ISRCTN45883421 https://doi.org/10.1186/ISRCTN45883421

Colles Fracture Brace prospective randomised study

Submission date	Recruitment status	[ ] Prospectively registered
27/10/2009	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
07/12/2009	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
02/10/2017	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
07/S0801/116

Study information

Scientific Title
Colles Fracture Brace prospective randomised study

Acronym
CFB Study

Study objectives
Treatment by a fracture brace will have a better outcome than standard care.

Ethics approval required
Old ethics approval format

Ethics approval(s)
North of Scotland Research Ethics Committee, 26/02/2008, ref: 07/S0801/116

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 to request a patient information sheet

Health condition(s) or problem(s) studied
Colles fracture

Interventions
Application of plaster of paris (POP) cast, brace or surgery.

Total duration of follow-up: 2 years

Intervention Type
Procedure/Surgery

Primary outcome measure
Clinical and radiological outcome at 6 months

Secondary outcome measures
Clinical outcome at 2 years

Overall study start date
16/11/2009

Completion date
16/05/2012

Eligibility

Key inclusion criteria
1. Both males and females, aged 18 and over
2. Colles fracture requiring treatment

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Both

Target number of participants
340

Key exclusion criteria
1. Compared fracture
2. Previous fracture
3. Unable to give informed consent

Date of first enrolment
16/11/2009

Date of final enrolment
16/05/2012

Locations

Countries of recruitment
Scotland

United Kingdom

Study participating centre
Woodend Hospital
Aberdeen

United Kingdom
AB15 6XS

Sponsor information

Organisation
NHS Grampian (UK)

Sponsor details
Research and Development
Foresterhill House Annexe
Foresterhill
Aberdeen
Scotland
United Kingdom
AB25 2ZB

Sponsor type
Hospital/treatment centre

Website
http://www.nhsgrampian.org

ROR
https://ror.org/00ma0mg56

Funder(s)

Funder type
Other

Funder Name
Local funding from endowments (derek.angus@nhs.net)

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