



Review article

The impact of Tai Chi and Qigong mind-body exercises on motor and non-motor function and quality of life in Parkinson's disease: A systematic review and meta-analysis



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ABSTRACT

Purpose: To systematically evaluate and quantify the effects of Tai Chi/Qigong (TCQ) on motor (UPDRS III, balance, falls, Timed-Up-and-Go, and 6-Minute Walk) and non-motor (depression and cognition) function, and quality of life (QOL) in patients with Parkinson's disease (PD).

Methods: A systematic search in 7 electronic databases targeted clinical studies evaluating TCQ for individuals with PD published through August 2016. Meta-analysis was used to estimate effect sizes (Hedges's g) and publication bias for randomized controlled trials (RCTs). Methodological bias in RCTs was assessed by two raters.

Results: Our search identified 21 studies, 15 of which were RCTs with a total of 735 subjects. For RCTs, comparison groups included no treatment ($n = 7$, 47%) and active interventions ($n = 8$, 53%). Duration of TCQ ranged from 2 to 6 months. Methodological bias was low in 6 studies, moderate in 7, and high in 2. Fixed-effect models showed that TCQ was associated with significant improvement on most motor outcomes (UPDRS III [ES = -0.444 , $p < 0.001$], balance [ES = 0.544 , $p < 0.001$], Timed-Up-and-Go [ES = -0.341 , $p = 0.005$], 6 MW [ES = -0.293 , $p = 0.06$], falls [ES = -0.403 , $p = 0.004$], as well as depression [ES = -0.457 , $p = 0.008$] and QOL [ES = -0.393 , $p < 0.001$], but not cognition [ES = -0.225 , $p = 0.477$]). I^2 indicated limited heterogeneity. Funnel plots suggested some degree of publication bias.

Conclusion: Evidence to date supports a potential benefit of TCQ for improving motor function, depression and QOL for individuals with PD, and validates the need for additional large-scale trials.

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1. Introduction

Parkinson's disease (PD) is one of the most common progressive neurodegenerative disorders, leading to loss of motor function and reduced quality of life. A growing body of evidence supports the role of exercise in improving both motor and non-motor outcomes

in PD [1–3]. In fact, exercise is now considered an integral part of the management of PD [4–6].

Tai Chi and Qigong (TCQ) are two increasingly popular mind-body interventions that have the potential to address a range of motor and non-motor symptoms associated with PD [7]. TCQ share a common history which includes elements of traditional Chinese medicine, martial arts conditioning, and Asian lifestyle philosophy. Both integrate balance, flexibility, and neuromuscular coordination training with a number of cognitive components, including heightened body awareness, focused mental attention, imagery, multi-tasking, and planned and goal-oriented training, which together may result in benefits to PD above and beyond conventional exercise [8]. In contrast to many styles of yoga or seated mind-body meditative practices, TCQ typically places a greater emphasis on standing and dynamic movements (e.g. pushing and lifting gestures linked to martial applications) that have the potential to impact gait, balance, and other functional activities. For these reasons, in the present work TCQ are grouped together and are considered equivalent interventions, paralleling other recent reviews [9–11].

The evidence for TCQ's effects on PD motor symptoms have been evaluated in a growing number of trials, with prior meta-analyses and systematic reviews generally showing positive support for clinical measures of balance [12–14]. Mixed or inconclusive results for other motor outcomes (e.g., gait, mobility, and falls) have typically been reported, in large part due to the few numbers of studies assessing these outcomes [12–14]. Even less attention has been devoted to systematically evaluating the effects of TCQ on non-motor outcomes [12], and no meta-analyses to date have evaluated key issues such as depression and cognitive function, which have been shown to be strongly linked to overall quality of life [15].

Research to date exploring possible neuromuscular and behavioral mechanisms underlying TCQ's effect on postural control add credibility to clinical findings, however, these studies have generally not included populations with PD. Studies largely focused on normal aging suggest that enhanced motor performance may be due to various underlying processes, including improved lower extremity strength and flexibility [16–22], proprioception and postural awareness [23–25], neuromuscular coordination and reaction time [26,27], executive function [28], and reduced fear of falling [29,30]. Less research has evaluated mechanisms that may contribute to improvements in non-motor function. TCQ related improvements in depression and anxiety have been hypothesized to be related to modest cardiovascular loading, breathing and imagery related changes in autonomic tone, cognitive restructuring leading to reduced rumination and catastrophizing, and indirect effects of enhanced motor confidence and enhanced self-efficacy [11,31–33]. These mechanisms, as well as multi-tasking and attention shifting training associated with TCQ, have been hypothesized to contribute to enhanced cognitive function [34], and some studies have reported correlations between TCQ-related improvements in cognitive function and brain neural architecture and resting state neural network activity [35,36]. However, these studies have not been conducted in PD populations.

Using a meta-analytic approach and incorporating a number of recent clinical trials not included in previous reviews [37–41], this study aimed to systematically evaluate and quantify the effects of TCQ on motor and non-motor function and quality of life in patients with PD.

2. Methods

2.1. Literature search

Electronic literature searches were performed using PubMed,

CINAHL, Web of Science, ProQuest Central, Science Direct, Scopus, and Cochrane Library for English language articles published until August 30, 2016. The search terms were Tai Chi, Taiji, Qigong, and Parkinson's disease. Additional manual searches based on references listed in the retrieved articles were performed to complete the search.

2.2. Eligibility criteria

Randomized controlled trials (RCTs) and prospective non-randomized controlled and observational studies published in English in which Parkinson's disease was the primary disease and Tai Chi and/or Qigong were the primary interventions were included. Formal meta-analysis was limited to RCTs, with non-RCTs used to qualitatively further inform synthesis of the overall evidence.

2.3. Study selection and data extraction

Study eligibility assessment was performed independently by two researchers (WG and KO) who applied eligibility criteria using a standardized protocol. Data were extracted by two reviewers (RS and MP) independently using a standardized template generated in Microsoft Excel. Data related to study design, duration and frequency of the intervention program, type of the control, sample size, and outcome measures were extracted for qualitative analysis in accordance with the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines (PRISMA) [42].

