Prospective randomised trial of treatment of fingertip injuries

| 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 |
| 19/02/2014 | Injury, Occupational Diseases, Poisoning | Record updated in last year |

Plain English Summary

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Jo Skillman

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0143136995

Study information

Scientific Title

Study hypothesis

Is it better to shorten or reconstruct the fingertip, in terms of feeling, function and appearance? Or does the nail regrow straighter if part of the bone is included in the reconstruction?

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)

Not Specified

Participant information sheet

Condition

Injury, Occupational Diseases, Poisoning: Fingertip injuries

Interventions

Randomised trial of treatment of zone 2 and 3 fingertip amputations. Demographic data, mechanism, level and time of injury will be recorded. Patient to choose between immediate finger shortening (terminalisation) or reconstruction. If reconstruction, they will be randomised to reconstruction with perionychial graft with or without part of the bone in the part of the fingertip that has been amputated.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

To find out which treatment of fingertip injuries gives the best results, terminalisation or reconstruction.

Secondary outcome measures

Not provided at time of registration

Overall study start date

15/01/2004

Overall study end date

01/04/2007

Eligibility

Participant inclusion criteria

60 patients admitted with total or subtotal fingertip amputations to be offered shortening of the finger or reconstruction.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

60

Participant exclusion criteria

Patients with multilevel or associated injuries will be excluded.

Recruitment start date

15/01/2004

Recruitment end date

01/04/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Plastic Surgery

Northwood United Kingdom HA6 2RN

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

Government

Funder Name

West Hertfordshire 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