

Basal osteoarthritis of the thumb: a controlled prospective randomised study

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 18/11/2013	Condition category Musculoskeletal Diseases	<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
Mr Neil Citron

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0112112974

Study information

Scientific Title

Study objectives

Trapeziumectomy and interposition arthroplasty versus the same operation combined with a simple suspension operation to limit migration of the thumb base and stabilise it during pinch grip.

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

Osteoarthritis of the thumb

Interventions

1. Trapeziumectomy and interposition arthroplasty
2. Trapeziumectomy and interposition arthroplasty combined with a simple suspension operation

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Which type of operation is most beneficial to this condition?

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/1999

Completion date

01/02/2006

Eligibility

Key inclusion criteria

All patients presenting to the hand clinic of Mr Citron were enrolled in the trial.

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/04/1999

Date of final enrolment

01/02/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Epsom and St Helier NHS Trust

Surrey

United Kingdom

SM5 1AA

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

Epsom and St Helier University 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