Suspected scaphoid fracture management Plaster of Paris versus Splint

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
08/09/2015	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

Mr Habib Dardouri

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers N0300129102

Study information

Scientific Title

Suspected scaphoid fracture management Plaster of Paris versus Splint

Study objectives

Is the Futura Splint an effective way of treating suspected scaphoid fracture?

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

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Injury, Occupational Diseases, Poisoning: Scaphoid fracture

Interventions

Patients are randomised to receive: plaster of paris or splint

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Pain, cost and patient comfort

Secondary outcome measures

Not provided at time of registration

Overall study start date

Completion date

01/04/2004

Eligibility

Key inclusion criteria

Patients with suspected scaphoid fracture aged 16-70 years.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/08/2002

Date of final enrolment

01/04/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Blackburn Royal Infirmary

Blackburn United Kingdom BB2 3LR

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

East Lancashire 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