

Trapezium implant versus sling procedure in the treatment of thumb carpometacarpal arthritis: a randomised, assessor-blinded controlled trial

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/06/2017	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

N0284126130

Study information

Scientific Title

Trapezium implant versus sling procedure in the treatment of thumb carpometacarpal arthritis: a randomised, assessor-blinded controlled trial

Study objectives

Does a trapezium sling offer no extra benefit in the treatment of thumb metacarpal arthritis over the sling procedure alone?

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

Musculoskeletal Diseases: Osteoarthritis of the thumb

Interventions

Trapezium sling offer vs sling procedure alone. Patients who have had the procedure and patients who have not had the procedure will be assessed.

Intervention Type

Procedure/Surgery

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

29/04/2002

Completion date

01/05/2004

Eligibility

Key inclusion criteria

1. Female
2. Aged 50-80 years

Participant type(s)

Patient

Age group

Mixed

Sex

Female

Target number of participants

40

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

29/04/2002

Date of final enrolment

01/05/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Wrightington, Wigan & Leigh NHS Trust

Wigan

United Kingdom

WN6 9EP

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

Hospital/treatment centre

Funder Name

Wrightington, Wigan and Leigh 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