

Original Article

Qigong Ameliorates Symptoms of Chronic Fatigue: A Pilot Uncontrolled Study

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Traditional Chinese Medicine practitioners consider that chronic fatigue reflects a disharmony and depletion in the supply of *qi* in the body. Qigong is one of the traditional complementary interventions used to strengthen *qi* through self-practice, and to manage the state of *qi* to prevent and cure disease. The aim of this study is to assess whether qigong could be used to manage the symptoms of chronic fatigue. Eighteen Caucasian, British female participants were recruited, taught a qigong routine during weekly classes over 6 months, and asked to practice it daily for 15 min. Participants completed the core set of the RAND Medical Outcomes Study questionnaire (RAND MOS) and a sleep diary during the 2-week baseline control period, and at 3 and 6 months following the start of the trial. The qigong intervention resulted in significant changes in sleep rate score and in the following subscales of the RAND MOS: SF36 Vitality, Sleep Problems, Social Activity, Social Activity Limitation due to Health, Health Distress, Mental Health Index and Psychological Well-being. Qigong seems to improve factors related to chronic fatigue such as sleep, pain, mental attitude and general mobility after 3 and 6 months. Qigong's positive effects indicate that it represents a potentially safe method of treatment for chronic fatigued patients. However, we cannot completely discount the possible influence of placebo effects, and more objective clinical measures are needed to reproduce our findings with long-term follow-up in a randomized, controlled study involving a larger number of subjects.

Keywords: chronic fatigue–sleep disturbance–qigong–quality of life

Introduction

Chronic fatigue syndrome (CFS) consists of a range of symptoms including physical and mental fatigue, headaches, sleep disturbances, concentration difficulties and muscle pain (1). In 2002 it was estimated that 240 000 people in the UK (~0.4%) were suffering from CFS, with the associated lost working hours, medication costs and medical support costs amounting to around £3.5 billion annually (2).

Common pharmacological treatments include the use of reduced nicotinamide, adenine dinucleotide (NADH) (3), sulbutiamine (4), fludrocortisone (5,6) and antidepressants (7,8), as well as certain pain and sleep medications. Because of the potential for adverse reactions to pharmacologic therapy in CFS patients, complementary and alternative interventions including massage (9), cognitive behavioral therapy (10–12) and osteopathy (13) are used to manage their symptoms. Traditional Chinese medicine considers chronic fatigue to reflect a disharmony and depletion in the supply of *qi*, with blockage, stagnation, imbalance or change in the pattern or organization of *qi* resulting in disease (14,15). Disruption to *qi* manifests in symptoms such as pain,

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fatigue and mood disturbances. Qigong is one of the traditional complementary interventions used to manage the deregulated state of *qi* to prevent and cure disease, to improve health and to strengthen *qi* by self-practice or by receiving *qi* from a practitioner.

Although neither *qi* itself nor the mechanism of its effects is understandable or explicable within any paradigm of modern medical science, its effects on the human body are apparent and some studies have been tried to find the underlying mechanism with laboratory experiments (16). Recent *in vitro* studies show that *qi* protected mitochondria from oxidative stress and calcium ions may play an important role in this protection (17). Another study suggest the possibility that *qi* may be beneficial for cancer patients because it suppresses cancer cell growth, and at the same time, it stimulates the patients immune functions (18,19).

Previous clinical research shows that qigong training significantly affects symptoms of asthma (20) and measurable parameters of bodily physiological processes, including levels of superoxide and growth hormones in elderly men (21), reductions in blood pressure and blood lipids in middle-aged adults with hypertension (22), improvement of immunological blood parameters (23) and cellular antigen immunity (24). Considering these effects, qigong can be applied to supportive care for symptoms of CFS such as vitality, sleep problems, social activity limitations due to health problems and psychological well-being. However, few clinical studies have examined the effects of qigong in chronically fatigued patients. Hence, this study aimed to determine whether qigong could help patients with chronic fatigue.

