

Anterior Cruciate Ligament (ACL) Reconstruction using two different types of Femoral Fixation i.e. Mitec Rigidfix femoral polylactide (PLA) cross pin and the Anthrex Bio Transfix implant

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 13/02/2017	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

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Additional identifiers

Protocol serial number
N0020129460

Study information

Scientific Title

Anterior Cruciate Ligament (ACL) Reconstruction using two different types of Femoral Fixation i. e. Mitec Rigidfix femoral polylactide (PLA) cross pin and the Anthrex Bio Transfix implant

Study objectives

Is there any difference in the outcome of the ACL reconstruction using two types of femoral fixation?

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Surgery: Anterior cruciate ligament reconstruction

Interventions

Patients with ACL deficiency to be reconstructed using hamstring graft (4 strands) with two types of fixture.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

International Knee Documentation Committee (IKDC) scoring system - tagner scoring system

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/09/2006

Eligibility

Key inclusion criteria

100 ligament reconstruction patients

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/09/2003

Date of final enrolment

01/09/2006

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Oldchurch Hospital

Romford

United Kingdom

RM7 0BE

Sponsor information**Organisation**

Department of Health

Funder(s)

Funder type

Government

Funder Name

Barking, Havering and Redbridge Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration