ISRCTN84362147 https://doi.org/10.1186/ISRCTN84362147

Quantifying the effects of early mobilisation and loading for Achilles tendon rupture

Submission date	Recruitment status
12/09/2003	No longer recruiting
Registration date	**Overall study status**
12/09/2003	Completed
Last Edited	**Condition category**
16/09/2014	Musculoskeletal Diseases

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] 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
N0544093625

Study information

Scientific Title

Study objectives
Quantifying the effects of early mobilisation and loading for Achilles tendon rupture.

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: Achilles tendon rupture

Interventions
1. Early loading in new orthosis
2. Traditional plaster cast treatment

A full quantitative assessment of the effects of early loading and mobilisation on Achilles tendon healing. The study is part of a three-centre randomised controlled trial comparing early loading in a new orthosis with traditional treatment in a plaster cast. Patients will be assessed with clinical, radiological, histological, anthropometric and questionnaire follow-up.

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
27/02/2001

Completion date
20/01/2004

Eligibility

Key inclusion criteria
48 subjects (PROJ 01/02/2001).

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
48

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
27/02/2001

Date of final enrolment
20/01/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Warwick Medical School
Coventry
United Kingdom
CV2 2DX

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
Cambridge Consortium - Addenbrooke's (UK)

Funder Name
British Medical Association (UK)

Alternative Name(s)
BMA

Funding Body Type
Private sector organisation

Funding Body Subtype
Associations and societies (private and public)

Location
United Kingdom

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	01/01/2006		Yes	No