Participants and Methods

Study Design

An uncontrolled 6-month trial with three measurement periods (baseline, 3 and 6 months) was conducted (Fig. 1).

Participants

Recruitment of volunteers was from patients that general practitioners (GPs) or practice nurses had identified and referred as suffering from chronic fatigue. Participants were included if they had demonstrated symptoms of fatigue (tiredness with an inability to sleep, inability to concentrate, bodily pain, irritability and emotional problems) over a period of 3 months and could stand unaided for at least 10 min. No attempt was made to analyze the reason for their fatigue. All the participants were visiting a GP regularly and continued to do so for the trial's duration. The study was carried out within a nurse-led GP in Derby, UK between January and October 2004. This practice is unusual in being a

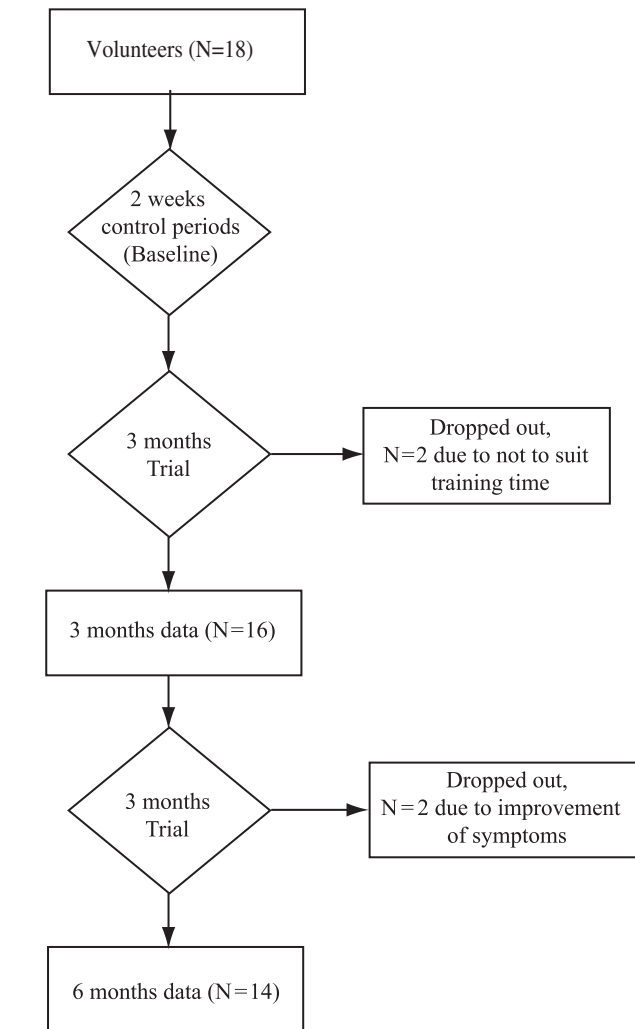


Figure 1. Diagram of study design showing the flow of participants.

(female only) nurse-led practice, and its nursing staff provided all recruitment, support and collected all data. Most patients were female and thus the volunteers recruited were also female. Their ages ranged from 25 to 55. No enquiry was made about whether they were pre- or post-menopausal or about any other factors such as social class, income or marital status because there was an insufficient number of participants to render any meaningful statistical analysis. The Southern Derbyshire Local Research Ethics Committee granted ethical approval (LREC REF: 0309/724). All participants agreed to participate and signed an informed consent form provided by this ethics committee.

Intervention Protocol

Over an initial 2-week period (Fig. 1), participants recorded pre-intervention outcome measures (i.e. sleep diaries and RAND MOS responses) so as to establish a stable baseline prior to the qigong intervention. During this period (and for the duration of the trial) they

continued to attend the GP practice and to take their prescribed medication.

