# Randomised controlled trial comparing the efficacy of ambulatory local anaesthetic infusion pump versus femoral nerve block in day case anterior cruciate ligament (ACL) reconstruction

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
30/09/2004	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2004	Completed	Results
Last Edited	Condition category	Individual participant data
24/08/2015	Signs and Symptoms	<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

Mr S J Shaw

#### Contact details

Orthopaedics Lancashire Teaching Hospitals NHS Trust Preston United Kingdom PR2 9HT

# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

## Secondary identifying numbers

N0188139053

# Study information

#### Scientific Title

Randomised controlled trial comparing the efficacy of ambulatory local anaesthetic infusion pump versus femoral nerve block in day case anterior cruciate ligament (ACL) reconstruction

## **Study objectives**

To determine the most effective form of postoperative analgesia in day case ACL surgery. This could enhance patient acceptance and satisfaction of day case ACL reconstruction in the future.

## 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

Signs and Symptoms: Post-operative pain

#### **Interventions**

Comparison of local anaesthetic infusion pump with the femoral nerve block for post operative analgesia in day case ACL surgery

## Intervention Type

Procedure/Surgery

## Primary outcome measure

Not provided at time of registration

## Secondary outcome measures

## Not provided at time of registration

# Overall study start date

01/01/2004

# Completion date

01/12/2006

# **Eligibility**

## Key inclusion criteria

Not provided at time of registration

# Participant type(s)

**Patient** 

## Age group

**Not Specified** 

#### Sex

**Not Specified** 

## Target number of participants

50 patients

# Key exclusion criteria

Not provided at time of registration

## Date of first enrolment

01/01/2004

## Date of final enrolment

01/12/2006

# Locations

# Countries of recruitment

England

**United Kingdom** 

# Study participating centre Lancashire Teaching Hospitals NHS Trust

Preston United Kingdom PR2 9HT

# Sponsor information

## Organisation

Department of Health

## Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

# Sponsor type

Government

## Website

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

# Funder(s)

# Funder type

Government

#### **Funder Name**

Lancashire Teaching Hospitals NHS Trust (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