Ultrasound-guided versus palpation-guided steroid injection for plantar fascitis

Submission date	Recruitment status	Prospectively registered
30/09/2004	No longer recruiting	∐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2004	Completed	☐ Results
Last Edited	Condition category	Individual participant data
14/06/2017	Musculoskeletal Diseases	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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0188130055

Study information

Scientific Title

Ultrasound-guided versus palpation-guided steroid injection for plantar fascitis: a randomised prospective trial

Study objectives

The aim of this study is to determine whether ultrasound-guided injection gives superior results to palpation-guided steroid injection in plantar fasciitis.

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)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Musculoskeletal Diseases: Plantar fasciitis

Interventions

Comparison of steroid injection for the treatment of plantar fasciitis without ultrasound scan guidance and with ultrasound scan guidance.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2003

Completion date

01/12/2006

Eligibility

Key inclusion criteria

Patients with clinical features of plantar fascitis who present in the orthopaedic clinic will be approached to participate in the study

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

40 patients

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/08/2003

Date of final enrolment

01/12/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Lancashire Teaching Hospitals NHS Trust

Preston United Kingdom PR2 9HT

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Lancashire Teaching 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