Participants were invited to attend a weekly qigong session held at their local GP, when they were taught a 10-min qigong routine (Tai Yi Shen Gong) and were then asked to sit quietly for 5 min with their eyes closed while resting their hands on their knees. A certified qigong master who had trained in qigong for about 20 years led the classes. He had studied qigong and Tai Chi in China at Anhui and Beijing TCM Universities between 1992 and 2006. Tai Yi Shen Gong is a qigong routine comprising sequential arm movements performed slowly in time with steady breathing, such as reaching downward, forward and above the head. Participants repeated the routine at weekly classes over a 3-month period. Following this initial phase, they then practiced the qigong routine daily at home in addition to attending the weekly qigong class for a further 3-month period.

Adverse Events

The instructor monitored participants before, during and after each qigong class for any sign of adverse events.

Outcome Measures

RAND Medical Outcomes Study Questionnaire

Participants completed the core set of the RAND Medical Outcomes Study questionnaire (RAND MOS) (25) during the 2-week control period, and 3 and 6 months following the start of the trial. They completed every question on the questionnaire, and every index therein was calculated. The core set of the RAND MOS is comprised of 116 questions that measured participants' functioning and well-being in 37 categories of concepts related to the following (25): Physical health (e.g. physical functioning, satisfaction with physical ability, mobility, pain effect, pain severity and role limitations due to physical health), Mental health (e.g. psychological distress/anxiety, depression, psychological well-being, positive effect, feeling of belonging, cognitive functioning and role limitations due to emotional problem), and General health (e.g. energy/fatigue, sleep problems, psychophysiological symptoms, social functioning, role functioning—unable to work, role functioning—unable to do housework, current health perceptions and health distress). The instrument's subscales were scored as summated rating scales on a 0–100 scale, where higher scores indicated better health. The validity and reliability of RAND MOS were examined in detail in a previous study (26).

Sleep Diary

Participants were also asked to keep a subjective sleep diary, which is generally used at UK National Health

Service hospitals (27). The following variables were computed from the diary entries: (i) interruptions to sleep, defined as number of times waking up at night; (ii) total sleep time, defined as number of hours actually slept out of the number in bed, and quality of sleep, anchored by: 1—very disrupted, 2—restless, 3—average, 4—sound and 5—very sound (subjectively rated by participants). The diaries were completed for an initial 2 weeks before the trial commenced and then for 1 week each month for the duration of the trial. Participants were given detailed instructions about how to complete the various measures in the diary.

Statistical Analysis

Data from the RAND MOS were converted into RAND indices using recommended procedures, and sleep-diary data were converted into means for each week. Data were analyzed using the intent-to-treat principle. All outcomes were compared using a non-parametric Friedman's test across treatment time (control, 3 and 6 months) for each subscale. Follow-up tests were performed with multiple comparisons of pairs of times (control versus 3 months, control versus 6 months and 3 months versus 6 months) using a Wilcoxon-singed rank test. The SPSS version 11.0 (SPSS, Inc., Chicago, IL, USA) software package was used for all statistical calculations.

Results

Attrition and Compliance

Details of attrition are shown in Fig. 1. Eighteen patients were approached and all agreed to participate. The trial commenced with 18 female volunteers (mean age \pm SD, 40.7 ± 9.5 years). Two of them withdrew after the first qigong session because the study's timing did not suit their personal circumstances. The remaining 16 completed the first 3-month phase of the study. In the second 3-month phase, two more participants stopped attending because of improvement in their condition (although they continued to practice qigong at home); hence 14 participants completed the study. After the initial period when two participants stopped attending, all of the remaining participants continued practicing qigong for the 6-month period, and no negative or adverse events were reported. The total attendance rate was 91.88% at the weekly sessions.

Outcome Measures: Qigong Improves Factors Related to Chronic Fatigue

Figure 2 shows the mean \pm SD values for subscales of the sleep parameters from the RAND MOS (Fig. 2A) and the sleep diary (Fig. 2B). The qigong intervention

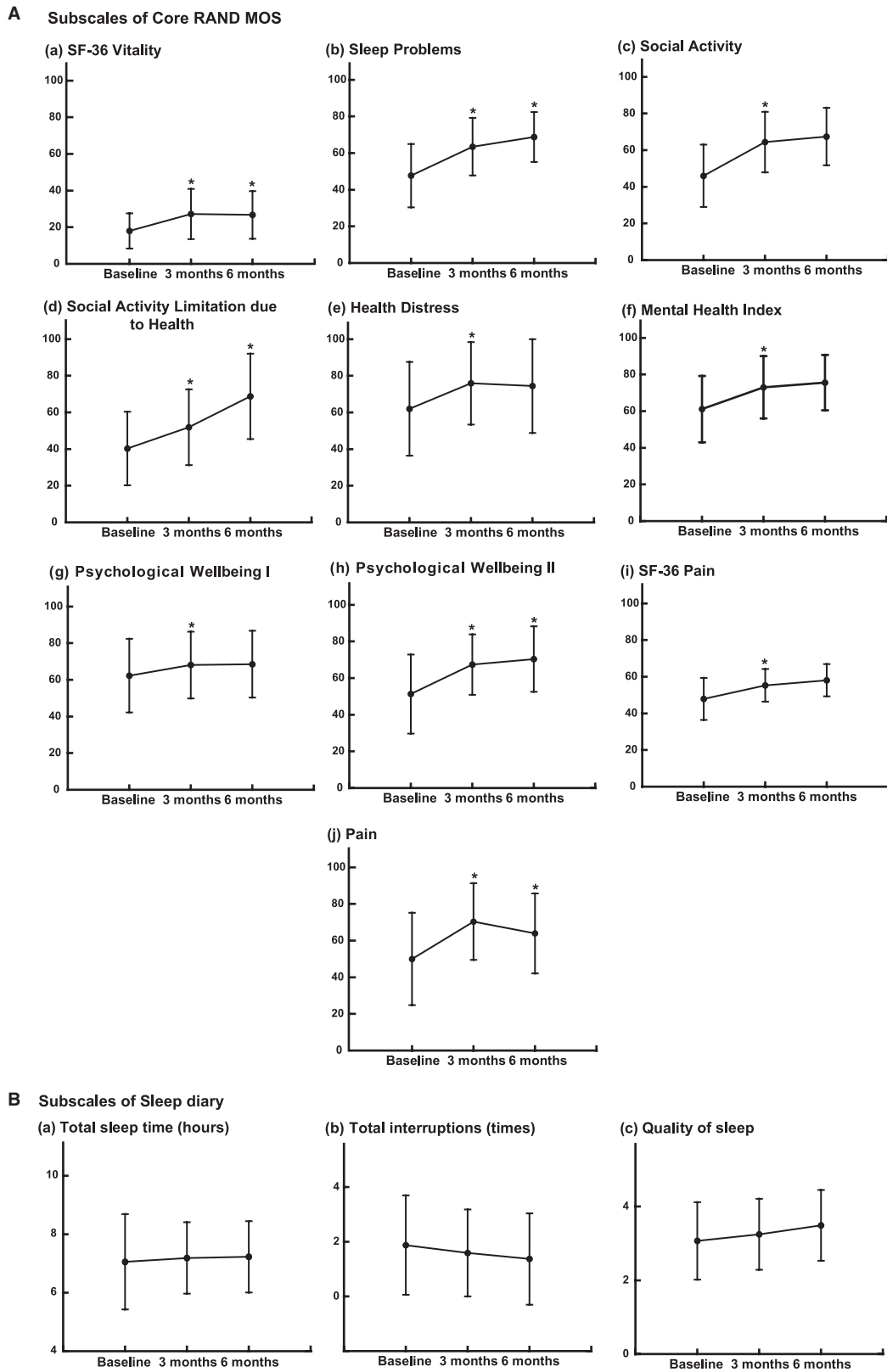


Figure 2. Mean changes of subscales of (A) RAND Medical Outcomes Study questionnaire (The subscales of the instrument were scored as summated rating scales on a 0–100 scale where higher scores indicated better health. All subscales were statistically significant ($P < 0.05$) by Friedman’s test. * $P < 0.05$ by Wilcoxon signed rank test compared with control period) and (B) Sleep diary. All results are presented as means \pm SD. The numbers of subject were 18, 16 and 14 for baseline, 3 months and 6 months, respectively (Quality of sleep were scored on a 0–5 scores where higher score indicate a good sleep. $P < 0.05$ by Friedman’s test).

resulted in significant changes in the following subscales of the RAND MOS: SF36 Vitality, Sleep Problems, Social Activity, Social Activity Limitation due to Health, Health Distress, Mental Health Index, Psychological Well-being and Psychological Well-being II; whereas Effects of Pain and SF-36 Pain did not change significantly. Significant improvement was also noted in the Sleep Rate score ($P < 0.005$), but not in the Total Sleep or Total Interruptions scores.

The participants differed significantly from the control period after 3 months in SF36 Vitality, Sleep Problems, Social Activity, Social Activity Limitation due to Health, Health Distress, Mental Health Index, Psychological Well-being I, Psychological Well-being II, Pain and SF-36 Pain (all $P < 0.05$).

Significant differences compared with the control period were also noted after 6 months in SF36 Vitality, Sleep Problems, Social Activity Limitation due to Health, Mental Health Index, Psychological Well-being I, Psychological Well-being II and Pain (P4) ($P < 0.05$). However, there were no significant differences in sleep parameters compared with the control period.

Discussion

Epidemiological evidence suggests that both a single session of qigong (acute effects) and prolonged qigong over several months (chronic effects) can produce significant and positive changes in psychological characteristics and in the neuroendocrine and immune systems (23,28–31). Our pilot study suggests that regular qigong training reduces pain and improves sleep, vitality and physical functioning in patients with chronic fatigue. This finding is consistent with results from previous studies. Astin *et al.* (32) reported that 8 weeks of qigong combined with mind–body intervention reduced multiple sclerosis patients' pain level. Another study also reported qigong's beneficial effects on general health measured with SF-36 in patients with muscular dystrophy (33).

