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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
Accident & Emergency Department
Blackburn Royal Infirmary
Bolton Road
Blackburn
United Kingdom
BB2 3LR

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

01/08/2002

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