Comparison of Internal Fixation of Metacarpal and Phalangeal Fractures with Metal or Bioabsorbable Implants

Recruitment status	Prospectively registered
Stopped	☐ Protocol
Overall study status	Statistical analysis plan
Stopped	Results
Condition category	Individual participant data
Injury, Occupational Diseases, Poisoning	Record updated in last year
	Overall study status Stopped Condition category

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

Protocol serial number N0295132798

Study information

Scientific Title

Study objectives

Internal fixation of metacarpal and phalangeal fractures with bio-absorbable implants negates the need for removal of the implant, as with metallic implants, at a second operation with benefit to the patient and financial implications.

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Injury, Occupational Diseases, Poisoning: Fractures

Interventions

Please note that as of 21/06/10 it has been confirmed that this trial was stopped due to lack of staff and resources.

Prospective Randomised Controlled Trial. Metal vs bioabsorbable implants.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

- 1. Total active range of motion (TAM) of affected digit
- 2. Grip strength measured with Jamar instrument
- 3. Complication rate
- 4. Fracture union rate
- 5. Implant removal rate
- 6. Financial cost of each treatment group

Key secondary outcome(s))

Not provided at time of registration

Completion date

30/10/2004

Reason abandoned (if study stopped)

Lack of staff/facilities/resources

Eligibility

Key inclusion criteria

50 patients with metacarpal and phalangeal fractures. Sample size will take 12-18 months to recruit at current admission rates for the fractures being studied.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/03/2004

Date of final enrolment

30/10/2004

Locations

Countries of recruitment

United Kingdom

England

Study participating centre c/o Secretary to Professor Griffin

Coventry United Kingdom CV1 4FH

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

University Hospitals Coventry and Warwickshire NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration