

Acupuncture in a group setting for chronic knee pain: ScrutiKnee

Submission date 18/05/2012	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 18/05/2012	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 20/10/2017	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Background and study aims

Osteoarthritis (OA) is a common cause of pain and disability, affecting one in five people aged over 50 in the UK. Previous studies show that acupuncture (a form of ancient Chinese medicine in which fine needles are inserted into the skin at certain points on the body) is effective in treating pain in people with mild to moderate OA of the knee. Studies also show that every person with osteoarthritis of the knee should be offered exercise and advice. What we don't yet know is whether exercise, advice and acupuncture can help people with severely painful knee osteoarthritis, who may otherwise be considered for joint replacement surgery. Another issue is cost. In some clinics in the NHS, acupuncture is being given to a group of patients at the same time to reduce its cost. There has been no research on how well acupuncture works when it is given to people who are part of a group, nor whether or not this reduces the cost of treatment. ScrutiKnee is a initial small study. So it is not designed to see if exercise, advice and acupuncture are effective, but it is designed to provide information for a larger study which will test whether these treatments are effective and cost-effective. ScrutiKnee will find out whether patients are interested in participating, and how we can best measure the changes we want to measure, such as pain and the use of health services.

Who can participate?

Men and women over the age of 45 can take part in ScrutiKnee, if they have had painful OA knees for at least a year, and if the pain has troubled them for most days in the previous month. The pain must be of a particular severity, as scored by the common scale, the Oxford Knee Score.

What does the study involve?

Patients who might be eligible are identified from their general practitioner's (GPs) databases and sent a letter. If patients are interested in taking part and seem eligible, they are phoned and sent an information leaflet about the study. Those still interested are invited to attend the research centre and consented to have their knees examined by a research nurse. Participants are selected at random to receive one of three treatments. All participants receive a booklet on self-management advice and exercises which is considered standardised care. The three treatment groups are:

1. The standardised care on its own
2. Standardised care plus acupuncture as part of a group with other participants, and

3. Standardised care plus acupuncture one-to-one with the practitioner.

The acupuncture clinic is in Plymouth (Teaching) Primary Care Trust. The acupuncture is provided by a Chartered Physiotherapist trained in acupuncture. A course of acupuncture consists of six to ten treatments over 12 weeks the treatment stops at six weeks if there is no improvement in the symptoms.

Participants are asked at the start and at the end of the treatment period to complete questionnaires about the impact their knee pain has on their life.

In addition, since this is a initial study arranged to help design the main study, participants are asked to write comments on their experiences in the study. Some participants are also invited for interview to report their experiences in greater detail.

What are the possible benefits and risks of participating?

Stuides shows that advice and exercise can produce long-term benefit for people with OA knee. It is not clear whether acupuncture provides any additional pain relief for people with severely painful OA knee. However there could be benefits to future patients by showing whether or not acupuncture is worth providing in the NHS for people with severely painful OA knee. The risks of taking part are minimal. Acupuncture is a very safe treatment when given by properly trained clinicians. Occasionally acupuncture can make people feel nauseous or faint or experience a temporary increase in pain either during or after treatment. Participants are warned of these potential side-effects before consenting to have acupuncture.

Where is the study run from?

The study is run from the Peninsula Medical School, University of Plymouth, and the Peninsula Clinical Trials Unit (PenCTU) based in Plymouth.

When is the study starting and how long is it expected to run?

February 2012 to September 2012

Who is funding the study?

National Institute for Health Research, Research for Patient Benefit (NIHR-RfPB) (UK)

Who is the main contact?

Dr Adrian White

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Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

11589

Study information

Scientific Title

Pilot study of acupuncture in a group setting for chronic knee pain: ScrutiKnee

Study objectives

ScrutiKnee is a pilot randomised controlled trial that will provide information to help develop a subsequent trial that will evaluate the effectiveness and cost-effectiveness of group acupuncture in severe knee osteoarthritis. ScrutiKnee is a feasibility study designed particularly to test recruitment rate and resources required, and to establish the best method to measure the use of health service resources.

More details can be found at <http://public.ukcrn.org.uk/Search/StudyDetail.aspx?StudyID=11589>

Ethics approval required

Old ethics approval format

Ethics approval(s)

Cornwall and Plymouth Ethics Committee First MREC approval date 24/11/2011, ref: 11/SW/0277

Study design

Randomised; Interventional; Design type: Treatment

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Osteoarthritis

Interventions

Potentially eligible patients will be identified from GP registers and sent an invitation to participate. Those who respond will be screened by telephone and sent a participant information leaflet. Those still interested will be invited to attend the research centre. After giving consent and providing baseline information, all participants will receive a booklet on self-management advice and exercises, which is considered standardised care. Participants will be randomised to three groups:

1. Standardised care alone

2. Standardised care plus group acupuncture, or
3. Standardised care plus individual acupuncture.

Acupuncture: Group or individual acupuncture using semi-standardised formula, exercise and advice leaflet, purpose-designed leaflet for patients with moderate to severe knee pain. A course of acupuncture consists of six to ten treatments over 12 weeks.

Knee symptoms will be assessed before and after the course of treatment using common outcome measures. Participants will also be asked to write comments on their experiences in the study, and some participants will also be invited for interview to report their experiences in greater detail to provide information that will optimise the design of the subsequent main study. The trial duration for each participant is 14 weeks.

Follow Up Length: 3 month(s); Study Entry : Single Randomisation only

Intervention Type

Procedure/Surgery

Primary outcome(s)

Western Ontario and McMaster Universities Arthritis Index (WOMAC) at baseline and 3 months

Key secondary outcome(s)

1. Oxford Knee Score at baseline and 3 months
2. Use of health and social service resources

Completion date

28/09/2012

Eligibility

Key inclusion criteria

1. Pain in one or both knees for at least a year, troublesome for most days in the previous month
2. Meets the clinical American College of Rheumatology (ACR) criteria for the diagnosis of osteoarthritis (at least three of the following):
 - 2.1. Age at least 45 years
 - 2.2. Stiffness, if present, lasts \geq 30 minutes
 - 2.3. Crepitus
 - 2.4. Bony tenderness
 - 2.5. Bony enlargement
 - 2.6. No palpable warmth
3. Not had Knee Joint Replacement (KJR) in index knee, nor currently referred for surgery
4. Oxford Knee Score (OKS) \geq 28
5. Able to travel to attend clinic
6. Able to complete outcome measures
7. Can be contacted by telephone; Target Gender: Male & Female ; Lower Age Limit 45 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Steroid injection or acupuncture to knee in the last two months
2. Hyaluronic acid injection, arthroscopy or serious injury to knee in last six months
3. Had Knee Joint Replacement (KJR) in index knee, or currently referred for consideration of surgery of the index knee
4. Clinical diagnosis of severe osteoarthritis of the ipsilateral hip
5. Not willing or able to provide informed consent
6. Bleeding disorder undiagnosed or severe
7. Concurrent medical conditions which would impair participation
8. Currently participating in any other interventional clinical trial

Date of first enrolment

22/02/2012

Date of final enrolment

28/09/2012

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

I T T C Building

Plymouth

United Kingdom

PL6 8BX

Sponsor information**Organisation**

NHS Plymouth (UK)

Funder(s)

Funder type
Government

Funder Name
National Institute for Health Research (NIHR) - Research for Patient Benefit (RFPB) (UK)

Results and Publications

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	16/02/2016		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Study website	Study website	11/11/2025	11/11/2025	No	Yes