2.4. Risk of bias assessment

Two researchers (RS and MP) independently assessed the methodological quality of RCTs using the 10 item Cochrane Collaboration Tool for assessing risk of bias. Criteria include: random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data (attrition rates or ITT), selective reporting, eligibility criteria, groups similar at baseline (primary outcome), rationale for sample size, and other bias [43]. A summed score of 5 points or more was set as the threshold to be excluded in the analysis. The evaluated domains were assessed as 0 (no bias), 1 (minimal bias), or 2 (moderate or severe bias) according to the established criteria [43]. Any discrepancies in the evaluations conducted by two authors were discussed and resolved.

2.5. Safety monitoring

Studies were reviewed to identify if formal protocols for systematically monitoring adverse events were described, and if adverse events reported in the study were associated with the intervention.

2.6. Data analysis

Outcome data for formal meta-analysis using identified 15 RCTs with 735 subjects with PD were systematically extracted and organized in a database. For each outcome, data extracted included the mean and standard deviation (SD) of the pre-test and post-test values for each group, mean and SD of change scores in each group, t score or p-value within groups, and sample size (N) in each group. When these data were not available, data in the form of standard errors, confidence intervals or medians with ranges were converted into mean and SD format using previously suggested statistical formulas [43,44].

Comprehensive Meta-Analysis software v3 (CMA v3, Biostat, Inc. USA) was used for data synthesis. When an outcome of interest

(e.g., depression) was measured with multiple instruments, a pooled effect size was calculated by CMA. For continuous data, Hedge's g and 95% confidence intervals (CI) using a fixed effect model were calculated to estimate mean differences between groups for all eligible trials. The Q value and I^2 statistics were used to assess heterogeneity for the variation in true effect sizes across the included studies; a low p value for the Q statistics and/or I^2 index above 40% indicated significant heterogeneity [43]. Publication bias was assessed using funnel plots.

Pooled effect size for each outcome variable of motor and non-motor function was calculated. Subgroup analyses were also conducted according to the comparison group; alternative active interventions (e.g., exercise, health education, meditation) or no treatment (e.g., no contact control, wait list).

One study [45] only reported data combining multiple outcomes; the authors for this study were contacted and provided the raw data needed to extract separate scores for each outcome.

3. Results

3.1. Study selection and characteristics

Fig. 1 summarizes the flow of the literature search and selection process following PRISMA guidelines. An initial search identified 382 records from multiple databases and manual searches. Removing duplicates resulted in 200 records. The title and abstracts of these records were screened according to the inclusion criteria. A total of 28 full text articles met eligibility criteria and were further reviewed, with an additional 7 being excluded due to duplicated data in separate publications, or inappropriate study design. Of the remaining 21 eligible studies, 15 were RCTs and 6 non-RCTs, of which 11 RCTs and 4 non-RCTs have been included in previous systematic reviews or meta-analysis [37–41].

Among the 15 RCTs, Li and colleagues [38,46,47] published 3 papers from the same parent trial, with each paper emphasizing different outcome variables. Outcome variables duplicated across publications were excluded in analyses. Li and colleagues [38,46,47] also had more than two comparison groups; data for each comparison group were separately entered in the analysis with the sample sizes adjusted (i.e., divided by 2) to minimize over-influence of one study. Amano et al. [48] reported two research studies within one publication, each including unique interventions and control groups; these were treated as separate studies in analyses.

3.1.1. Participants characteristics and study setting

Table 1 summarizes the 21 studies (15 RCTs and 6 non-RCTs) which include a total of 838 subjects. Study participants all had diagnoses of idiopathic Parkinson's disease with Hoehn & Yahr scores ranging from 1 to 4 with an average of 2.3. The average age of the participants was 67.5 y; 58.4% of participants were male. All RCTs included patients on a stable PD medication regimen. All but two studies conducted testing in the on-medication state. The two studies evaluating outcomes in the off-medication state utilized a 12-hour pre-testing washout period [39,40].

3.1.2. Intervention and control group characteristics

Tai Chi was used as an intervention in 16 studies. In nearly all cases, a modified, simplified, or short form of Tai Chi was used. All interventions were delivered in a group setting, but one study [49] utilized home training sessions following only four 45-min group trainings. The duration of interventions ranged from 5 to 24 weeks. Individual intervention sessions ranged from 45 to 90 min, and the frequency of classes varied between 1 and 3 times per week. Studies varied in their prescriptions for home practice regimens. Nine studies compared TCQ with other alternative active

interventions (e.g., exercise, resistance training, walking or dance); 7 studies employed a no-treatment or wait-list control group, and 5 studies were uncontrolled.

3.1.3. Outcome measures

For motor function, UPDRS Part III was most commonly assessed (14 studies), followed by balance (9 studies), Timed-Up-and-Go (8 studies), 6 MW (6 studies), and falls (2 studies). For non-motor outcomes, quality of life was most frequently measured (9 studies), followed by depression (3 studies) and cognition (3 studies).

3.1.4. Adverse effects

None of the 15 RCTs reported protocols for monitoring adverse events. Two studies did mention safety was monitored without providing detail [50,51], but no safety related results were reported. Three separate studies reported no adverse events related to interventions were observed [41,47,52]. No serious adverse events were reported in any study.

3.1.5. Risk of bias assessment

The fifteen RCTs were assessed for risk of bias (Table 2). Randomization procedures were described in all studies, but 6 of them provided no specific details of procedures used. Allocation concealment was not mentioned in 6 studies (60%), while 14 studies (93%) reported blinding of outcome assessment. Seven studies (46%) reported either intention to treat approach or no dropouts. Homogeneity of baseline data was confirmed by most studies, but one study [51] reported significant differences in baseline values of an outcome variable. Twelve studies (80%) did not provide a rationale for the sample size. Overall rating of bias assessment indicated that 6 studies were methodologically strong (0–1 risk), 7 studies moderate (2–3 risk), and 2 studies were weak (4 risk). No studies met the pre-defined threshold for exclusion from the analyses.

