ISRCTN85092388 https://doi.org/10.1186/ISRCTN85092388

Influence of continuous femoral analgesia after anterior cruciate ligament repair on postoperative pain and range of motion: a pilot study

Submission date 11/08/2006	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 29/01/2007	**Overall study status** Completed	
Last Edited 06/02/2008	**Condition category** Musculoskeletal Diseases	

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
2005-000237-37

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Acronym
FemoX

Study objectives
Patients with a catheter-based analgesia at the femoral nerve have less pain compared to patients receiving a Patient-Controlled Analgesia (PCA).

Ethics approval required
Old ethics approval format

Ethics approval(s)
Ethics approval received, no details provided as of 29/01/2007.

Study design
Single-blinded, randomised study.

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Quality of life

Participant information sheet

Health condition(s) or problem(s) studied
Ruptured anterior cruciate ligament

Interventions
Peripheral nerve catheter at the femoral nerve versus PCA.

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
Pain levels.

Secondary outcome measures
Physiotherapeutic options.

Overall study start date
01/01/2006

Completion date
31/12/2006

Eligibility

Key inclusion criteria
Patients receiving a anterior cruciate ligament repair.

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
90

Key exclusion criteria
1. Pregnancy
2. Redo-Surgery
3. Neurologic or psychiatric diseases
4. Other diseases potentially putting the patient at risk
5. Body Mass Index of more than 35
6. Allergic to local anesthetics

Date of first enrolment
01/01/2006

Date of final enrolment
31/12/2006

Locations

Countries of recruitment
Germany

Study participating centre
Charité Universitätsmedizin Berlin
Berlin
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Sponsor information

Organisation
Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin) (Germany)

Sponsor details
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Sponsor type
University/education

Website
http://www.charite.de/ch/anaest/

ROR
https://ror.org/001w7jn25

Funder(s)

Funder type
University/education

Funder Name
Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin), Medical School Research Grant (Germany)

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