

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	<input type="checkbox"/> Prospectively registered
Registration date 29/01/2007	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 06/02/2008	Condition category Musculoskeletal Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

Clinical Trials Information System (CTIS)

2005-000237-37

Protocol serial number

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

Study type(s)

Quality of life

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(s)

Pain levels.

Key secondary outcome(s))

Physiotherapeutic options.

Completion date

31/12/2006

Eligibility

Key inclusion criteria

Patients receiving a anterior cruciate ligament repair.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

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

Germany

10098

Sponsor information**Organisation**

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

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

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration