

Mobilization of the thumb flexor tendon

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 14/04/2015	Condition category Surgery	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0237109529

Study information

Scientific Title

Mobilization of the thumb flexor tendon

Study objectives

Postoperative 'Controlled Active Movement' mobilization method gives a better outcome than the 'Kleinert' mobilisation method after surgical repair of a lacerated thumb flexor tendon. The assessor will be blinded.

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

Health condition(s) or problem(s) studied

Surgery: Repair of thumb tendon

Interventions

Postoperative 'Controlled Active Movement' mobilization method vs 'Kleinert' mobilisation

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

The outcome at 12 weeks post repair of injury.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/02/2002

Completion date

08/12/2005

Eligibility

Key inclusion criteria

80 patients over 16

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

80

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/02/2002

Date of final enrolment

08/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

St Helens & Knowsley Hospitals NHS Trust

Prescot

United Kingdom

L35 5DR

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

St Helens and Knowsley 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