W.) worked with the tai chi instructors (3 certified instructors with 10+ years' experience) to modify existing protocols by increasing aerobic intensity over time (ie, by gradually adjusting the pace of practices [slightly faster over time] and depth of stances [slightly lower over time]).<sup>33</sup>

Intervention fidelity was assessed according to the Treatment Fidelity Workgroup guidelines.<sup>40</sup> Tai chi sessions were video recorded, and the study auditor (P.M.W.) reviewed 10% of videos for protocol consistency using a checklist developed for the study.

## Blinding

Participants were informed they would be randomized to 1 of 2 doses of tai chi and, thus, were not blinded to their dose allocation. Study personnel, other than the research assistant

responsible for participant randomization and monitoring of adverse events, were blinded.

## Measures

Assessments were performed at baseline and at 3, 6, and 9 months after enrollment. Data were collected using Access software for direct data entry from study participant into electronic surveys. Study data were collected between January 2015 and December 2017 and analyzed in March–April 2017.

## Primary Outcomes

Indicators of feasibility were retention rates, tai chi session attendance (recorded by the research assistant at each session), and individual tai chi home practice (self-reported using a log to be kept daily and collected in class each week).

Acceptability was evaluated at the end of the intervention using a self-reported rating scale ranging from 0 (did not like) to 4 (liked very much).<sup>29</sup>

For safety, the PA Readiness questionnaire was used to determine before each session whether the participant had experienced any chest pain, shortness of breath, dizziness, or muscle injury since the previous class.<sup>41</sup> At each study visit, we collected information about unplanned clinic visits and hospital readmissions. Adverse events were validated with information from medical records whenever possible.

## Secondary Outcomes

PA was measured using accelerometers. Accelerometers were not worn during tai chi classes. Measures includes number of valid days, wear time, min/week of moderate-to-vigorous intensity PA (MVPA), min/week of MVPA recorded in bouts of at least 10 minutes, and min/week of light activity. Data were processed using ActiLife software, with minimum wear time set at 10 h/d and nonwear time set at 60 minutes. Given the increased burden associated with participants wearing the accelerometer for a second week in case of mechanical failure of the accelerometer, the 7-day PA Recall Questionnaire was used as a backup.<sup>34</sup> Although absolute amounts of PA differ between these 2 measures, there is a good correlation between PA Recall Questionnaire and accelerometry collected data.<sup>42</sup>

For aerobic fitness, oxygen consumption in metabolic equivalents was calculated in all patients from stress test parameters, according to the protocol used in the test. Formulas for the Bruce Protocol and Modified Bruce Protocol were based on the American College of Sports Medicine's metabolic equations,<sup>43,44</sup> which are functions based on sex and time on the treadmill. Because of logistical problems, we were able to obtain direct measures of oxygen uptake ( $\text{VO}_2$ ) at

peak exercise (mL/kg per minute) using breath-by-breath expired gas analysis only in a subset of participants (n=21). There were no significant differences in baseline demographics between those with and without a direct measure of VO<sub>2</sub> (all  $P>0.05$ ).

Body weight was measured on a calibrated balance beam scale. QOL was assessed using a short version (8 items) of the Health Survey Short Form, a self-reported measure widely used for routine monitoring and assessment of care outcomes in adult patients.<sup>45</sup>

Covariates included sociodemographic characteristics (age, sex, race/ethnicity, education, and income) measured using validated forms; medical history (coronary risk factors, coronary revascularization procedures, ejection fraction, New York Heart Association class, comorbidities, and medications), abstracted from medical records; body mass index (calculated as weight [kg]/height [meters squared]); depression (using the Hospital Anxiety and Depression Scale,<sup>46</sup> a self-administered questionnaire with 2 subscales [score range, 0–21] measuring anxiety and depression, with higher scores indicating greater psychological morbidity); and patients' expectations about the intervention.<sup>47</sup> We also gathered information about reasons for not attending CR using a questionnaire developed for this study, as in the study by Dunlay et al.<sup>48</sup>

## Statistical Analyses

All analyses were conducted according to the intention-to-treat principle (with the exception of the attendance outcome, for which both intention-to-treat and per-protocol analyses were conducted). *t* Tests, nonparametric tests, and  $\chi^2$  tests, as applicable, were used to compare between-group baseline differences as well as retention rates, attendance rates, and individual home tai chi practice.

Unadjusted average min/week of MVPA over time were calculated separately by group and between-group differences in unadjusted mean and median MVPA at each time point and were tested using *t* tests and Mann-Whitney tests, respectively. To estimate treatment effects, we used a longitudinal mixed-effects regression model to simultaneously regress absolute MVPA (min/week) at each follow-up on treatment group, time, group×time, and baseline MVPA. Models included a subject-specific intercept to account for repeated measures within individual over time. A statistically significant interaction indicates that between-group differences vary across time points.<sup>49</sup> Contrasts were used to test between-group differences in MVPA at each follow-up time, controlling for baseline. Baseline variables that differed between groups and were predictive of PA were included as covariates in the multivariate model.<sup>50</sup> Because of the large variances in mean MVPA over time, we attempted to transform towards

normality. Because there were no substantial differences in the resulting model, we present the untransformed version in the Results, for ease of interpretation. Similar mixed-effects models were used to estimate mean between-dose differences for light activity and other secondary outcomes. All study analyses were performed using SAS, version 9.3, statistical software. Before study inception, we established to deem either dose feasible if at least 80% of participants were retained at the end of each intervention, if they attended at least 70% of the planned classes, and if they completed at least 70% of the prescribed home tai chi exercises. The intervention was considered acceptable if 80% of participants rated it a 4 (liked very much).

## Results

Participants were on average 67.9 (SD, 10.3) years old, 27% were female, and 93% were white, with almost two thirds reporting at least some college education (Table 1). Overall, this was a high-risk population: half of the study participants were diabetic, two thirds had high cholesterol levels or hypertension, more than half were obese, and one third were smoking at enrollment. More than 60% had a history of myocardial infarction, and 80% had had a percutaneous coronary intervention. Although baseline characteristics were overall balanced by randomization, LITE participants were slightly older, had a higher body mass index, were less physically active, and were more likely to have heart failure. The most common reasons for not having attended CR included dislike of intensive exercise and the perception that CR was dangerous (indicated by more than half of study participants), followed by insurance, scheduling, and transportation issues.

Of the 214 subjects initially contacted, 108 were ineligible, and 46 were not interested in participating. The most common reasons for refusal were schedule conflicts and lack of interest. Of the 33 randomized participants, 4 left the study immediately after randomization (Figure 1) and 29 started the intervention (13 in the PLUS group and 16 in the LITE group).

## Primary Outcomes

Retention rates were 90% (94% in the LITE group and 88% in the PLUS group) at the 9-month follow-up visit, with no significant differences by condition. Three participants were unavailable for follow-up, and 2 discontinued the intervention because of serious adverse events but remained in the study. Among the intention-to-treat sample, 39% of PLUS participants and 69% of LITE participants attended at least 70% of planned classes (Table 2). Participants' ratings showed high satisfaction, with 95% indicating that tai chi was highly enjoyable (score,  $\geq 3$  on a range of 0–4) and 100% reporting

**Table 1.** Baseline Characteristics of the Study Sample

Characteristics	Overall (N=29)	PLUS Group (n=13)	LITE Group (n=16)
Age, mean (SD), y	67.9 (10.3)	65.2 (10.5)	70.1 (9.9)
Female sex	7 (26.9)	3 (25.0)	4 (28.6)
Race			
American Indian or Alaskan Native	1 (3.4)	1 (7.7)	0
Asian or Pacific Islander	0	0	0
Black	1 (3.4)	1 (7.7)	0
Hispanic/Latino	0	0	0
White (not Hispanic)	27 (93.1)	11 (84.6)	16 (100)
Education			
High school or less	8 (27.6)	5 (38.5)	3 (18.8)
College or some college	15 (51.7)	7 (53.8)	8 (50.0)
Postgraduate	6 (20.7)	1 (7.7)	5 (31.3)
Current smoker	8 (27.6)	3 (23.1)	5 (31.2)
Diabetes mellitus	14 (48.3)	6 (46.2)	8 (50.0)
High cholesterol	22 (75.9)	8 (61.5)	14 (87.5)
BMI, mean (SD), kg/m <sup>2</sup>	30.4 (5.8)	28.3 (3.8)	32.2 (6.8)
Normal (<25 kg/m <sup>2</sup> )	4 (20.0)	3 (33.3)	1 (9.1)
Overweight (25–<30 kg/m <sup>2</sup> )	7 (35.0)	2 (22.2)	5 (45.5)
Obese (≥30 kg/m <sup>2</sup> )	9 (45.0)	4 (44.4)	5 (45.5)
Hypertension	23 (79.3)	9 (69.2)	14 (87.5)
Angina	23 (79.3)	11 (84.6)	12 (75.0)
Myocardial infarction	17 (58.6)	9 (69.2)	8 (50.0)
Heart failure	7 (24.1)	2 (15.4)	5 (31.3)
CABG	9 (31.0)	7 (53.8)	2 (12.5)
PCI	24 (82.8)	8 (61.5)	16 (100)
Implantable cardioverter defibrillator	3 (10.3)	1 (7.7)	2 (12.5)
Cerebrovascular disease	3 (10.3)	2 (15.4)	1 (6.3)
Peripheral artery disease	2 (6.9)	1 (7.7)	1 (6.3)
Cancer	4 (13.8)	1 (7.7)	3 (18.8)
COPD	2 (6.9)	0	2 (12.5)
Depression, mean (SD) HADS scores	4.5 (3.5)	4.6 (2.3)	4.5 (4.3)
Medications			
Antiplatelets	27 (93.1)	12 (92.3)	15 (93.8)
Anticoagulants	3 (10.3)	1 (7.7)	2 (12.5)
β blockers	26 (89.7)	12 (92.3)	14 (87.5)
ACE inhibitors	11 (37.9)	6 (46.2)	5 (31.3)
Statins	24 (82.8)	11 (84.6)	13 (81.3)
Intervention expectation scores, mean (SD)	12.6 (2.2)	12.1 (2.3)	12.9 (2.2)
Reasons for not attending CR*			
Does not like high-intensity exercise	18 (62.1)	7 (53.9)	11 (68.8)
CR too dangerous	17 (58.6)	6 (46.2)	11 (68.7)
CR schedule not convenient	12 (41.3)	5 (38.5)	7 (43.8)

Continued

Table 1. Continued

Characteristics	Overall (N=29)	PLUS Group (n=13)	LITE Group (n=16)
No insurance coverage	8 (27.5)	4 (30.8)	4 (25.0)
No transportation	7 (24.1)	2 (15.4)	5 (31.3)
Professional/personal commitments	6 (20.7)	1 (7.7)	5 (31.2)

Values are n (%) unless otherwise indicated. ACE indicates angiotensin-converting enzyme; BMI, body mass index; CABG, coronary artery bypass graft; COPD, chronic obstructive pulmonary disease; CR, cardiac rehabilitation; HADS, Hospital Anxiety and Depression Scale; LITE, 2 sessions/week for 12 weeks; PCI, percutaneous coronary intervention; and PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks.

\*Those responding “agree” to “strongly agree.”

they would recommend it to a friend. Average acceptability scores were similar between conditions (Table 2). Three serious adverse events were observed (1 in the LITE group and 2 in the PLUS group); none was related to the tai chi intervention (Table 3). No cardiac adverse events occurred during the study. The most common adverse events were musculoskeletal, and none was deemed related to tai chi practice.

## Secondary Outcomes

Complete accelerometer data were available for all participants. Unadjusted means of MVPA (min/week) at each time point are presented in Table 4 and Figure 2. We found significant between-group differences in mean MVPA at baseline, 3 months, and 6 months ( $P=0.04$ ,  $P=0.04$ , and  $P=0.02$ , respectively) in favor of the PLUS group. Because PA outcomes are subject to large variances (particularly in small samples), we also examined between-group differences in median MVPA over time. Results suggested significant differences in unadjusted median MVPA at 9 months ( $P=0.03$ ).

Regression models adjusting for baseline MVPA and including subject-specific intercepts indicated significant treatment effects on mean MVPA at 3 and 6 months favoring the PLUS groups. Specifically, between-group differences in MVPA were 100.33 min/week at 3 months (95% confidence interval [CI], 15.70–184.95 min/week) and 111.62 min/week at 6 months (95% CI, 26.17–197.07 min/week). Effects at 9 months trended towards significance ( $P=0.08$ ) in favor of the PLUS group. In addition, MVPA did not significantly decrease between 6 and 9 months in the aggregated sample ( $P>0.05$ ). We also tested a model that controlled for accelerometer wear time and variables not balanced by randomization, but model fit was not improved and these variables were removed from the final model.

Unadjusted average maximum  $VO_2$  ( $VO_2$  max) at baseline and 3 months is presented in Table 5 (calculated and measured). As shown by similarly low baseline  $VO_2$  max values, both groups were severely deconditioned, confirming this was a particularly frail population. There were no

significant between-group differences in calculated  $VO_2$  max at 3 months, controlling for protocol type and baseline differences between groups ( $\beta=0.22$ ; 95% CI,  $-1.91$  to  $2.35$ ). For the subset of participants in whom we were able to obtain direct measures of  $VO_2$  max, we also found no significant between-group difference in mean  $VO_2$  max at 3 months, controlling for baseline ( $\beta=-0.30$ ; 95% CI,  $-2.92$  to  $2.31$ ).

Unadjusted findings for weight and QOL are shown in Table 6. Longitudinal models suggest significant effects in favor of the PLUS arm for percentage change in body weight<sup>51</sup> at 3 months, controlling for baseline ( $\beta=3.31$ ; 95% CI,  $0.05$ – $7.09$ ). Specifically, mean percentage weight loss was higher in the PLUS group (2.86 kg; SD, 3.35 kg) than in the LITE group ( $-0.45$  kg; SD, 4.99 kg). Weight loss was maintained from 3 to 9 months.

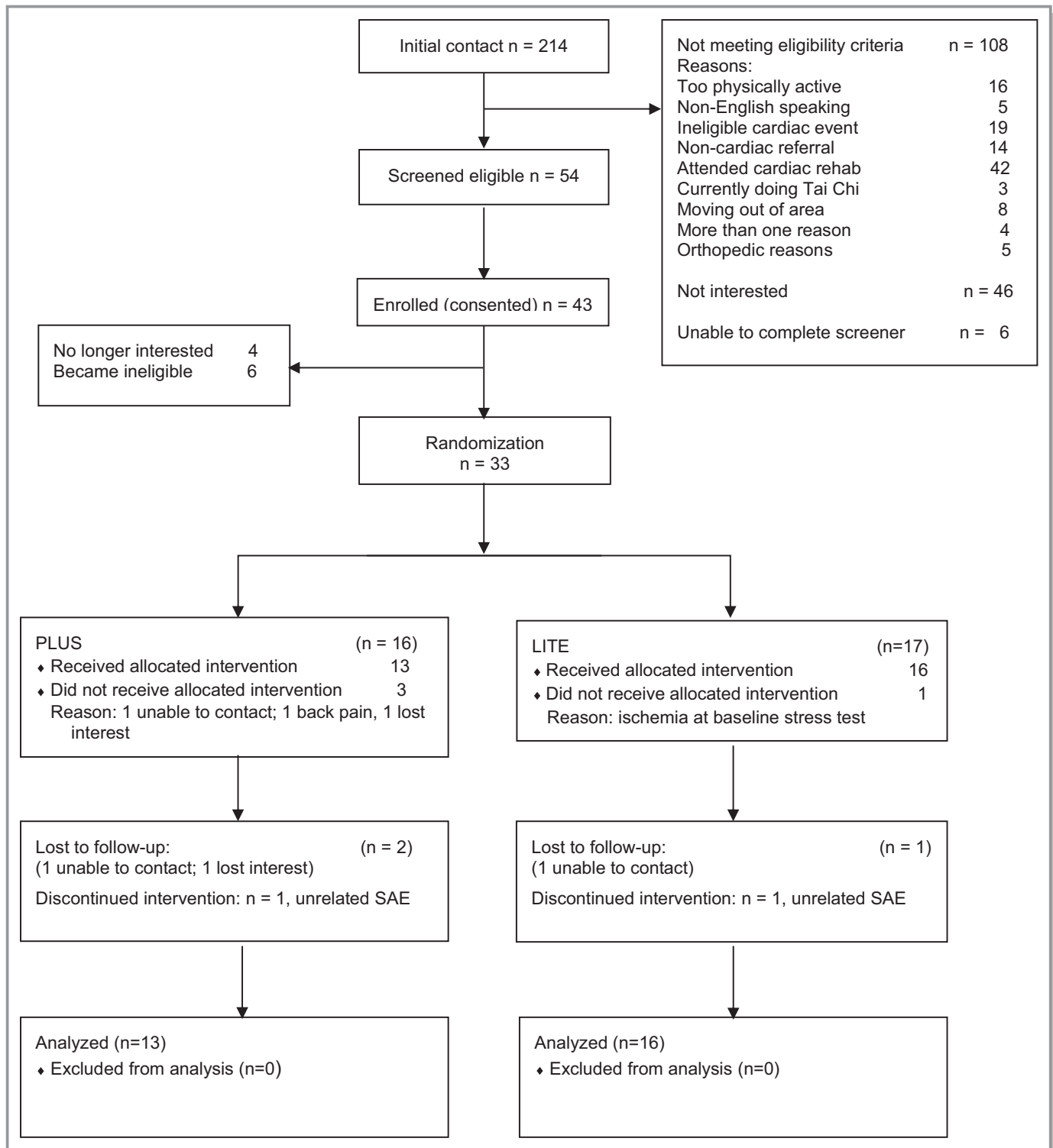
Models for QOL (where lower scores indicate better QOL) show a significant difference in favor of the PLUS condition at 3 months ( $\beta=-0.69$ ; 95% CI,  $-1.34$  to  $-0.04$ ). Effects at 6 and 9 months were in the same direction, albeit not significant ( $P=0.16$  and  $P=0.27$ , respectively).

## Post Hoc Analyses

Because this population was particularly deconditioned, we thought it would be important to assess the effect of the intervention on light activity. Models showed a significant effect on light activity at 3 and 6 months favoring the PLUS group (mean between-group differences in light activity were 397.88 [95% CI, 78.50–717.27] min/week and 376.75 [95% CI, 53.34–700.15] min/week), respectively. We observed a trend towards significance at 9 months ( $P=0.09$ ) in favor of the PLUS arm.

## Discussion

In this community-based sample of patients who declined enrollment in CR, we found that a longer (6-month) tai chi exercise intervention improved objectively assessed PA, weight, and QOL compared with a shorter (3-month) intervention. Overall, both interventions were safe, acceptable, and



**Figure 1.** Consort diagram. LITE indicates 2 sessions/week for 12 weeks; PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks; and SAE, serious adverse event.

feasible, with the exception of class attendance, which was slightly lower in the PLUS group.

Notably, once patients enrolled in the tai chi program, 90% were retained until the end of the study; those who dropped out

did so for serious adverse events and not because they disliked the program. This high retention rate compares favorably with dropout rates reaching up to 60% in CR.<sup>20</sup> With regard to safety, and consistent with a recent systematic review,<sup>23</sup> our findings

**Table 2.** Class Attendance and Acceptability Scores

Variable	Overall	LITE Group	PLUS Group	P Value
<b>ITT analysis</b>				
Attendance (classes attended/planned), %	66.0	69.4	53.2	0.19
Months 1–3, %		69.4	54.0	0.21
Months 4–6, %		NA	50.0	NA
Classes attended, mean (SD)		16.6 (7.2)	25.1 (16.4)	0.07
<b>Per-protocol analysis</b>				
Attendance (classes attended/planned), %	66.5	73.3	61.2	0.35
Months 1–3, %		73.3	62.5	0.36
Months 4–6, %		NA	59.9	NA
Classes attended, mean (SD)		17.5 (6.3)	29.2 (14.9)	0.01
Acceptability score, mean (SD)	3.6 (0.6)	3.6 (0.7)	3.7 (0.5)	0.61

ITT indicates intention to treat; LITE, 2 sessions/week for 12 weeks; NA, not applicable; and PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks.

indicate that tai chi is safe among deconditioned patients with coronary heart disease provided that participants are screened following the same criteria adopted for CR.

We found that participation in a longer tai chi program improved both moderate and light PA, effects that trended toward significance at 9 months in the PLUS group. Such a trend is important because it suggests a more enduring behavioral change that persisted even when support from the instructor and group was no longer present. This finding is particularly remarkable considering that our population was clearly resistant to behavioral change, as shown by the high proportion of current smokers and obese individuals in our study population.

Both fear of exercise and the perception of CR as dangerous were the most commonly reported reasons for declining participation in CR. Tai chi can clearly overcome these barriers because it is a different form of exercise. During training, participants are constantly reminded they do not need to strive or struggle to achieve predetermined goals in terms of heart rate or exercise intensity. Instead, they are invited to focus their attention on the breath and/or on the movements of the body. As a result, participants do not see tai chi exercise as threatening, and this may result in improvements in exercise self-efficacy. In fact, an analysis of data from a randomized controlled trial of tai chi for patients with heart failure has shown that increases in exercise self-efficacy mediated changes in functional status.<sup>52</sup>

**Table 3.** Adverse Events

Event	LITE Group	PLUS Group	Relatedness
Vaginal bleeding*	1	0	Definitely unrelated
Abdominal discomfort*	0	1	Definitely unrelated
Muscular chest pain*	0	1	Definitely unrelated
Sore throat	1	0	Definitely unrelated
Bruise on left thumb	0	1	Unlikely related
Bruise on left forearm	0	1	Definitely unrelated
Muscle cramps	0	1	Possibly related
Thoracic pain	1	1	Definitely unrelated
Back pain	1	0	Definitely unrelated
Cellulitis, right foot	1	0	Definitely unrelated
Fall	1	1	Definitely unrelated
Bronchitis	0	1	Definitely unrelated
Total	6	8	

LITE indicates 2 sessions/week for 12 weeks; and PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks.

\*Serious adverse events (participants were admitted to the hospital, but no serious condition was diagnosed as the cause of the patient’s complaint).

To our knowledge, this is the first study that examined the effect of tai chi on objectively assessed PA among patients not attending CR. One study conducted in outpatients with class I to III systolic heart failure found that participation in a 12-week tai chi training program improved self-reported PA (CHAMPS (Community Healthy Activities Model Program for Seniors/CHAMPS activities questionnaire)) compared with controls.<sup>29</sup> Another study reported similar results with a slightly longer intervention (16 weeks).<sup>30</sup> A second element of novelty is the use of tai chi as a possible alternative to CR-based exercise programs. To date, studies have examined the role of tai chi as an adjunct to CR exercise training,<sup>53,54</sup> whereas, to our knowledge, no study has investigated the possible role of tai chi as an alternative exercise option for patients not attending traditional CR.

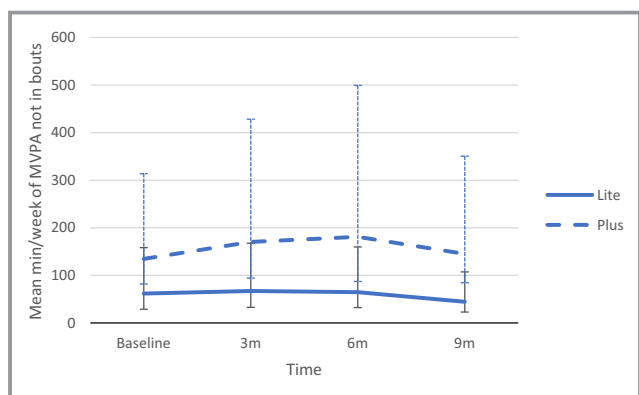
Increasing class frequency (2 versus 3 classes) did not affect aerobic fitness after 3 months of tai chi training

**Table 4.** Unadjusted MVPA Levels (Accelerometry)

MVPA, min/week	Overall	LITE Group	PLUS Group
Baseline	94.4 (96.9)	61.8 (64.8)	134.6 (116.0)
3 mo	105.8 (122.7)	67.1 (67.4)	170.3 (167.0)
6 mo	117.6 (154.2)	64.8 (63.7)	181.0 (205.7)
9 mo	87.6 (103.5)	44.4 (42.2)	145.1 (133.4)

Values are mean (SD). LITE indicates 2 sessions/week for 12 weeks; MVPA, moderate-to-vigorous physical activity; and PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks.





**Figure 2.** Unadjusted moderate-to-vigorous physical activity (MVPA), by study condition. LITE indicates 2 sessions/week for 12 weeks; and PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks.

compared with baseline. Findings from a recent meta-analysis suggest that tai chi produces modest improvements in  $VO_2$  max<sup>28</sup>; however, the duration of the tai chi intervention in the studies included in this meta-analysis ranged from 1 to 4.7 years.<sup>55–57</sup> It is possible that the duration of tai chi training may positively affect aerobic fitness; in fact, it has been suggested that a longer duration of tai chi training is needed to affect fitness.<sup>58</sup> For logistical reasons, we were unable to perform a third stress test at intervention completion among PLUS participants and, thus, we do not know if aerobic fitness would have increased after 6 months of training in this group. Furthermore, 1 of the aforementioned studies<sup>55</sup> assessed exercise intensity during steady-state tai chi performance and found average heart rates exceeding 70% of the maximal heart rate. Although we did not measure participants’ heart rate during exercise, another explanation for our results is that the intensity of tai chi intervention may not have been sufficient to improve peak  $VO_2$ . Finally, aerobic fitness did not improve despite improvements in overall PA. This finding is not surprising, given that several factors,

**Table 5.** Unadjusted Calculated and Measured  $VO_2$  Max

$VO_2$ max, mL/kg per minute	Overall	LITE Group	PLUS Group
<b>Calculated</b>			
Baseline	15.0 (7.5)	13.7 (8.0)	16.6 (6.8)
3 mo	15.0 (6.4)	13.8 (6.8)	16.6 (5.8)
<b>Measured</b>			
Baseline	11.8 (3.0)	10.8 (3.2)	13.0 (2.2)
3 mo	12.2 (3.6)	11.4 (3.8)	13.4 (3.2)

Values are mean (SD). n=28 for the calculated group, and n=21 for the measured group. LITE indicates 2 sessions/week for 12 weeks; PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks; and  $VO_2$  max, maximum oxygen uptake.

