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Infra clavicular plexus block - Lateral (coracoid) approach vs vertical approach - a comparison of techniques

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2005	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
18/10/2017	Surgery	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0249142464

Study information

Scientific Title
Infra clavicular plexus block - Lateral (coracoid) approach vs vertical approach - a comparison of techniques

Study objectives
Which of two infraclavicular approaches to the brachial plexus is preferable

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)
Not specified

Study type(s)
Not Specified

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Surgery: Arm

Interventions
Randomised Controlled Trial:
1. Lateral approach
2. Vertical approach

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Degree of sensory block, complications

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/12/2001

Completion date
09/09/2004

Eligibility

Key inclusion criteria
40 patients undergoing distal upper limb surgery

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
40

Key exclusion criteria
1. Age under 16
2. Uunable to give informed consent
3. Neurovascular compromise of limb
4. Allergy to amide local anaesthetic agents
5. Coagulopathy (INR>2)
6. Chest or shoulder deformities
7. Healed but disclosed clavicle fracture

Date of first enrolment
01/12/2001

Date of final enrolment
09/09/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Specialist Registrar in Anaesthesia
Taunton
United Kingdom
TA1 5DA

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
Hospital/treatment centre

Funder Name
Taunton and Somerset Research and Development Consortium (UK)

Funder Name
NHS R&D Support Funding (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