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

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
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	<ul><li>Record updated in last year</li></ul>

# 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

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

# 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

# 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