Investigation into patient characteristics and treatment factors associated with short-term and medium-term outcome following Anterior Cruciate Ligament Reconstruction

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year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Mr Fares S Haddad

Contact details

Department of Trauma & Orthopaedics University College Hospital 235 Euston Road London United Kingdom NW1 2BU

Additional identifiers

Protocol serial number N0209181918

Study information

Scientific Title

Investigation into patient characteristics and treatment factors associated with short-term and medium-term outcome following Anterior Cruciate Ligament Reconstruction

Study objectives

What are the factors pre-operatively, intra-operatively and post-operative and related to patient characteristics that have the most influence on outcomes of ACL reconstruction? Particular focus on post operative/rehabilitation factors (knee swelling and initial rehabilitation) and patient psychological factors (adherence to rehabilitation, motivation and sense of control).

Ethics approval required

Old ethics approval format

Ethics approval(s)

Added 28/08/09:

Joint UCL/UCLH Committees on the Ethics of Human Research (Committee A), Research & Development Directorate, 1st Floor Maple House, 149 Tottenham Court Road, London W1P 9LL. REC reference number: 05/Q0505/120. Meeting date 26th January 2006, approval letter dated 9th February 2006.

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Surgery: Anterior cruciate ligament reconstruction

Interventions

Randomised between 2 groups: normal management vs measures to reduce swelling

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

Questionnaire - International Knee Documentation committee (IKDC) form

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/03/2010

Eligibility

Key inclusion criteria

100 patients recruited from UCLH Foundation Trust, in need of Arthroscopic anterior cruciate ligament reconstruction (ACL) for uncomplicated ACL reconstruction.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

- 1. Concurrent surgery (meniscus repair, posterior cruciate ligament reconstruction, other ligament injuries)
- 2. Patients with gross pre-operative swelling/inflammation, pre-operative loss of normal movement (esp. leg straightening), heavy manual/impact type of work, compulsory return to work before 4 weeks post op).

Date of first enrolment

01/04/2006

Date of final enrolment

01/03/2010

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Department of Trauma & Orthopaedics

London United Kingdom NW1 2BU

Sponsor information

Organisation

Funder(s)

Funder type

Government

Funder Name

UCL Hospitals NHS Foundation Trust (UK), NHS R&D Support Funding

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type Details Date created Date added Peer reviewed? Patient-facing?

Participant information sheet Participant information sheet 11/11/2025 11/11/2025 No Yes