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To assess two methods of surgical repair of posterior vaginal wall prolapse

Submission date	Recruitment status	[_] Prospectively registered
30/09/2004	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
30/09/2004	Completed	[_] Results
Last Edited	**Condition category**	[_] Individual participant data
18/07/2016	Urological and Genital Diseases	[_] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0241124939

Study information

Scientific Title
To assess two methods of surgical repair of posterior vaginal wall prolapse

Study objectives
To assess two methods of surgical repair of posterior vaginal wall prolapse.

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)
Treatment

Participant information sheet
Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied
Surgery: Gynaecological

Interventions
Patients randomised into two groups, one group will undergo a fascial repair, the other group will undergo a standard levator plication.

Intervention Type
Procedure/Surgery

Primary outcome measure
Comparison of results and questionnaire

Secondary outcome measures
Not provided at time of registration

Overall study start date
08/07/2003

Completion date
31/03/2006

Eligibility

Key inclusion criteria
100 patients with posterior vaginal wall prolapse requiring surgical repair

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
100

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
08/07/2003

Date of final enrolment
31/03/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
St Mary's Hospital
London
United Kingdom
W2 1NY

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
Hospital/treatment centre

Funder Name
St Mary's 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