3.2. Effects of TCQ on motor function

3.2.1. Unified Parkinson's disease rating scale (UPDRS) part III

Motor symptoms were assessed with the UPDRS III in 13 studies [37,39,40,45,46,48,49,51,53–57], including 11 RCTs. Lower scores reflect less severe motor symptoms. Meta-analysis of the 11 RCTs employing a fixed effect model indicated an overall small effect size (Hedges's $g = -0.444$, 95% CI -0.606 to -0.282 , $p < 0.001$) supporting that TCQ significantly improved motor function relative to all control groups. Q -value ($p = 0.732$) and I^2 (0%) indicate no heterogeneity. The subgroup analysis limited to 6 studies with active control groups revealed a slightly lower but statistically significant effect size ($ES = -0.368$, $p = 0.001$). The effect size limited to the 6 studies with a no-treatment control group was relatively larger ($ES = -0.555$, $p < 0.001$). Lack of heterogeneity persisted in subgroup analyses. Publication bias was not suspected based on the funnel test plot asymmetry. Of note, the two non-RCTs that assessed UPDRS did not report statistically significant within-group improvements [39,54] (see Fig. 2-A).

3.2.2. Balance

Balance was reported in 10 studies [40,41,45,46,49,50, 55,58–60], including 7 RCTs. Balance was assessed using the Berg Balance Scale, posturography, or single leg standing time. Results from a fixed effect meta-analysis model indicate that TCQ had greater improvements in balance than control groups with an overall medium effect size (Hedges's $g = 0.544$, 95% CI 0.356 to 0.730 , $p < 0.001$). Heterogeneity was not substantial based on Q value ($p = 0.156$) and I^2 (34.12%). Subgroup analysis limited to 3

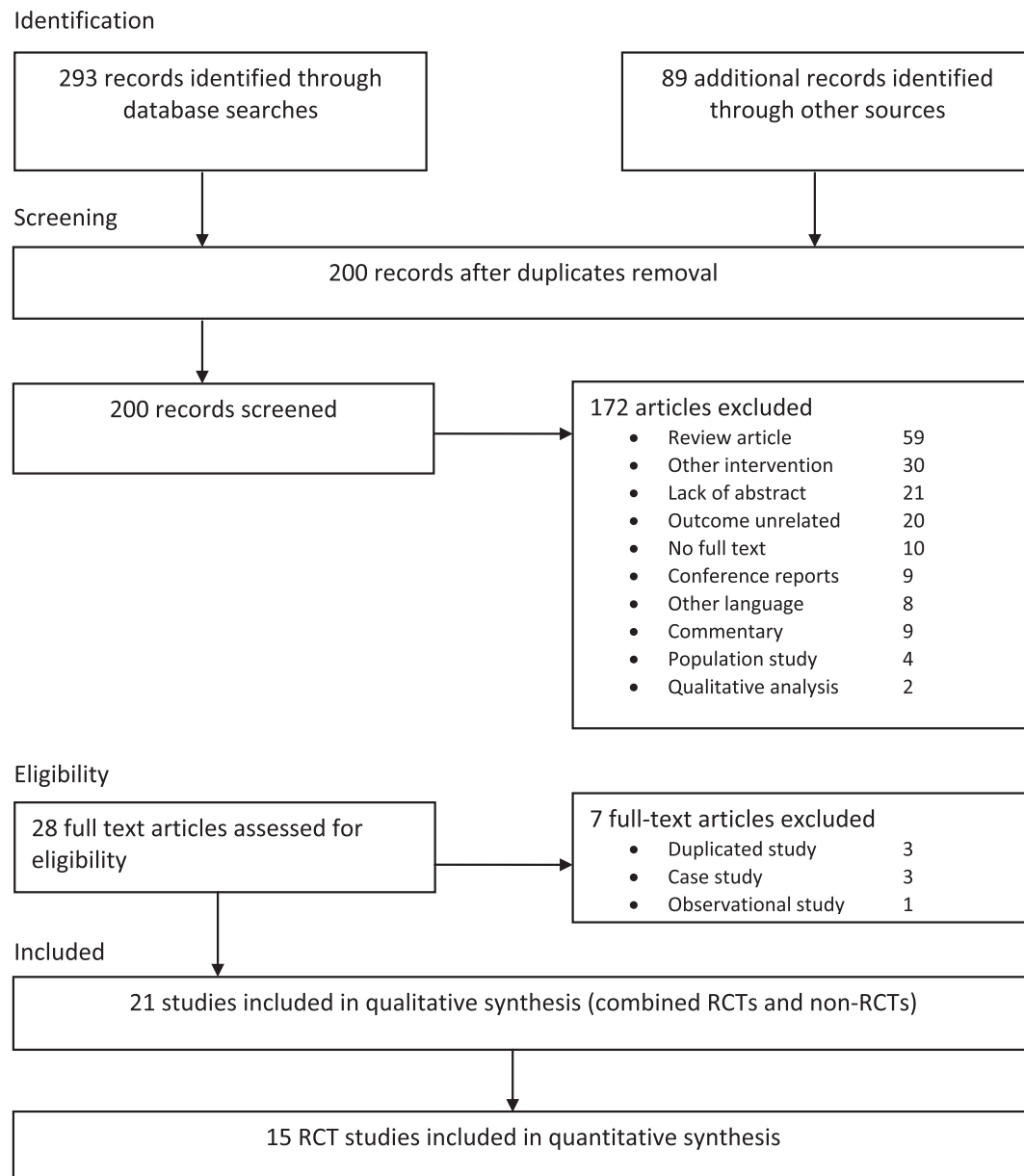


Fig. 1. Summary of the flow of our literature search according to the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) guidelines.

studies employing active control groups also showed a medium effect size ($ES = 0.477$, $p = 0.001$), as did subgroup analyses limited to 5 studies using no treatment control groups ($ES = 0.544$, $p < 0.001$). Lack of heterogeneity persisted in subgroup analysis. Three additional non-controlled studies [58–60] also reported within-group improvements in balance (see Fig. 2-B).

3.2.3. Timed-Up-and-go

Timed-Up-and-Go (TUG) was assessed in 7 studies [40,41,45,49,50,55,61], including 6 RCTs. Results of a fixed-effect model of RCTs indicate that TCQ significantly improved TUG compared with control groups, with a small effect size (Hedges's $g = -0.341$, 95% CI -0.578 to -0.104 , $p = 0.005$). Q value ($p = 0.277$) and I^2 (20.78%) indicate limited heterogeneity. Due to the small number of studies with active control groups, subgroup analysis was not performed. One non-RCT study [61] also reported within-group improvement in TUG (see Fig. 2-C).

3.2.4. 6-Minute walk (6 MW)

Four RCTs [40,45,53,55] assessed the effect of TCQ on the 6 MW test. A fixed-effect model indicated that TCQ did not significantly improve 6 min walking speed compared to control groups (Hedges's $g = -0.293$, 95% CI -0.603 to 0.017 , $p = 0.064$), although most studies showed greater improvement in performance in the TCQ group. Q value ($p = 0.356$) and I^2 (7.49%) showed limited heterogeneity. Due to the small number of studies, subgroup analyses were not performed (see Fig. 2-D).

3.2.5. Falls

Only 2 RCTs [38,49] assessed the effect of TCQ on the number of fall rates. A fixed-effect model indicated that TCQ significantly reduced fall episodes compared to control group (Hedges's $g = -0.403$, 95% CI -0.677 to -0.129 , $p = 0.004$). Q value ($p = 0.623$) and I^2 (0%) confirms limited heterogeneity. One non-controlled study [60] also reported that the PD patients experienced fewer

Table 1

Summary of Tai Chi/Qigong studies for individuals with Parkinson's disease.

Main Author (country)	Type of study	Sample (mean age)	Gender (M/F)	Med	Intervention	Frequency (per week)	Duration (weeks)	Control group	Measured outcomes
Burini 2006 (Italy)	RCT with cross-over	26 (65.2)	9/16	on	Qigong	50m, 3x	7	aerobic exercise	UPDRS, 6 MW*, MBS, PDQ, BDI, CPET
Amano 2013 (USA)	RCT	21 (66.0) 24 (66.0)	14/7 14/10	on	Tai Chi	1 h, 2x	16	Qigong meditation	UPDRS, gait
Choi 2013 (Korea)	RCT	22 (60.8)	NS	on	Tai Chi	1 h, 3x	12	no treatment	UPDRS, reaction time*, gait, TUG, 6 MW, OLS*
Kurlan 2015 (USA)	RCT	44 (72.0)	27/17	on	Tai Chi	1 h, 1x	16	no treatment	UPDRS, S&E, GDS, PDQ, falls
Li 2014 (USA)	RCT	195 (68.7)	122/73	on	Tai Chi	1 h, 2x	24	resistance training or stretching	UPDRS, 50-ft walk, VPS*, PDQ*
Li 2015 (USA)	RCT	176 (68.7)	122/73	on	Tai Chi	1 h, 2x	24	resistance training or stretching	Falls*, PDQ*, cost effectiveness*
Li 2012 (USA)	RCT	195 (68.7)	122/73	on	Tai Chi	1 h, 2x	24	resistance training or stretching	UPDRS*, gait*, TUG*, FRT*, muscle strength*
Schmitz-Hubsch 2006 (Germany)	RCT	56 (63.5)	43/13	on	Qigong	1.5 h/1x	8	no treatment	UPDRS*, MADRS, PDQ
Xiao 2015 (China)	RCT	96 (67.5)	67/29	off	Qigong	3 h training + 15m, 4x at home	24	walking	UPDRS*, 6 MW*, BBS*, PDSS*, TUG*, PFS, gait*
Zhang 2015 (China)	RCT	40 (54.0)	24/16	on	Tai Chi	1 h, 2x	12	multimodal exercise	UPDRS, BBS, TUG, gait
Gao 2014 (China)	RCT	76 (69.5)	50/26	on	Tai Chi	1 h, 3x	12	no treatment	UPDRS, TUG, BBS*, falls*
Hackney 2008 (USA)	RCT	26 (63.0)	21/5	on	Tai Chi	1 h, 2x	10	no treatment	UPDRS*, TUG, TS*, 6 MW*, OLS, BBS*, gait
Gladfelter 2011 (USA)	RCT	17 (72.0)	12/5	on	Tai Chi	1 h, 2x	12	no treatment	PDQ, BBS*, FRT*, TUG
Hackney 2009 (USA)	RCT	71 (66.0)	45/16	on	Tai Chi	1 h, 2x	13	tango, waltz, no treatment	UPDRS, PDQ
Nocera 2014 (USA)	RCT	21 (66.5)	11/10	on	Tai Chi	1 h, 3x	16	no treatment	cognition, PDQ*, TFES
Cheon 2013 (Korea)	non-RCT	23 (64.2)	0/23	on	Tai Chi	1 h, 3x	8	combined exercise or no treatment	UPDRS, QOL, BDI, S&E, functional fitness
Li 2007 (USA)	one group pre-post test	17 (71.5)	6/11	on	Tai Chi	1.5 h, 5x	5	None	50-ft walk, TUG, FRT
Kim 2014 (Korea)	one group pre-post test	12 (65.3)	2/10	on	Tai Chi	1 h, 3x	12	None	COP
Kim 2011 (Korea)	one group pre-post test	10 (78.5)	1/9	on	Tai Chi	1 h, 3x	12	None	COP
Wassom 2015 (USA)	one group pre-post test	7 (66.9)	3/4	off	Qigong	1 h, 1x + 40m daily at home	6	None	UPDRS, PDSS, PFS, cognition, gait
Loftus 2014 (USA)	one group pre-post test	34 (66.0)	17/17	on	Qigong	1 h, 1x	12	None	BBS, PDFFP, Pull test

Note. Med, anti-Parkinson medication ('off' refers to medication off during measurement). UPDRS, Unified Parkinson's disease rating scale; 6 MW, 6 min walking; MBS, modified Borg scale; PDQ, Parkinson's disease questionnaire; BDI, Beck depression inventory; CPET, cardiopulmonary exercise test; TUG, timed up and go; OLS, one leg standing; S&E, Schwab and England activities of daily living scale; GDS, geriatric depression scale; VPS, vitality plus scale; FRT, functional reach test; MADRS, Montgomery-Asperg depression rating scale; BBS, Berg balance scale; PDSS, Parkinson's disease sleep scale; PFS, Parkinson's fatigue scale; TS, tandom stance; TFES, Tinetti's falls efficacy scale; HRQOL, health related quality of life; PDFFP, Parkinson's disease fall profile; MMSE, Mini Mental State Examination.

*Indicates outcome significantly improved by Tai Chi/Qigong relative to control group.

Table 2

Quality assessment of the included studies.

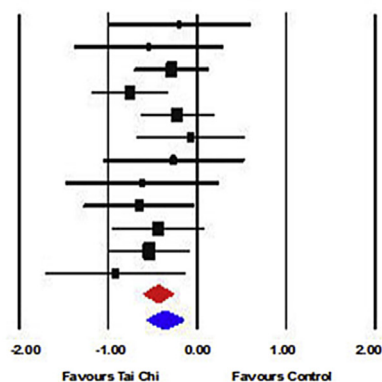
Study	Selection Bias		Performance bias		Attrition bias		Reporting bias		Eligibility criteria	Difference at baseline	Rational for sample size	Other bias	Total*
	Random sequence	Allocation concealment	Blinding of participants	Blinding of outcome assessment	Incomplete outcome data (ITT)	Selective reporting							
Burini2006	0	0	0	0	1	0	0	0	0	0	1	0	2
Amano2013	NS	NR	NR	0	0	1	0	0	0	0	1	0	3
Choi2013	NS	NR	NR	0	1	0	0	0	0	0	1	0	3
Kurlan2015	NS	NR	NR	0	1	0	0	0	0	0	1	UGS	3
Li2014	0	0	NR	0	0	1	0	0	0	0	0	0	1
Li2015	0	0	NR	0	0	1	0	0	0	0	0	0	1
Li2012	0	0	NR	0	0	1	0	0	0	0	0	0	1
Schmitz-Hubsch2006	0	0	NR	0	0	0	0	0	0	0	1	0	1
Xiao2015	NS	NR	NR	0	1	0	0	0	0	0	1	0	3
Zhang2015	0	0	NR	0	0	0	0	0	0	0	1	0	0
Gao2014	0	NR	NR	0	1	0	0	0	0	0	1	0	2
Hackney2008	0	NR	NR	0	0	0	0	0	0	0	1	0	1
Gladfelter2011	0	NR	0	0	1	0	0	0	0	0	1	0	2
Hackney2009	NS	NR	0	NR	1	0	0	0	1	1	1	0	4
Nocera2014	NS	NR	NR	0	1	1	0	0	0	0	1	0	4

Note. NS (not specified, counted as 1); NR (not reported); UGS (unequal group size).

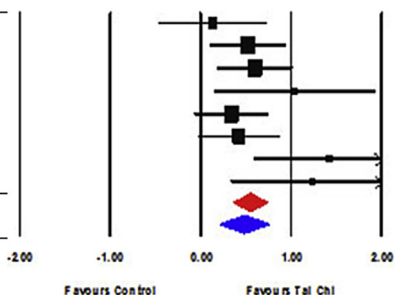
*Higher scores represent more bias.

A. UPDRS III

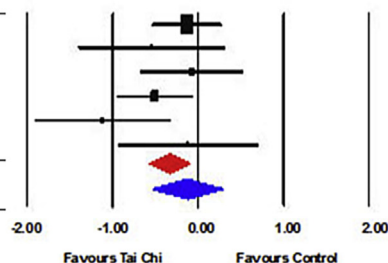
Study	Weight	Hedges's g	LL	UL	p
Burini 2006	4.04	-0.201	-1.007	0.605	0.625
Amano 2013-1	3.67	-0.544	-1.389	0.302	0.208
Li 2012-R	14.76	-0.291	-0.712	0.131	0.177
Li 2012-S	14.28	-0.755	-1.184	-0.326	0.001
Xiao 2015	15.39	-0.222	-0.635	0.191	0.292
Zhang 2015	7.11	-0.076	-0.683	0.532	0.807
Amano 2013-2	4.09	-0.268	-1.069	0.534	0.512
Choi 2013	3.51	-0.621	-1.487	0.244	0.159
Kurlan 2015	6.68	-0.655	-1.282	-0.027	0.041
Schmitz-Hubsch 2006	9.42	-0.440	-0.968	0.088	0.102
Gao 2014	12.78	-0.539	-0.992	-0.086	0.020
Hackney 2008	4.26	-0.918	-1.703	-0.133	0.022
Fixed effects model		-0.444	-0.606	-0.282	0.000
Subgroup with Active control		-0.368	-0.579	-0.158	0.001

**B. Balance**

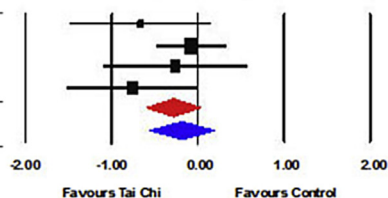
Study	Weight	Hedges's g	LL	UL	p
Zhang 2015	9.53	0.132	-0.476	0.740	0.670
Li 2012-R	19.40	0.522	0.096	0.949	0.016
Li 2012-S	19.61	0.599	0.175	1.023	0.006
Choi 2013	4.32	1.040	0.137	1.943	0.024
Xiao 2015	20.48	0.339	-0.076	0.754	0.109
Gao 2014	17.39	0.425	-0.025	0.876	0.064
Hackney 2008	5.01	1.421	0.582	2.260	0.001
Gladfelter 2011	4.26	1.237	0.327	2.147	0.008
Fixed effects model		0.544	0.356	0.731	0.000
Subgroup with Active control		0.477	0.207	0.746	0.001

**C. Timed Up & Go**

Study	Weight	Hedges's g	LL	UL	p
Xiao 2015	32.99	-0.132	-0.544	0.281	0.215
Choi 2013	7.58	-0.544	-1.404	0.317	0.532
Zhang 2015	15.19	-0.072	-0.680	0.536	0.817
Gao 2014	27.41	-0.506	-0.958	-0.053	0.029
Hackney 2008	8.68	-1.116	-1.920	-0.312	0.007
Gladfelter 2011	8.14	-0.118	-0.948	0.712	0.781
Fixed effects model		-0.341	-0.578	-0.104	0.005
Subgroup with Active control		-0.132	-0.544	0.281	0.532

