

A prospective randomised double-blind controlled trial for efficacy of lignocaine with hyaluronidase as a local anaesthetic for carpal tunnel decompression.

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 16/04/2015	Condition category Surgery	<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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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0155151587

Study information

Scientific Title

A prospective randomised double-blind controlled trial for efficacy of lignocaine with hyaluronidase as a local anaesthetic for carpal tunnel decompression.

Study objectives

Does lignocaine with hyaluronidase give a better field of anaesthesia and decrease tissue deformity for carpal tunnel decompression?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Prospective randomised double-blind 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: Anaesthesia

Interventions

Lignocaine with hyaluronidase vs standard practice

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

Lignocaine with hyaluronidase

Primary outcome measure

Field of anaesthesia and tissue deformity

Secondary outcome measures

Not provided at time of registration

Overall study start date

30/04/2004

Completion date

31/12/2005

Eligibility

Key inclusion criteria

60 patients undergoing carpal tunnel decompression

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

60

Key exclusion criteria

Patients with a known contra-indication to lignocaine or lignocaine hyaluronidase and patients undergoing hand surgery additional to carpal tunnel decompression

Date of first enrolment

30/04/2004

Date of final enrolment

31/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

North Manchester General Hospital
Manchester
United Kingdom
M8 5RB

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Government

Funder Name

Pennine Acute Hospitals NHS Trust (UK), Own Account, NHS R&D Support Funding

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