

A randomised prospective comparison of two anterior cruciate ligament graft fixation techniques

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 27/02/2020	Condition category Surgery	<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

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0234098041

Study information

Scientific Title

A randomised prospective comparison of two anterior cruciate ligament graft fixation techniques

Study objectives

Comparing two different methods of fixing an anterior cruciate ligament (ACL) graft to the tibia in patients undergoing ACL reconstruction. To identify any difference in knee stability, range of movement or return to pre-injury levels of activity between the two groups.

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

Anterior cruciate ligament (ACL) graft

Interventions

Randomised controlled trial. Patients randomised to one of two methods of fixation of ACL grafts.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Post operative knee laxity

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2001

Completion date

31/05/2003

Eligibility

Key inclusion criteria

80 Patients undergoing ACL reconstruction.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

80

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/06/2001

Date of final enrolment

31/05/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Orthopaedic Surgery

Bristol

United Kingdom
BS16 1LE

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

North Bristol 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