Distal forearm torus fractures - do they need a splint?

| Submission date | Recruitment status | Prospectively registered |
|-------------------|--|---|
| 28/09/2007 | No longer recruiting | ☐ Protocol |
| Registration date | Overall study status | Statistical analysis plan |
| 28/09/2007 | Completed | Results |
| Last Edited | Condition category | Individual participant data |
| 19/10/2011 | 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

Dr Michelle Jacobs

Contact details

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Additional identifiers

Protocol serial number N0143185333

Study information

Scientific Title

Study objectives

To demonstrate that a torus fracture, treated with no splintage at all, is not significantly more painful than one treated with splintage and allows the child to start using the limb sooner.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Pilot Randomised Controlled Trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Injury, Occupational Diseases, Poisoning: Fractures

Interventions

A pilot study recruiting for 1 year, using patients randomised into two groups after x-ray diagnosis. Pain, satisfaction and treatment evaluation will be conducted at 1 week using patient and carer responses. At 4 weeks, patient's carer will do a similar evaluation again.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

- 1. Difference in pain levels at weeks 1 and 4 between the 2 groups
- 2. Time to return to normal use of affected limb with and without splintage
- 3. Patient and carer satisfaction level in the 2 groups.

Key secondary outcome(s))

Not provided at time of registration

Completion date

08/08/2007

Eligibility

Key inclusion criteria

All children under 17 presenting with a torus fracture of the distal end of the radius and or ulna.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Upper age limit

17 years

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

08/08/2006

Date of final enrolment

08/08/2007

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

A&E

Watford United Kingdom WD18 0HB

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Funder(s)

Funder type

Government

Funder Name

Hertfordshire Hospitals Research and Development Consortium (UK)

Funder Name

West Hertfordshire Hospitals NHS Trust

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration