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

Submission date	Recruitment status	Prospectively registered
12/09/2003	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
12/09/2003	Completed	Results
Last Edited	Condition category	Individual participant data
14/05/2018	Surgery	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Mr JH Newman

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Knee surgery

Interventions

Prospective randomised trial.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

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

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/2000

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

Age group

Other

Sex

Both

Target number of participants

100

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

England

United Kingdom

Study participating centre Winford Unit - AOC

Bristol United Kingdom BS10 5NB

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