

Individualising exercise for knee pain: developing an evidence-based impairment-targeted intervention

Submission date 24/06/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 24/06/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 01/02/2016	Condition category Signs and Symptoms	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

5142

Study information

Scientific Title

Individualising exercise for knee pain: developing an evidence-based impairment-targeted intervention

Acronym

TargET-Knee-Pain

Study objectives

The overall aim of this study is to investigate the principle of impairment-targeted exercises as treatments for older adults with knee pain. The study has a single-group design. It has the following objectives:

Objective 1:

To test the proof-of-principle that simple home-based impairment-targeted exercises can improve those impairments in older adults with knee pain.

Objective 2:

To investigate whether improvements in these impairments are reflected in improvements in self-reports of physical function.

Objective 3:

To assess the feasibility of this intervention approach and its acceptability to patients by qualitative means.

Ethics approval required

Old ethics approval format

Ethics approval(s)

MREC approved on the 7th November 2008 (ref: 08/H1202/179)

Study design

Single centre non-randomised interventional treatment trial

Primary study design

Interventional

Secondary study design

Non randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Topic: Primary Care Research Network for England, Musculoskeletal; Subtopic: Not Assigned, Musculoskeletal (all Subtopics); Disease: Musculoskeletal, All Diseases

Interventions

This will be a single-centre study. Participants will be invited to participate in the current study at the end of their visit for final clinical assessment before the termination of the CAS-K study.

Exercise programme: consists of an individualised home-based impairment-targeted exercise programme for 12 weeks (fortnightly home visits alternating with fortnightly telephone calls) and involves daily self-directed formal exercises.

Recruitment will last for the duration of the final clinical assessment clinics, which are planned to take place between January and December 2009. Duration of follow-up will be 12 weeks.

Study entry: registration only

Intervention Type

Other

Phase

Phase IV

Primary outcome measure

Degree of knee flexion, measured with a continuous scale outcome

Secondary outcome measures

1. Isometric quadriceps strength at 90° flexion, measured with a continuous scale outcome
2. The four-test balance scale, measured with an ordinal scale (range 0 - 5), including feet together stand, semi-tandem stand, tandem stand and one-leg stand

Overall study start date

05/05/2009

Completion date

09/02/2010

Eligibility

Key inclusion criteria

1. Aged 56 years and older, either sex
2. A history of painful osteoarthritis of the knees
3. One or more of the target impairments
4. Recruited from the longitudinal population-based Clinical Assessment Study of the Knee (CAS-K)

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

Planned sample size: 80; UK sample size: 80

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

05/05/2009

Date of final enrolment

09/02/2010

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

Primary Care Musculoskeletal Research Centre

Newcastle

United Kingdom

ST5 5BG

Sponsor information**Organisation**

Keele University (UK)

Sponsor details

Keele

Newcastle

England

United Kingdom

ST5 5BG

Sponsor type

University/education

Website

<http://www.keele.ac.uk/>

ROR

<https://ror.org/00340yn33>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) programme (ref: PB-PG-0107-10612)

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	07/01/2011		Yes	No
Results article	results	29/01/2016		Yes	No