**D. 6 minute walking**

Study	Weight	Hedges's g	LL	UL	p
Burini 2006	14.01	-0.665	-1.493	0.162	0.115
Xiao 2015	56.53	-0.075	-0.487	0.337	0.722
Choi 2013	13.36	-0.266	-1.114	0.581	0.538
Hackney 2008	16.09	-0.757	-1.529	0.016	0.055
Fixed effects model		-0.293	-0.603	0.017	0.064
Subgroup with Active control		-0.192	-0.561	0.177	0.397

**E. Falls**

Study	Weight	Hedges's g	LL	UL	p
Li 2015	63.47	-0.351	-0.695	-0.006	0.046
Gao 2014	36.53	-0.492	-0.944	-0.040	0.033
Fixed effects model		-0.403	-0.677	-0.129	0.004

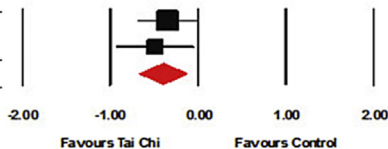


Fig. 2. Effects of Tai Chi/Qigong on motor function. Note 1. Data values indicate weight, effect size (Hedges's g) and confidence interval of LL (lower limit) to UL (upper limit). 2. Plots to the left of zero indicate negative effect sizes for all outcomes in favor of Tai Chi (i.e. fewer symptoms), except for balance where plots to the right of zero represent a positive effect.

falls at the beginning vs. at the end of a 6-month Qigong program (see Fig. 2-E).

3.3. Effect of TCQ on non-motor function and quality of life

3.3.1. Depression

Depression was assessed in 5 studies [37,45,53,54,56], including 4 RCTs. Depression was assessed using the Beck Depression Inventory, Geriatric Depression Scale, and Montgomery-Asperg Depression Rating Scale. A fixed-effect model indicated that TCQ significantly reduced depression scores compared to control groups, with an overall medium effect size (Hedges's $g = -0.457$, 95% CI -0.795 to -0.118 , $p = 0.008$). Q value ($p = 0.739$) and I^2 (0%) indicated limited heterogeneity. Due to the small number of studies, subgroup analyses were not performed. One non-randomized controlled study [54] also reported statistically significant improvements in depression when Tai Chi was compared to a no treatment group (see Fig. 3-A).

3.3.2. Cognitive function

Three studies [39,45,52], including 2 RCTs, assessed the effect of TCQ on cognition. Cognition in the RCTs was assessed with UPDRS Part I and Trail Making Tests. A fixed-model indicated that TCQ did not significantly improve cognition when compared to control groups (Hedges's $g = -0.225$, 95% CI -0.845 to -0.396 , $p = 0.477$), though both studies reported modest benefits in favor of TCQ. Q value ($p = 0.776$) and I^2 (0%) indicated limited heterogeneity.

One non-controlled study [39] also reported no significant improvements in cognition following 6 weeks of Qigong training (see Fig. 3-B).

3.3.3. Quality of life

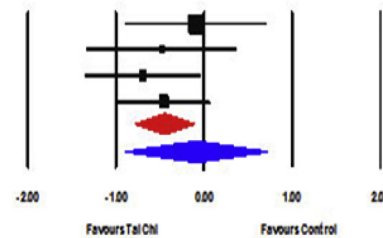
Quality of life was evaluated in 7 studies [47,50–54,56], including 6 RCTs. Five of the 6 RCTs assessed QOL using the Parkinson's Disease Questionnaire-39 (PDQ-39); 1 used the PDQ-8 short version. A fixed-effect model indicated that TCQ significantly improved quality of life (lower scores) compared with control groups, with an overall small effect size (Hedges's $g = -0.393$, 95% CI -0.612 to -0.174 , $p < 0.001$). Q value ($p = 0.527$) and I^2 (0%) indicated limited heterogeneity. Subgroup analysis limited to the 3 studies that employed active control groups indicated medium effect size ($ES = -0.502$, $p < 0.001$); the effect size was slightly smaller for the 4 studies that employed a no treatment control group ($ES = -0.230$, $p = 0.207$). Lack of heterogeneity persisted for subgroup analyses. One non-randomized controlled study [54] also reported statistically significant improvements in quality of life when Tai Chi was compared to a no treatment group (see Fig. 3-C).

4. Discussion

Due to recent changes in the understanding of the nature of PD, exercise is now widely recommended to individuals with PD for maintaining balance, mobility and overall quality of life. Nonetheless, few evidence-based guidelines are available to inform optimal

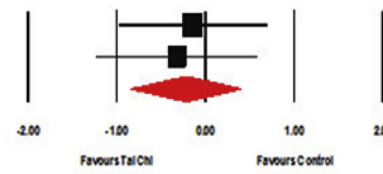
A. Depression

Study	Weight	Hedges's g	LL	UL	p
Burini 2006	17.70	-0.093	-0.897	0.711	0.821
Choi 2013	15.60	-0.481	-1.338	0.376	0.271
Kurlan 2015	25.70	-0.697	-1.365	-0.030	0.041
Schmitz-Hubsch 2006	41.00	-0.454	-0.982	0.075	0.092
Fixed effects model		-0.457	-0.795	-0.118	0.008
Subgroup with Active control		-0.093	-0.897	0.711	0.821



B. Cognition

Study	Weight	Hedges's g	LL	UL	p
Choi 2013	53.81	-0.142	-0.987	0.703	0.742
Nocera 2014	46.19	-0.322	-1.236	0.592	0.490
Fixed effects model		-0.225	-0.845	0.396	0.477



C. Quality of life

Study	Weight	Hedges's g	LL	UL	p
Burini 2006	7.42	-0.070	-0.874	0.734	0.864
Li 2013-R	25.94	-0.663	-1.093	-0.233	0.003
Li 2013-S	27.11	-0.466	-0.887	-0.045	0.030
Schmitz-Hubsch 2006	17.62	-0.041	-0.563	0.481	0.877
Gladfelter 2011	8.03	-0.083	-0.856	0.690	0.833
Hackey 2009	8.14	-0.693	-1.461	0.075	0.077
Nocera 2014	5.74	-0.331	-1.246	0.583	0.477
Fixed effects model		-0.393	-0.612	-0.174	0.000
Subgroup with Active control		-0.502	-0.784	-0.220	0.000

