A study of acupuncture, physiotherapy or nonintervention in management of painful (OA) knee

Submission date 12/09/2003	Recruitment status Stopped	 Prospectively registered Protocol
Registration date 12/09/2003	Overall study status Stopped	 Statistical analysis plan Results
Last Edited 25/10/2011	Condition category Musculoskeletal Diseases	 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 Dr Jeremy Mc Nally

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0199114670

Study information

Scientific Title

Study objectives

To evaluate the effectiveness of acupuncture, as compared with physiotherapy and no treatment in management of painful knee.

Ethics approval required Old ethics approval format

Ethics approval(s) Not provided at time of registration

Study design Randomised controlled trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Not specified

Study type(s) Treatment

Participant information sheet

Health condition(s) or problem(s) studied Musculoskeletal Diseases: Osteoarthritis (OA)

Interventions

Randomly assigned to three groups

This trial was stopped due to a lack of funding.

Intervention Type Other

Phase Not Specified

Primary outcome measure

1. Intensity of pain by Visual Analogue Scale

2. HAD

3. Western Ontario MacMaster Questionnaire (WOMAC)

4. Timed 50 yard walk

5. Clinician's assessment of knee pain

Secondary outcome measures Not provided at time of registration

Overall study start date 01/10/2001

Completion date 31/12/2006

Reason abandoned (if study stopped) Lack of funding

Eligibility

Key inclusion criteria

 150 patients fulfilling American College of Rheumatology (ACR) criteria for OA of knee, unilateral or bilateral OA knee pain for 3 months
 Never had acupuncture or physiotherapy for knee pain before

Participant type(s) Patient

Age group Adult

Sex Both

Target number of participants 150

Key exclusion criteria Not provided at time of registration

Date of first enrolment 01/10/2001

Date of final enrolment 31/12/2006

Locations

Countries of recruitment England

United Kingdom

Study participating centre Department of Rheumatology Reading United Kingdom RG30 1AG

Sponsor information

Organisation Department of Health (UK)

Sponsor details Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type Government

Website http://www.doh.gov.uk

Funder(s)

Funder type Government

Funder Name Royal Berkshire and Battle Hospitals NHS Trust (UK)

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