Assuming that qigong is a potentially useful treatment option for chronic fatigue related symptoms, its possible mechanism of action may be of interest. Possible mechanisms include increases in oxygen and decreases in carbon dioxide concentrations in the blood, which may enhance the removal of pain-inducing substances (such as metabolic waste products) from the tissues (34). Qigong may also enhance the circulation of pain-killing substances such as endorphins and other agents that control pain (35). In general, qigong, which consists of steady breathing, slow bodily movements and sitting quietly, affects the muscular systems, resulting in muscular adaptation, and ultimately leads to increased muscle strength and relaxation, if performed regularly. These effects also improve physical and psychological functioning,

and therefore beneficially affect fatigue, sleep disturbances, concentration difficulties and muscle pain.

However, this study has several possible limitations. One potential limitation is its use of a small sample and the absence of an active control group. Moreover, other objective outcome measures that are clinically related to symptoms, including the level of muscle strength, other physical activities and instrument-related fatigue and sleep disturbances should be used in addition to the measurements employed in this study. Furthermore, because all the participants were women, we cannot assume that the results are representative across the whole population.

In summary, the present results suggest that qigong can improve factors related to CFS such as sleep, pain, mental attitude and general mobility after 3 and 6 months. Qigong's positive effects indicate that it represents a potentially safe method of treating chronic fatigued patients. However, we cannot completely discount the possibility that the placebo effect during the intervention caused improvements in symptoms related to chronic fatigue. The reproducibility of our findings should also be tested using objective measurements and long-term follow-up in a randomized, controlled study involving a larger number of subjects.

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