

Randomised, controlled, multicenter, pilot study comparing qigong and back school for elderly patients with chronic neck pain

Submission date
09/05/2006

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
23/05/2006

Overall study status
Completed

☐ Statistical analysis plan

☐ Results

Last Edited
06/02/2008

Condition category
Injury, Occupational Diseases, Poisoning

☐ Individual participant data

☐ Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

Acronym

QIBANE

Study objectives

Primary hypothesis: there is a significant difference between the qigong and the waiting list groups for the primary outcome (visual analogue scale [VAS]) after 3 months.

Secondary hypothesis: there is a significant difference between the qigong and the back school groups for the primary outcome (VAS) after 3 months.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Yes, date of approval: 23/02/06; reference number: EA1/265/05

Study design

Multicenter, three-armed, randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Chronic neck pain

Interventions

Patients are randomized into the qigong and back school exercise groups (each 24 sessions of 45 min over a period of three months)

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Pain measured on the VAS (visual analogue scale) after three months

Key secondary outcome(s))

1. Neck pain (Neck Pain and Disability Scale, Wheeler)
2. Depression (Depression Scale, Hautzinger)
3. Quality of life (Short-form Questionnaire-36 [SF-36], Bullinger)
4. Frequency of falls and anxiety to fall
5. Sleep quality and satisfaction
6. Overall treatment effect (OTE) (Wiklund)

Completion date

31/10/2006

Eligibility

Key inclusion criteria

1. Age ≥ 55 years of age
2. Chronic neck pain (duration > 6 months)
3. Average pain during the last 7 days before randomisation ≥ 20 mm on 100 mm VAS
4. Informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Senior

Sex

All

Key exclusion criteria

1. Serious acute or chronic disease which would not allow the exercises to be performed
2. Planned start of physiotherapy for chronic neck pain during study participation
3. Participation in another study during the last 6 months before study entry

Date of first enrolment

09/03/2006

Date of final enrolment

31/10/2006

Locations

Countries of recruitment

Germany

Study participating centre

Institute for Social Medicine

Berlin

Germany

10098

Sponsor information

Organisation

Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin) (Germany)

ROR

<https://ror.org/001w7jn25>

Funder(s)

Funder type

University/education

Funder Name

Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin), Institute for Social Medicine, University Medical Center (Germany)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration