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Why do many patients with unicompartmental knee replacements have difficulty kneeling after surgery?

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 30/09/2004	**Overall study status** Completed	
Last Edited 27/09/2011	**Condition category** Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0176127672

Study information

Scientific Title

Study objectives
What is the patients' perception of the barrier to kneeling post operatively? Is there a relationship between the scar position and kneeling ability? What anatomical structure is knelt on post operatively? Can a one-off physiotherapy intervention improve patients' kneeling ability?

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)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Surgery: Knee replacement

Interventions
Post operative follow up patients who agree to take part within the study will complete a questionnaire about kneeling at 6 weeks post op and 12 months post op.
Half will receive the intervention at 6 weeks. The intervention is advice on kneeling techniques. The other half will receive the intervention at 12 months and the results will be compared.

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
Change in kneeling questionnaire and in Oxford knee score.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/2003

Completion date
31/08/2005

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
50 patients

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/09/2003

Date of final enrolment
31/08/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Physiotherapy Dept
Oxford
United Kingdom
OX3 7LD

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Oxford Radcliffe Hospitals 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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/09/2008		Yes	No