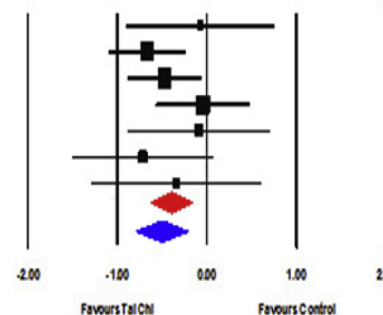


Fig. 3. Effects of Tai Chi/Qigong on non-motor function. Note 1. Data values indicate weight, effect size (Hedges's g) and confidence interval of LL (lower limit) to UL (upper limit). 2. Plots to the left of zero indicate negative effect sizes for all outcomes in favor of Tai Chi (i.e. fewer symptoms), except for balance where plots to the right of zero represent a positive effect.

exercise regimens. Extending prior meta-analyses and systematic reviews by including 5 recent studies (4 RCTs) not considered in prior reviews, our findings support the idea that TCQ may be effective in reducing multiple motor and non-motor symptoms for people with PD. With respect to motor symptoms, our meta-analyses indicate clinically relevant medium effect sizes in favor of TCQ for Part III of the UPDRS, clinical measures of balance, and fall reduction, and a small effect size for TUG. For non-motor outcomes, meta-analyses indicated clinically relevant small-to-medium effect sizes on quality of life and depression. Of note, for Part III of the UPDRS, balance, and quality of life, significant effect sizes persisted even when comparisons were limited to active controls. This finding suggests that the benefits of TCQ are likely to not solely be due to attention or psychosocial support factors, but instead, are the result of mind-body exercise-specific activities. In fact, the results of two trials support that TCQ may be superior to other conventional exercise interventions, though further comparative effectiveness studies are needed to confirm this observation and inform mechanisms leading to relative superiority [46,51]. Finally, although the evaluation of TCQ's safety within most trials that we included was not systematically assessed and was poorly described, there were no serious adverse events reported. This finding suggests that TCQ is likely to be safe for people with PD across multiple stages of disease progression [62].

4.1. Gaps in previous reviews

The overall conclusions of our study parallel conclusions reported in other recent reviews, but in addition to including updated evidence, the methods employed in our study differed in important ways. First, a number of prior reviews have been limited to qualitative synthesis and have not also included quantitative meta-analytical methods [63,64]. For studies that did include meta-analysis, one in 2014 focused only on Tai Chi (excluding Qigong) and included 9 trials in quantitative synthesis (vs. 15 in the present study) [7]. Another 2014 meta-analysis focused only on motor outcomes and included 8 trials [13]. A 2015 review that evaluated both Tai Chi and Qigong included 13 RCTs, however, 4 of these were published in Chinese and thus are not included in our review [14]. Finally, one 2016 review including meta-analytic methods employed broader inclusion criteria, synthesizing results from multiple mind-body modalities, including yoga and dance in addition to Tai Chi [12].

4.2. Effects of TCQ on motor function

The UPDRS motor subscale was the most commonly assessed measure of motor function in TCQ trials. This is the most widely used clinical scale to measure disease progression and severity and the response of motor function to therapy [65]. In many pharmacology and exercise studies, it is the primary outcome, thus it is worth discussing this outcome in more depth. For Part III of the UPDRS, we found an overall small effect size of -0.44 ; when analyses were limited to the 6 trials with active controls, effect size remained statistically significant (-0.37). When we narrowed our focus to the 4 randomized trials deemed to have lowest bias in our review (i.e., bias scores ≤ 1 out of 10), absolute changes in UPDRS following TCQ ranged from -6.4 to -1.5 (average = -3.7). This average magnitude of change meets the threshold for established clinically important differences (minimum of 2.5 for UPDRS to 3.5 for MDS-UPDRS) with variation in reported meaningful differences due to instrument used (UPDRS vs. MDS-UPDRS) and severity of disease progression [66,67]. The magnitude of effects of TCQ on Part III of the UPDRS that we observed are similar to those reported for physiotherapy [68], but lower than those reported following

training in Argentine Tango [69], or for patients taking dopamine medications [57] or undergoing deep brain stimulation [70].

Because of the significant variability in intervention duration for studies evaluating UPDRS motor subscale, we also conducted an exploratory subgroup analysis based on TCQ exposure length. TCQ protocols varied from 5 to 24 weeks across studies. A comparison of shorter-term (12 weeks or less) vs. longer-term (13 weeks or more) protocols indicated effect sizes of -0.121 ($p = 0.625$) and -0.426 ($p < 0.001$), respectively. This suggests that longer exposures may be more effective, and emphasizes the need for further dosing studies, including studies that explore multiple domains of dosing, such as combined hours of class and home practice and measures of progressions in proficiency of performance [71].

In addition to overall motor performance assessed with part III of the UPDRS, we also found significant improvements in clinical measures of balance and mobility. The 6 MW speed also showed trends towards improvement following TCQ training, but this effect was not statistically significant. Only two studies evaluated fall frequency, but given the public health concerns and impact of falls in PD, the positive finding we observed ($ES = -0.40$) warrants comment. Both included trials had low risks of bias. In the smaller of the two studies ($n = 76$) conducted by Gao and colleagues [49], 12 weeks of Tai Chi training resulted in a significant drop in fall incidence assessed 6 months post training, with 21.6% of Tai Chi group reporting falling at least once, compared to 48.7% in the control group. In the study by Li and colleagues [46] ($n = 195$), 24 weeks of Tai Chi training resulted in an average of 12% monthly fall rates in the Tai Chi group (estimated over a 9-month period); in comparison, resistance training and seated stretching controls exhibited 23% and 38% average fall rates, respectively. A subsequent analysis of the data of Li and colleagues concluded that Tai Chi is a cost effective approach for fall prevention in PD [38].