**Table 6.** Unadjusted Weight, Sleep Quality, and Quality-of-Life Scores

Variable	Baseline		
	Entire Sample	LITE Group	PLUS Group
<b>Weight, kg</b>			
Baseline	88.4 (24.3)	97.9 (29.1)	77.4 (9.8)
3 mo	90.5 (23.5)	98.0 (26.8)	79.3 (11.1)
6 mo	89.7 (24.0)	98.4 (27.9)	78.5 (11.0)
9 mo	90.3 (25.0)	98.7 (28.9)	78.7 (11.6)
<b>Sleep quality scores</b>			
Baseline	7.7 (4.3)	7.9 (4.8)	7.5 (3.6)
3 mo	6.7 (3.2)	6.8 (3.1)	6.7 (3.5)
6 mo	6.9 (3.6)	6.3 (3.4)	7.3 (3.8)
9 mo	6.8 (4.6)	6.0 (4.5)	8.0 (4.7)
<b>Quality-of-life scores</b>			
Baseline	2.7 (1.0)	2.9 (1.0)	2.5 (1.1)
3 mo	2.7 (0.9)	3.1 (0.9)	2.2 (0.8)
6 mo	2.9 (0.9)	3.1 (0.5)	2.6 (1.1)
9 mo	3.0 (0.7)	3.1 (0.5)	2.7 (0.9)

Values are mean (SD). LITE indicates 2 sessions/week for 12 weeks; and PLUS, 3 sessions/week for 12 weeks, then maintenance classes for 12 additional weeks.

including, among others, heritability,<sup>59</sup> age and adiposity,<sup>60</sup> and smoking,<sup>61</sup> influence aerobic fitness in addition to PA.

Our work has several important strengths. First, this was the first study to examine the effect of tai chi on objectively measured PA. Second, we were able to enroll a community-based sample of severely deconditioned, high-risk patients who had declined enrollment in CR. Third, retention rates at the end of the study were superior to those commonly reported in CRs. Last, we included rigorous assessments of intervention fidelity.

Our study has also some limitations. First, as expected, given the type of population selected, recruitment was a challenging aspect of this study. Because these patients had already declined a recommended treatment (CR), it is not surprising that they were hard to enroll in a research study of a new approach. Despite their history of coronary heart disease, one third of our patients were still smoking and 80% were overweight or obese, indicating that we were targeting a high-risk group with high resistance to behavioral change. Competition with CR in our catchment area was another problem, because an intensive enrollment campaign was launched while our study was ongoing. Second, the sample size was small, which, despite randomization, resulted in groups being unbalanced for certain baseline characteristics. However, baseline differences in activity and other characteristics were accounted for in analyses. Third, another

limitation is the underrepresentation of women and minorities. Fourth, our exercise protocol may not have been intense enough to increase aerobic fitness. Once participants feel more confident, improvements in aerobic fitness could be achieved (eg, by increasing the pace of exercises, modifying body position during practice [ie, squatting or raising hands over head], or simply providing patients with a heart rate monitor to self-monitor pace during practice). Finally, this study measured only aerobic fitness, whereas other important determinants of overall physical fitness (eg, balance, agility, and muscular strength) were not assessed.

## Conclusion

In sum, tai chi is a promising and safe exercise alternative for patients with coronary heart disease who are unable or unwilling to attend traditional CR, in particular for older people, women, and deconditioned individuals. If our findings are confirmed in larger studies, tai chi could be offered either as an alternative exercise training option within existing CR programs or within the context of community-based rehabilitation programs.

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## Disclosures

Salmoirago-Blotcher, Dunsiger, Bock, Breault, Wu, and Yeh have no conflict of interest to declare. Wayne is the founder and sole owner of the Tree of Life Tai Chi Center. Wayne's interests were reviewed and managed by the Brigham and Women's Hospital and Partner's HealthCare in accordance with their conflict of interest policies.

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**Tai Chi Is a Promising Exercise Option for Patients With Coronary Heart Disease Declining Cardiac Rehabilitation**

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