Physical exercise for rehabilitation of neck /shoulder muscle pain: how little is enough?

Prospectively registered Submission date Recruitment status 28/07/2009 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 18/09/2009 Completed [X] Results [] Individual participant data **Last Edited** Condition category 20/11/2012 Musculoskeletal Diseases

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N/A

Study information

Scientific Title

Dose-response of specific strength training for rehabilitation of neck/shoulder muscle pain: a randomised single-blind controlled trial

Study objectives

Participants randomised to specific strength training for 5×2 minutes per week and 5×12 minutes per week will report better relief of neck/shoulder muscle pain compared with a control group.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Local Ethical Committee of Copenhagen and Frederiksberg, Denmark, approved in November 2008 (ref: HC-2008-103). Approval of supplementary protocol: 9th March 2009.

Study design

Randomised single-blind controlled intervention trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Musculoskeletal disorders

Interventions

The intervention will last 10 weeks. Employees randomised to specific strength training will be offered 5 x 2 minutes per week or 5 x 12 minutes per week for 10 weeks of specific strength training for the neck/shoulder muscles. Participants randomised to the control group will receive information on various aspects of general health.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

- 1. Subjective pain, measured weekly throughout the 10 week intervention period
- 2. Clinical findings, measured twice; before and after the intervention period
- 3. Muscle strength, measured twice; before and after the intervention period

Secondary outcome measures

- 1. Other subjective health complaints, measured twice; before and after the intervention period
- 2. Compliance, measured weekly throughout the 10 week intervention period

Overall study start date

01/08/2009

Completion date

15/12/2009

Eligibility

Key inclusion criteria

- 1. Generally healthy workers
- 2. Aged 25 65 years, either sex
- 3. Palpable tenderness of the neck/shoulder muscles
- 4. An anamnestic history of neck/shoulder pain for at least 30 days during last year
- 5. Reported neck/shoulder pain intensity of at least 2 on a scale of 0 10

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

180

Key exclusion criteria

- 1. Trauma
- 2. Pregnancy
- 3. Life threatening diseases
- 4. Blood pressure above 160/100 mmHg
- 5. Other known serious disorders, e.g. fibromyalgia or rheumatoid arthritis

Date of first enrolment

01/08/2009

Date of final enrolment

15/12/2009

Locations

Countries of recruitment

Denmark

Study participating centre Lersø Parkalle 105

Copenhagen Denmark 2100

Sponsor information

Organisation

The National Research Centre for the Working Environment (Denmark)

Sponsor details

Lersø Parkalle 105 Copenhagen Denmark 2100

Sponsor type

Research organisation

Website

http://www.arbejdsmiljoforskning.dk

ROR

https://ror.org/03f61zm76

Funder(s)

Funder type

Research organisation

Funder Name

The National Research Centre for the Working Environment (Denmark)

Funder Name

The Danish Rheumatism Association (Denmark) (ref: R68-A993)

Funder Name

Hygenic Corporation (USA)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/09/2012		Yes	No