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Study of timing of ankle block versus inflation of thigh tourniquet in forefoot surgeries

Submission date	Recruitment status	[ ] Prospectively registered
28/09/2007	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
28/09/2007	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
24/10/2016	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
N0155182857

Study information

Scientific Title
Study of timing of ankle block versus inflation of thigh tourniquet in forefoot surgeries

Study objectives
Is there any difference between timing of inflation of thigh tourniquet and ankle block on post-operative analgesia?

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
Forefoot

Interventions
Randomised controlled trial.
Ankle block before inflation of tourniquet or ankle block after inflation of tourniquet.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Amount of post operative analgesia required: time to first pain perception and amount of analgesia in first 24 hours.

Secondary outcome measures

Visual analogue scores for pain.

Overall study start date
01/06/2006

Completion date
31/12/2006

Eligibility

Key inclusion criteria
1. Patients undergoing forefoot surgery
2. Aged 18-60years old
3. Able to provide informed consent

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Upper age limit
60 Years

Sex
Not Specified

Target number of participants
100

Key exclusion criteria
1. Simultaneous iliac crest bone surgery
2. Surgery on the hind foot
3. Patients on long term analgesics for a reason other than foot pain
4. Patients unable to give consent

Date of first enrolment
01/06/2006

Date of final enrolment
31/12/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
J Block
Oldham
United Kingdom
OL1 2JH

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details
The Department of Health, 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

Funder Name
Own Account

Funder Name
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