Function versus position of the wrist following distal radial fracture

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
01/05/2015	Injury, Occupational Diseases, Poisoning	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 N0244124359

Study information

Scientific Title

Function versus position of the wrist following distal radial fracture

Study objectives

- 1. To investigate whether the end result following k-wire fixation of the distal radius changes depending upon the position of the hand.
- 2. To establish whether time to recovery differs significantly depending upon the position of the hand.

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

Injury, Occupational Diseases, Poisoning: Radial fracture

Interventions

Patients with distal radial fractures undergoing reduction and k-wire fixation are randomised to dorsi or palmar flexion in plaster of Paris (POP). Upon removal of POP, grip test is measured using a Jamar Dynamometer. At 8 weeks, grip strength is measured again and the test is repeated at 17 weeks.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1. Grip strength
- 2. Range of motion
- 3. Radiological outcomes

Secondary outcome measures

Not provided at time of registration

Overall study start date

06/03/2003

Completion date

31/12/2004

Eligibility

Key inclusion criteria

Consecutive patients with displaced distal radial fracture undergoing reduction and k-wire fixation.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

- 1. Patients with severe rheumatoid arthritis
- 2. Previous articular surgery
- 3. Previous injury to the same wrist
- 4. Less than 20°-30° flexion or extension after k-wire fixation

Date of first enrolment

06/03/2003

Date of final enrolment

31/12/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Stepping Hill Hospital Stockport

United Kingdom SK2 7JE

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

Government

Funder Name

Stockport 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 summaryNot provided at time of registration