

Who benefits from primary hip replacement? A feasibility study for a waiting list randomised controlled trial

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

N0264118306

Study information

Scientific Title

Who benefits from primary hip replacement? A feasibility study for a waiting list randomised controlled trial

Study objectives

Assess the feasibility of undertaking a full-scale waiting-list trial of hip replacement surgery.

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)

Other

Participant information sheet

Health condition(s) or problem(s) studied

Surgery: Hip replacement

Interventions

Feasibility study for a randomised controlled trial.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Better management of waiting lists, based on patient need rather than time waited.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2002

Completion date

01/06/2003

Eligibility

Key inclusion criteria

30 Patients from all age groups: 15 randomised for immediate surgery; 15 randomised to waiting list.

Participant type(s)

Patient

Age group

Other

Sex

Both

Target number of participants

30

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/06/2002

Date of final enrolment

01/06/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

c/o Research and Development Office

Bristol

United Kingdom

BS2 8HW

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

United Bristol Healthcare 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