4.3. Effects of TCQ on non-motor outcomes

An important finding from this review is that Tai Chi positively impacts multiple non-motor symptoms, in addition to motor symptoms. The most common non-motor outcome evaluated in TCQ trials was QOL assessed in all RCTs using the PDQ-39 [72]. For QOL, we found an overall effect size of -0.39 ; when analyses were limited to the 3 trials with active controls, effects size actually increased slightly to -0.50 . In the largest trial conducted by Li et al. [46], QOL in the Tai Chi group improved by 38%; in comparison, QOL in the resistance training and stretching improved by 15% and $<1\%$, respectfully. This is similar to the magnitude of QOL improvements following deep brain stimulation (14–38%) [70]. Meta-analyses also indicated significant improvements in depression. In contrast, the effects of TCQ on cognition was small and not statistically significant, however this estimate was based on 2 small studies [45,52]. As cognitive deficits are now recognized as an undisputable and highly prevalent feature of PD, both due to natural disease progression and side effects of some medication [73], additional study of the effects of TCQ in cognition are warranted. In particular, given the demonstrated importance of executive function in gait and balance in PD [74,75], and the potential of Tai Chi to improve executive function in normal aging [34], studies evaluating the impact of TCQ on executive function and dual task performance may be particularly informative. Other outcomes not systematically addressed in TCQ studies to date, but that warrant further study based on the promise of mind-body interventions in other populations, include sleep [76] and autonomic function [77].

It has been suggested that the broad, multi-system and multi-symptom benefits of TCQ may result from its multi-component approach, incorporating training in motor control, postural awareness, cognition, breathing, and stress reduction [8,78]. In

normal aging, some progress has been made in identifying mechanisms of TCQ that may improve balance and gait, and reduce falls, including improving lower extremity strength and flexibility [16–22], proprioception and postural awareness [23–25], neuromuscular coordination and reaction time [26,27], executive function [28], and reducing fear of falling [29,30]. Preliminary studies also support a neural basis for enhancing cognitive and motor processes [35,36]. The mechanistic exploration of how TCQ impacts motor and non-motor symptoms in PD is a rich area for further research.

4.4. Methodological issues and limitations

An important methodological decision in this study was to use fixed-vs. random-effects models for meta-analyses. Our decision was based on a number of factors and follows established guidelines [79]. First, we began with the statistical assumption that the sample of studies we evaluated was similar enough to represent the true effect sizes for each outcome assessed in trials of TCQ for PD. This assumption is supported by the facts that within each motor and non-motor outcome, there was strong similarity in instruments used, and that study populations across studies were similar with respect to disease progression. This assumption is also supported by I^2 and Q statistics that indicated low between-study heterogeneity in effect size. Finally, it has been suggested that random-models only be used when >5 studies are included in analyses [79]. As this was not the case in some subgroup analyses, we chose a fixed-effect model approach. While not reported in our results, exploratory analyses of effect sizes for all outcomes based on random models were found to be either equal to or larger than the fixed-effect results we reported, with only minor qualitative differences in statistical significance.

A general limitation of this review relates to the pluralistic nature of TCQ. Both Tai Chi and Qigong are represented by many different styles (e.g. Chen, Yang, Wu or Sun styles of Tai Chi) and within each style there exist many training protocols (e.g. short-vs. long-forms), with some training regimens developed specifically for research purposes. This heterogeneity limits comparisons between, and generalizations across, studies [71]. Additionally, because of the limited number of studies available evaluating TCQ for PD, as well as the heterogeneity of PD populations evaluated across studies, we did not attempt to quantify or draw general conclusions of effectiveness based on specific styles or regimens. As results of additional larger-scale trials become available, future studies should consider individual patient-data meta-analyses that can evaluate the impact of training regimen while controlling for stage of PD progression and other relevant confounding factors [80].

There are a number of methodological limitations with our study. First, our study only included trials published in the English language. Future studies should include other languages for a better global estimate. Second, because of the small number of relatively small studies available to assess most outcomes, these findings should be considered suggestive, but not definitive. This concern is reinforced by limitations in methodological quality and by the heterogeneity of both TCQ interventions and controls. Third, we did not distinguish outcomes that were assessed on and off-medication, as has been done in prior reviews [7,14]. Fourth, there were too few studies to compare the effects of TCQ on subgroups of PD patients, such as early vs. later disease progression or tremor dominant vs. postural instability and gait difficulty. Future TCQ trials targeting or stratifying by specific PD subgroups would contribute significantly to the current evidence base. Finally, our conclusion that TCQ is 'likely to be safe for people with PD' warrants further research. As is the case in trials of many other non-

pharmacological therapies, AE reporting in studies of TCQ for PD is poor [62]. Nevertheless, findings of a comprehensive review of 153 Tai Chi studies found that, when conclusions were limited to studies with valid AE monitoring and reporting protocols, Tai Chi was found to be a safe exercise option [62].

5. Conclusions

RCT evidence supports a potential benefit of TCQ for improving multiple motor and non-motor outcomes for individuals with PD. Multiple studies and small-to-medium effect sizes support clinically meaningful effects in motor function, balance, and quality of life. More limited evidence also supports positive effects on fall risk and depression. However, all findings must be interpreted cautiously due to limitations in both the quantity and quality of available evidence. Additional large, rigorous trials are warranted to better characterize the effects of TCQ in PD and to guide selection of optimal doses and specific protocols for individuals with different PD subtypes and symptom burdens.

Author's role

Research project	A) conception	B) organization	C) execution
Statistical analysis	A) design	B) execution	C) review and critique
Manuscript	A) writing of the first draft	B) review and critique	

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Kamila Osypuk,	1B, 1C, 3B
G.P. Vergara-Diaz,	1A, 3B
Paolo Bonato,	1A, 3B
Jeffrey Hausdorff,	1A, 3B
Mike Fox,	1A, 3B
Lewis R. Sudarsky,	1A, 3B
Eric A. Macklin,	2A, 2C, 3B
Peter M. Wayne,	1A, 1B, 2A, 2C, 3A, 3B

Conflict of interest

Financial disclosure

Peter Wayne is the founder and sole owner of the Tree of Life Tai Chi Center. Peter 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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