

Trial of patellofemoral (PF) arthroplasty versus total knee joint replacement (TKR)

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
12/09/2003	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
12/09/2003	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
14/05/2018	Surgery	<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

N0234088724

Study information

Scientific Title

Trial of patellofemoral (PF) arthroplasty versus total knee joint replacement (TKR)

Study objectives

To study the differences in patient based outcome measures, as a comparison between patellofemoral arthroplasty and total knee replacement in isolated osteoarthritis of the patellofemoral compartment.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled parallel group trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Knee surgery

Interventions

Prospective randomised trial.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

Patient based outcome measures to determine which type of prosthesis is best.

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/09/2003

Eligibility

Key inclusion criteria

Patients who are suitable for patellofemoral replacement and are on the waiting list for surgery.
50 Patients in each group aged over 50 years.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Other

Sex

All

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/10/2000

Date of final enrolment

30/09/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Winford Unit - AOC

Bristol

United Kingdom

BS10 5NB

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration