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

Contact name

Mr M Sundar

Contact details

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Additional identifiers

Protocol serial number

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 postoperative 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

Study type(s)

Treatment

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(s)

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

Key secondary outcome(s))

Visual analogue scores for pain.

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

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

60 years

Sex

Not Specified

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

United Kingdom

England

Study participating centre

J Block

Oldham

United Kingdom

OL1 2JH

Sponsor information

Organisation

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

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration